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# Amino Acid Analysis Protocols

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# Amino Acid Analysis

## *An Overview*

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### 1. Importance and Utility

Amino acids are found either in the free state or as linear chains in peptides and proteins. There are 20 commonly occurring amino acids in proteins, which are shown in **Table 1**. Amino acid analysis has an important role in the study of the composition of proteins, foods, and feedstuffs. Free amino acids are also determined in biological material, such as plasma and urine, and in fruit juice and wine. When it is performed on a pure protein, amino acid analysis is capable of identifying the protein (2,3, and Chapter 8 in this volume), and the analysis is also used as a prerequisite for Edman degradation and mass spectrometry and to determine the most suitable enzymatic or chemical digestion method for further study of the protein. It is also a useful method for quantitating the amount of protein in a sample (*see* Chapter 2 in this volume) and can give more accurate results than colorimetric methods.

### 2. Historical View

The earliest experiments on the acid hydrolysis of proteins were performed by Braconnot in 1820, in which concentrated sulphuric acid was used to hydrolyze gelatin, wool, and muscle fibers (4). Various reagents for performing protein hydrolysis were tried over the next 100 years, with 6 M HCl becoming the most widely accepted reagent. In 1972, Moore and Stein (5) were awarded the Nobel Prize for developing an automated instrument for separation of amino acids on an ion-exchange resin and quantitation of them using ninhydrin.

More recently, high-performance liquid chromatographs (HPLCs) have been configured for amino acid analysis. Some methods use postcolumn deriv-

**Table 1**  
**Common Amino Acids**

| Name                | Symbol   |          | Essential<br>for<br>humans (1) |
|---------------------|----------|----------|--------------------------------|
|                     | 3 letter | 1 letter |                                |
| Acidic amino acids  |          |          |                                |
| Aspartic acid       | Asp      | D        | No                             |
| Glutamic acid       | Glu      | E        | No                             |
| Neutral amino acids |          |          |                                |
| Alanine             | Ala      | A        | No                             |
| Asparagine          | Asn      | N        | No                             |
| Cysteine            | Cys      | C        | No                             |
| Glutamine           | Gln      | Q        | No                             |
| Glycine             | Gly      | G        | No                             |
| Isoleucine          | Ile      | I        | Yes                            |
| Leucine             | Leu      | L        | Yes                            |
| Methionine          | Met      | M        | Yes                            |
| Phenylalanine       | Phe      | F        | Yes                            |
| Serine              | Ser      | S        | No                             |
| Threonine           | Thr      | T        | Yes                            |
| Tryptophan          | Trp      | W        | Yes                            |
| Tyrosine            | Tyr      | Y        | No                             |
| Valine              | Val      | V        | Yes                            |
| Basic amino acids   |          |          |                                |
| Arginine            | Arg      | R        | Yes                            |
| Histidine           | His      | H        | Yes                            |
| Lysine              | Lys      | K        | Yes                            |
| Imino acid          |          |          |                                |
| Proline             | Pro      | P        | No                             |

atization in which the amino acids are separated on an ion-exchange column followed by derivatization with ninhydrin (**6**, and Chapter 2 in this volume), fluorescamine (**7**), or o-phthalaldehyde (**8**). Another approach has been to derivatize amino acids prior to separation on a reversed-phase HPLC column. Examples of this technique are dansyl (**9**), phenylisothiocyanate (PITC) (**10**, and Chapters 12 and 13 of this volume), 9-fluorenylmethyl chloroformate (Fmoc) (**11**), and 6-aminoquinolyl-N-hydroxysuccinimyl carbamate (AQC) (**12**, and Chapters 4 and 8 in this volume).

### 3. Sensitivity

Amino acid analysis can be performed accurately at the fmol level by methods employing fluorescence detection, whereas for derivatives detected by

**Table 2**  
**Comparison of Different Derivatization Chemistries**  
**for Amino Acid Analysis**

|                                  | Ninhydrin | OPA   | OPA  | PITC | Fmoc | AQC  |
|----------------------------------|-----------|-------|------|------|------|------|
| Derivatization type <sup>a</sup> | postc     | postc | prec | prec | prec | prec |
| Detection mode <sup>b</sup>      | c         | f     | f    | UV   | f    | f    |
| Sensitivity                      | pmol      | fmol  | fmol | pmol | fmol | fmol |
| Chromatography <sup>c</sup>      | i.e.      | i.e.  | r.p. | r.p. | r.p. | r.p. |

OPA, orthophthalaldehyde; PITC, phenylisothiocyanate; Fmoc, 9-fluorenylmethyl chloroformate; AQC, 6-aminoquinolyl-N-hydroxysuccinimidyl carbamate.

<sup>a</sup>postc, postcolumn; prec, precolumn.

<sup>b</sup>c, colorimetry; f, fluorescence; UV, ultraviolet.

<sup>c</sup>i.e, ion exchange; r.p., reversed-phase HPLC.

ultraviolet (UV) light, the analysis is at the pmol level. **Table 2** gives a comparison of the various derivatization chemistries and their sensitivities. Annual studies comparing the various methods have been carried out by the Association of Biomolecular Resource Facilities (ABRF) (**13,14**). Strydom and Cohen (**15**) have compared AQC and PITC chemistries and found AQC derivatives to be more stable.

## 4. Difficult Amino Acids

### 4.1. Tryptophan

Tryptophan is destroyed in acid hydrolysis. Alkaline hydrolysis with NaOH, Ba (OH)<sub>2</sub>, or LiOH have been used particularly in the hydrolysis of food and feedstuffs (**16,17**). However, acid hydrolysis is still needed to determine the other amino acids.

There have been a number of methods published for the determination of tryptophan that use the standard 6 M HCl hydrolysis in the presence of additives, some of which include thioglycolic acid (**18**), beta-mercapto ethanol (**19**), and mercaptoethanesulfonic acid (**20**).

### 4.2. Cysteine and Cystine

Cysteine and cystine are unstable during acid hydrolysis, particularly in the presence of carbohydrate. The total content of cysteine and cystine can be determined by oxidizing the protein with performic acid, which converts both forms to cysteic acid and methionine to methionine sulphone. The protein is then hydrolyzed with 6 M HCl (**17**).

Disulphide compounds such as dithiopropionic acid and dithiobutyric acid have been proposed as protecting agents for cysteine and cystine during acid

hydrolysis (21). The use of dithiodiglycolic acid during the acid hydrolysis step, followed by phenylisothiocyanate derivatization, allows cysteine and cystine plus all common hydrolysate amino acids (excluding tryptophan) to be determined (22).

Reduction of disulphide bridges, followed by alkylation of cysteine with iodoacetic acid or 4-vinylpyridine is also used to determine the cysteine-plus-cystine content of proteins (23). Alkylation with acrylamide produces cysteine-S-propionamide, which is converted to cysteine-S-propionic acid during acid hydrolysis (24).

### 4.3. Asparagine and Glutamine

Asparagine and glutamine are amide derivatives of aspartic acid and glutamic acid, respectively. During acid hydrolysis, which cleaves amide bonds, asparagine is converted to aspartic acid and glutamine to glutamic acid. Thus, the amount determined for aspartic acid represents the total of aspartic acid and asparagine and similarly for glutamic acid and glutamine.

### 4.4. D amino Acids

The D-amino acid content of a protein or peptide can be determined by employing a short partial acid hydrolysis, followed by an enzymatic hydrolysis with pronase, and then with leucine aminopeptidase and peptidyl-D-amino acid hydrolase (25).

## 5. Modified Amino Acids

Phosphorylated amino acids are able to be analyzed using a variety of different chemistries (26), but the ABRF 1993 study found that precolumn methods were more successful (27). Phosphoserine (28,29), phosphothreonine (29), and phosphotyrosine (29) have varying stabilities. Highest recoveries for phosphoserine and phosphotyrosine are produced with hydrolysis time of 60 min or less at 110°C, whereas for phosphothreonine, a hydrolysis time of 2 h gave better results (26). Chapter 14 in this volume covers the analysis of phosphoamino acids more extensively.

There are many other rarer amino acids and derivatives that can be analyzed. These include hydroxyproline (17,30, and Chapter 16 in this volume) and hydroxylysine (17,30, and Chapter 2 of this volume), found in collagen. Taurine has dietary importance and can be readily determined in infant formulas, pet food, plasma, urine, and tissue extracts (17, and Chapter 10 of this volume). Posttranslational modifications, including glycosylated amino acids (31, and Chapters 2 and 7 of this volume) and glycated amino acids (32,33), are important in studying protein function. Chapter 18 of this volume describes the application of mass spectrometry to the analysis of glycated amino acids.

## 6. Limitations—Contaminants and Precautions

The accuracy of amino acid analysis is very dependent on the integrity of the sample. Cleanliness of all surfaces the sample contacts is essential, as is the purity of all reagents used. Traces of salts, metals, or detergents can effect the accuracy of results.

The hydrolysis step is particularly important, as was demonstrated in the ABRF 1994 AAA collaborative study (34). Many laboratories now satisfactorily perform a 1-h hydrolysis in 6 M HCl at 150°C under vacuum. The traditional method uses 6 M HCl for 20–24 h at 110°C under vacuum. Losses of serine, threonine, and to a lesser extent, tyrosine may occur under these conditions. During acid hydrolysis, some amide bonds between aliphatic amino acids are more difficult to cleave. The Ala–Ala, Ile–Ile, Val–Val, Val–Ile, Ile–Val, and Ala–Val linkages are resistant to hydrolysis and may need a longer hydrolysis time of 48 or 72 h at 110°C (35).

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## Amino Acid Analysis, Using Postcolumn Ninhydrin Detection, in a Biotechnology Laboratory

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### 1. Introduction

Although lacking the speed and sensitivity of more widely heralded techniques such as mass spectrometry, amino acid analysis remains an indispensable tool in a complete biotechnology laboratory responsible for the analysis of protein pharmaceuticals.

Moore and Stein developed the first automated amino acid analyzer, combining cation-exchange chromatographic separation of amino acids with postcolumn ninhydrin detection (**1**). Commercial instruments based on this design were introduced in the early 1960s, though many manufacturers have abandoned this technology in favor of precolumn amino acid derivatization with separations based on reversed-phase chromatography (**2–4**) (*see Note 1*).

In our product development role, we still rely on amino acid analysis to generate key quantitative and qualitative data. Amino acid analysis after acid hydrolysis remains the best method for absolute protein/peptide quantitation, limited in accuracy and precision only by sample handling. We produced an Excel macro to process these data; the macro transfers and converts the amino acid molar quantities into useful values such as composition (residues per mol) and concentration. In addition, we employ several specialized amino acid analysis applications to monitor structural aspects of some of our recombinant products.

*De novo* biosynthesis of leucine in bacteria will lead to a minor amount of norleucine (Nle) production (**5**), particularly if recombinant proteins are produced in fermentations that have been depleted of leucine (**6**). The side-chain of Nle ( $-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_3$ ) is similar enough to methionine ( $-\text{CH}_2-\text{CH}_2-\text{S}-\text{CH}_3$ ) that some of the tRNA<sup>Met</sup> will be acylated by Nle, leading to incorporation of Nle at

Met positions (6,7). When this occurs, Nle may be incorporated at a low level at every Met position, and amino acid analysis is often the only method able to detect this substitution.

Hydroxylysine (Hyl) is a common modification of lysine residues found at *-Lys-Gly-* positions in collagens and collagen-like domains of modular proteins (8). This modification is also found at certain solvent-accessible *-Lys-Gly-* sites in noncollagenous proteins, usually at substoichiometric levels (9). Amino acid analysis is a useful screening technique for the identification of Hyl-containing recombinant proteins produced by mammalian cells.

The analysis of recombinant proteins using carboxypeptidases may still be required to assign the C-terminus when the polypeptide chain is extensively modified, thus ruling out making a C-terminal assignment based solely on mass and N-terminal analyses, or in cases where the C-terminal peptide cannot be assigned in a peptide map. When carboxypeptidase analyses are needed, a modified amino acid analysis program is needed to resolve Gln and Asn (which are not found in acid hydrolysates) from other amino acids.

Assignment of Asn-linked glycosylation sites is greatly facilitated by prior knowledge of the *-Asn-Xaa-Thr/Ser/Cys-* consensus sequence sites (10), and specific endoglycosidases, such as peptide:N-glycosidase F can be employed to quantitatively release all known types of Asn-linked oligosaccharides (11). O-linked sites are harder to assign, as these are found in less-stringent sequence motifs (12–14), and there is no universal endoglycosidase for O-glycans except for endo- $\alpha$ -*N*-acetylgalactosaminidase, which can only release the disaccharide Gal( $\beta$ 1 $\rightarrow$ 3)GalNAc. In addition, O-glycosylation is often substoichiometric.

In mammalian cell products, at least two N-acetylglucosamine (GlcNAc) residues are found in Asn-linked oligosaccharides, whereas N-acetylgalactosamine (GalNAc) is found at the reducing terminus of the most common (mucin-type) O-linked oligosaccharides. A cation-exchange-based amino acid analyzer can easily be modified for the analysis of the amino sugars glucosamine (GlcNH<sub>2</sub>) and galactosamine (GalNH<sub>2</sub>) from acid hydrolysis of GlcNAc and GalNAc, respectively, allowing confirmation of the presence of most oligosaccharide types. In glycoproteins, HPLC fractions from peptide digests can be screened using amino sugar analysis to identify glycopeptides for further analysis.

Regulated biotechnology products are usually tested for identity using HPLC maps after peptide digestion (15,16). A key aspect of the digestion step for most proteins is obtaining complete reduction of all disulfide bonds, followed by complete alkylation of cysteines without the introduction of artifacts (e.g., methionine S-alkylation) (17). Amino acid analysis can be used to monitor cysteine alkylation levels for reduced proteins, such as are obtained after alkylation with iodoacetic acid, iodoacetamide or 4-vinylpyridine.

## 2. Materials

### 2.1. Equipment

1. 1-mL hydrolysis ampoules (Bellco, Vineland, NJ; part number 4019-00001) (*see Note 2*).
2. Savant SpeedVac.
3. Oxygen/methane flame.
4. Glass knife (Bethlehem Apparatus Co., Hellertown, PA).
5. 1/4" ID  $\times$  5/8" OD Tygon tubing.
6. Model 6300 analyzers (Beckman Instruments, now Beckman Coulter, Fullerton CA). The sum of the 440 nm and 570 nm absorbances is converted to digital format using a PE Model 900 A/D converter, and the data are collected by a PE Turbochrom Model 4.1 data system (*see Notes 3–5*).
7. Lithium-exchange column (Beckman part number 338075, 4.6  $\times$  200 mm).

### 2.2. Reagents and Solutions

1. Constant boiling (6 *N*) HCl ampoules are obtained from Pierce (Rockland, IL) (*see Note 6*).
2. Mobile phase buffers purchased from Beckman Instruments include sodium citrate buffers Na-D, Na-E, Na-F, Na-R, and Na-S; lithium citrate buffers include Li-A, Li-B, Li-C, Li-D, Li-R, and Li-S.
3. Ninhydrin kits (Nin-Rx) are also purchased from Beckman; these must be mixed thoroughly before use (usually 2 h at room temperature), and care must be taken to avoid skin discoloration because of contact with ninhydrin-containing materials.
4. Dialysis may be used to desalt samples into dilute acetic acid prepared from deionized water (Milli-Q, Millipore) and Mallinckrodt U.S.P. grade glacial acetic acid.
5. Amino acid standards: are diluted from the stock Beckman standard (part number 338088) with Na-S buffer to final concentration of 40 nmol/mL or 20 nmol/mL (*see Note 7*).
6. 2 *N* glacial acetic acid.

## 3. Methods

### 3.1. Sample Preparation

Proteins should be desalted to obtain optimal compositional data. Dialysis against 0.1% acetic acid removes salts while keeping proteins in solution, but quantitative data will often require direct hydrolysis (i.e., without dilution or sample losses introduced during dialysis). When proteins must be analyzed without desalting, neutral buffers such as 50 mM Tris can be used without compromising the results. Excipients to avoid include urea (which generates abundant ammonia during hydrolysis), sugars (which caramelize during hydrolysis), and detergents such as the polysorbate and Triton types that can

**Table 1**  
**Standard Amino Acid Analysis**

| Time (min) | Event                   | Conditions            |
|------------|-------------------------|-----------------------|
| 0.0        | Sample injection        | Na-E buffer, 48°C     |
| 8.5        | Start temp. gradient    | 48°C to 60°C in 8 min |
| 24.5       | Buffer change           | Na-E to Na-F          |
| 41.0       | Buffer change           | Na-F to Na-D          |
| 78.0       | Reagent pump            | Ninhydrin to water    |
| 79.0       | Buffer change           | Na-D to Na-R          |
| 80.0       | Buffer change           | Na-R to Na-E          |
| 82.5       | Temperature change      | 60°C to 48°C          |
| 84.0       | Reagent pump            | Water to ninhydrin    |
| 97.0       | Recyle (start next run) |                       |

Buffer pump: 16 mL/h.

Reagent pump: 8 mL/h.

damage cation-exchange columns. Samples in enzyme-linked immunosorbent assays (ELISA)-type buffers should be avoided as they typically contain albumin or gelatins, whose amino acids cannot be distinguished after hydrolysis from the protein of interest. Peptides generally can be desalted by reverse phase (RP)-HPLC using volatile solvents such as 0.1% TFA in water/acetonitrile.

1. Place samples in hydrolysis ampoules (*see Note 8*), then dry under vacuum using a Savant SpeedVac.
2. Place approx 100  $\mu$ L of 6 N HCl in the lower part of the ampoule (*see Note 9*). Freeze in a dry ice/ethanol bath, attached to a vacuum system via 1/4" ID  $\times$  5/8" OD Tygon tubing, then slowly thaw and evacuate to < 150 mtorr.
3. Use an oxygen/methane flame to seal the neck of the tube at the constriction.
4. Place the sealed ampoules in a 110°C oven for 24 h (*see Note 10*), then allow to cool before opening after scoring them with a glass knife.
5. Remove the acid by vacuum centrifugation, again using a Savant system, with a NaOH trap inserted between the centrifuge and cold trap.
6. After hydrolysis and acid removal, samples that contain 0.5–10  $\mu$ g of protein, or 0.1–1 nmol of peptide fractions should be reconstituted with 60–200  $\mu$ L of Na-S sample buffer (*see Note 11*).

### 3.2. Protein/Peptide Quantitation

1. Subject triplicate samples containing 0.5–10  $\mu$ g of protein or 0.1–1 nmol of peptide to 24-h hydrolysis *in vacuo* as aforementioned.
2. Follow the standard operating conditions given in **Table 1** (*see Note 12*). A standard chromatogram containing 2 nmol of each component is shown in **Fig. 1**.

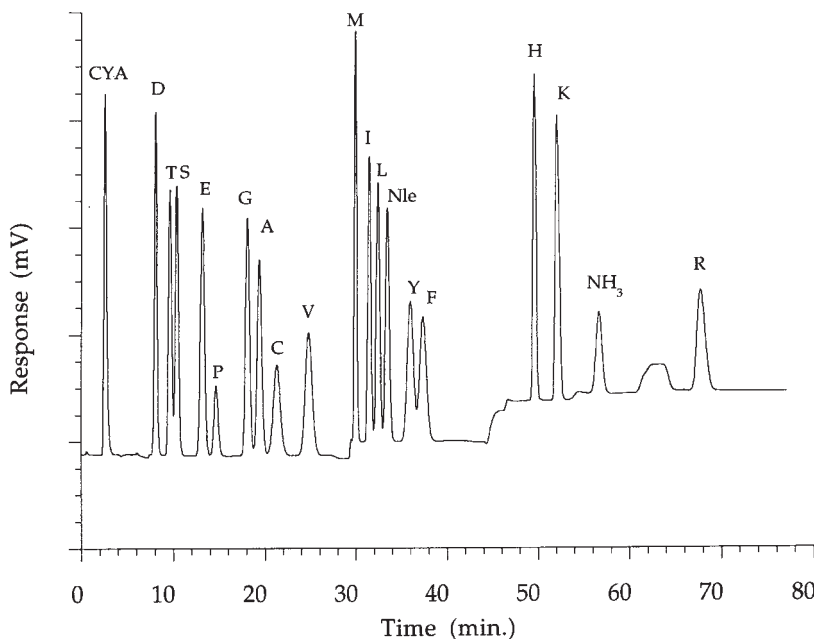


Fig. 1. Analysis of a standard amino acid mixture. The standard contains 2 nmol of each component except for  $\text{NH}_3$ . Operating parameters are given in **Table 1**.

3. Peak area data from the Turbochrom system are converted to nmol values by external standard calibration; internal standards are not necessary if a reliable autosampler is used.
4. The amino acid nmol values are also automatically converted to .txt files that can be imported into a custom Microsoft Excel program called the AAA MACRO (**Table 2**) for analysis using a PC-based computer.
5. The first step in running the AAA MACRO is to open a template, such as the example "protein.xls" given in **Table 3**. The residues per mol and molecular mass calculations must be modified and saved for each different protein/peptide; Asn and Asp are reported as Asx, whereas Gln, Glu and pyroglutamate are reported as Glx.
6. The macro asks for some background information (e.g., requestor's name, sample name, number of replicates), sample prep information (e.g., volumes of hydrolysate loaded vs reconstitution volume, original sample volume), then processes the data, providing a single-page report showing calculated compositions and concentration, as shown in **Fig. 2** (see **Note 13–15**).

### 3.3. Norleucine Incorporation

1. Detection of trace Nle levels in *Escherichia coli*-derived proteins require 24-h hydrolysis of 25–100  $\mu\text{g}$  of protein (see **Note 16**).

**Table 2**  
**Amino Acid Analysis Data Conversion Macro**

Commands

|                                                                                      |                                                        |
|--------------------------------------------------------------------------------------|--------------------------------------------------------|
| Macro 4(a)                                                                           |                                                        |
| =ACTIVATE("MACRO4A.XLM")                                                             |                                                        |
| =HIDE()                                                                              |                                                        |
| =\TC4\DATA                                                                           |                                                        |
| =SELECT(!B4)                                                                         |                                                        |
| =INPUT("Requestor's Name?",2,"Name", "")                                             | Asks for Requestor's Name                              |
| =IF(A7=FALSE,HALT())                                                                 | Halts macro if Cancel button is clicked                |
| =FORMULA(A7)                                                                         | Returns name to data worksheet cell B4                 |
| =SELECT(!B5)                                                                         |                                                        |
| =INPUT("Requestor's Extension?",1,"Telephone extension", "")                         | Asks for Requestor's Extension                         |
| =IF(A11=FALSE,HALT())                                                                | Halts macro if Cancel button is clicked                |
| =FORMULA(A11)                                                                        | Returns extension to data worksheet cell B5            |
| =SELECT(!E4)                                                                         |                                                        |
| =INPUT("Sample to be analyzed?",2,"Sample name", "")                                 | Asks for protein to be analyzed                        |
| =IF(A15=FALSE,HALT())                                                                | Halts macro if Cancel button is clicked                |
| =FORMULA(A15)                                                                        | Returns protein's name to data worksheet cell E4       |
| =SELECT(!E5)                                                                         |                                                        |
| =INPUT("Requestor's Mail Stop?",1,"Mail Stop", "")                                   | Asks for Requestor's MailStop                          |
| =IF(A19=FALSE,HALT())                                                                | Halts macro if Cancel button is clicked                |
| =FORMULA(A19)                                                                        | Returns Requestor's MailStop to data worksheet cell E5 |
| =SELECT(!B28)                                                                        |                                                        |
| =INPUT("Molecular Mass of protein to be analyzed?",1, "Molecular Mass (g/mole)", "") | Asks for MW                                            |
| =IF(A23=FALSE,HALT())                                                                | Halts macro if Cancel button is clicked                |
| =FORMULA(A23)                                                                        | Returns MW to data worksheet cell B28                  |
| =SELECT(!F29)                                                                        |                                                        |
| =INPUT("µL in Ampoule?",1,"Ampoule volume (µL)", "")                                 | Asks for amount sample put in ampoule                  |
| =IF(A27=FALSE,HALT())                                                                |                                                        |
| =FORMULA(A27)                                                                        |                                                        |
| =SELECT(!F30)                                                                        |                                                        |
| =INPUT("Sample reconstitution volume?",1,"Reconstituted volume (µL)", "")            | Asks for reconstitution volume of sample               |
| =IF(A31=FALSE,HALT())                                                                | Halts macro if Cancel button is clicked                |
| =FORMULA(A31)                                                                        |                                                        |
| =SELECT(!F31)                                                                        |                                                        |

```

=INPUT("Dilution Factor?",1,"Dilution factor", "")
=IF(A35=FALSE,HALT())
=FORMULA(A35)
=SELECT(!B1)
=INPUT("How many replicates will you be analyzing today?", 1,"Number of replicates", "")
=IF(A39=FALSE,HALT())
=FORMULA(A39)
=SET.NAME("counter1",1)
=SELECT(!B41)
=DIRECTORY("\\TC41\data")
=FILES("*.*)"
=OPEN?("*.*",0,FALSE,2)
=SELECT("R38C6:R57C6")
=COPY()
=CLOSE()
=ACTIVATE.NEXT()
=SELECT("RC[1]")
=PASTE()
=SET.NAME("counter1",counter1+1)
=IF(counter1<=(B1),GOTO(A46))

=SELECT(!C40)
=INPUT("What is the name of your first replicate?",2, "Name of 1st replicate", "")
=IF(A56=FALSE,HALT())
=FORMULA(A56)
=SELECT(!D40)
=INPUT("What is the name of your second replicate?",2, "Name of 2nd replicate", "")
=IF(A60=FALSE,HALT())
=FORMULA(A60)
=SELECT(!E40)
=INPUT("What is the name of your third replicate?",2, "Name of 3rd replicate", "")
=IF(A64=FALSE,HALT())
=FORMULA(A64)
=SAVE.AS?(,1)
=IF(A67=FALSE,HALT())
=ACTIVATE("MACRO4A.XLM")
=UNHIDE()
=ACTIVATE.NEXT()
=RETURN()

```

Asks if samples were diluted prior to analysis  
Halts macro if Cancel button is clicked

Selects cell B1 on worksheet  
Asks how many replicate samples will be processed  
Halts macro if Cancel button is clicked  
Places users sample number in cell B2  
Resets counter1  
Selects active cell to be B6  
Selects disk in drive as AAA directory  
Opens AAA disk  
Top of loop and Select data file from listed files  
Select data file to open  
Copies AAA nanomole data  
Closes data file

Pastes nanomole values into worksheet  
Adds value of 1 to the counter1  
Checks to see what value = counter1 if >=1 then loops up to the  
top of the loop, if 0 then proceeds downward

Asks for name of first data file chosen

Asks for name of second data file chosen

Asks for name of third data file chosen

Asks if you want to save data  
Halts macro if Cancel button is clicked

Halts the macro

---

**Table 3**  
**AAA Macro Template**

| A                             |                                                   | B                                   |                    | C                                   |  | D |  |
|-------------------------------|---------------------------------------------------|-------------------------------------|--------------------|-------------------------------------|--|---|--|
| AAA Macro                     |                                                   | 3                                   |                    | Protein Template                    |  |   |  |
| Name:                         |                                                   | Researcher:                         |                    | Protein Name:                       |  |   |  |
| Extension:                    | 0                                                 |                                     |                    | Mail Stop:                          |  |   |  |
| Amino Acid                    | Theoretical Composition                           | = (C66)                             | = (D66)            |                                     |  |   |  |
| CyA                           | 0                                                 | = C67                               | = D67              |                                     |  |   |  |
| Asx                           | 35                                                | = C68                               | = D68              |                                     |  |   |  |
| Thr                           | 37                                                | = C69                               | = D69              |                                     |  |   |  |
| Ser                           | 59                                                | = C70                               | = D70              |                                     |  |   |  |
| Glx                           | 41                                                | = C71                               | = D71              |                                     |  |   |  |
| Pro + Cys SH                  | 24                                                | = C72                               | = D72              |                                     |  |   |  |
| Gly                           | 33                                                | = C73                               | = D73              |                                     |  |   |  |
| Ala                           | 30                                                | = C74                               | = D74              |                                     |  |   |  |
| 1/2 Cys-Cys                   | 10                                                | = C75                               | = D75              |                                     |  |   |  |
| Val                           | 35                                                | = C76                               | = D76              |                                     |  |   |  |
| Met                           | 3                                                 | = C77                               | = D77              |                                     |  |   |  |
| Ile                           | 12                                                | = C78                               | = D78              |                                     |  |   |  |
| Leu                           | 32                                                | = C79                               | = D79              |                                     |  |   |  |
| Nle                           | 0                                                 | = C80                               | = D80              |                                     |  |   |  |
| Tyr                           | 22                                                | = C81                               | = D81              |                                     |  |   |  |
| Phe                           | 13                                                | = C82                               | = D82              |                                     |  |   |  |
| His                           | 7                                                 | = C83                               | = D83              |                                     |  |   |  |
| Lys                           | 27                                                | = C84                               | = D84              |                                     |  |   |  |
| Arg                           | 12                                                | = C86                               | = D86              |                                     |  |   |  |
| Molecular Mass (g/mole)       | 47503.01                                          |                                     |                    |                                     |  |   |  |
|                               | ul in Ampoule<br>smp recon(ul)<br>dilution factor |                                     |                    |                                     |  |   |  |
| column load (ul)              | 50                                                |                                     |                    |                                     |  |   |  |
|                               | nMoles Protein                                    | = F62                               | = G62              |                                     |  |   |  |
|                               | Protein Concentration mg/mL                       | = [(C33*B28*0.001*F30/B32)/F29]*F31 |                    | = [(D33*B28*0.001*F30/B32)/F29]*F31 |  |   |  |
| Amino Acid                    | Theoretical Composition                           | data file #1                        | data file #2       |                                     |  |   |  |
| CyA                           | = B7                                              | 0.01                                | 0.011              |                                     |  |   |  |
| Asx                           | = B8                                              | 5.132                               | 5.07               |                                     |  |   |  |
| Thr                           | = B9                                              | 5.146                               | 5.086              |                                     |  |   |  |
| Ser                           | = B10                                             | 7.692                               | 7.58               |                                     |  |   |  |
| Glx                           | = B11                                             | 5.875                               | 5.8                |                                     |  |   |  |
| Pro + CySH                    | = B12                                             | 4.415                               | 4.227              |                                     |  |   |  |
| Gly                           | = B13                                             | 4.831                               | 4.76               |                                     |  |   |  |
| Ala                           | = B14                                             | 4.545                               | 4.474              |                                     |  |   |  |
| 1/2 Cys-Cys                   | = B15                                             | 1.263                               | 1.375              |                                     |  |   |  |
| Val                           | = B16                                             | 4.922                               | 4.851              |                                     |  |   |  |
| Met                           | = B17                                             | 0.39                                | 0.365              |                                     |  |   |  |
| Ile                           | = B18                                             | 1.68                                | 1.662              |                                     |  |   |  |
| Leu                           | = B19                                             | 4.68                                | 4.812              |                                     |  |   |  |
| Nle                           | = B20                                             | 0                                   | 0                  |                                     |  |   |  |
| Tyr                           | = B21                                             | 3.11                                | 3.066              |                                     |  |   |  |
| Phe                           | = B22                                             | 1.916                               | 1.893              |                                     |  |   |  |
| His                           | = B23                                             | 1.039                               | 1.022              |                                     |  |   |  |
| Lys                           | = B24                                             | 3.971                               | 3.906              |                                     |  |   |  |
| NH4                           | 0                                                 | 7.145                               | 6.989              |                                     |  |   |  |
| Arg                           | = B25                                             | 1.753                               | 1.736              |                                     |  |   |  |
| Total nMoles                  | = SUM(B42:B60)                                    | = SUM(C41:C60)-C59                  | = SUM(D41:D60)-D59 |                                     |  |   |  |
| Total nMoles/Total # residues |                                                   | C61/B61                             | D61/B61            |                                     |  |   |  |



|                                     | E                   | F                   | G                   | H |    |
|-------------------------------------|---------------------|---------------------|---------------------|---|----|
|                                     |                     |                     |                     |   | 1  |
|                                     |                     |                     |                     |   | 2  |
|                                     |                     |                     |                     |   | 3  |
|                                     |                     |                     |                     |   | 4  |
|                                     | Protein X           |                     |                     |   |    |
| 0                                   |                     |                     |                     |   | 5  |
| = (E66)                             | Averages            |                     |                     |   | 6  |
| = E67                               |                     | = AVERAGE(C7:E7)    |                     |   | 7  |
| = E68                               |                     | = AVERAGE(C8:E8)    |                     |   | 8  |
| = E69                               |                     | = AVERAGE(C9:E9)    |                     |   | 9  |
| = E70                               |                     | = AVERAGE(C10:E10)  |                     |   | 10 |
| = E71                               |                     | = AVERAGE(C11:E11)  |                     |   | 11 |
| = E72                               |                     | = AVERAGE(C12:E12)  |                     |   | 12 |
| = E73                               |                     | = AVERAGE(C13:E13)  |                     |   | 13 |
| = E74                               |                     | = AVERAGE(C14:E14)  |                     |   | 14 |
| = E75                               |                     | = AVERAGE(C15:E15)  |                     |   | 15 |
| = E76                               |                     | = AVERAGE(C16:E16)  |                     |   | 16 |
| = E77                               |                     | = AVERAGE(C17:E17)  |                     |   | 17 |
| = E78                               |                     | = AVERAGE(C18:E18)  |                     |   | 18 |
| = E79                               |                     | = AVERAGE(C19:E19)  |                     |   | 19 |
| = E80                               |                     | = AVERAGE(C20:E20)  |                     |   | 20 |
| = E81                               |                     | = AVERAGE(C21:E21)  |                     |   | 21 |
| = E82                               |                     | = AVERAGE(C22:E22)  |                     |   | 22 |
| = E83                               |                     | = AVERAGE(C23:E23)  |                     |   | 23 |
| = E84                               |                     | = AVERAGE(C24:E24)  |                     |   | 24 |
| = E86                               |                     | = AVERAGE(C25:E25)  |                     |   | 25 |
|                                     |                     |                     |                     |   | 26 |
|                                     |                     |                     |                     |   | 27 |
|                                     |                     | 20                  |                     |   | 28 |
|                                     |                     | 150                 |                     |   | 29 |
|                                     |                     | 1                   |                     |   | 30 |
|                                     |                     |                     |                     |   | 31 |
| = H62                               |                     |                     |                     |   | 32 |
|                                     |                     |                     |                     |   | 33 |
| = ([E33*B28*0.001*F30/B32]/F29)*F31 |                     | = AVERAGE(C25:E25)  |                     |   | 34 |
|                                     |                     |                     |                     |   | 35 |
| data file #3                        | Ave nM cal          | Ave nM cal          | Ave nM cal          |   | 40 |
| 0.011                               |                     |                     |                     |   | 41 |
| 5.075                               | = C42/B42           | = D42/B42           | = E42/B42           |   | 42 |
| 5.091                               |                     |                     |                     |   | 43 |
| 7.586                               |                     |                     |                     |   | 44 |
| 5.801                               | = C45/B45           | = D45/B45           | = E45/B45           |   | 45 |
| 4.417                               |                     |                     |                     |   | 46 |
| 4.765                               |                     |                     |                     |   | 47 |
| 4.481                               | = C48/B48           | = D48/B48           | = E48/B48           |   | 48 |
| 1.182                               |                     |                     |                     |   | 49 |
| 4.86                                |                     |                     |                     |   | 50 |
| 0.401                               |                     |                     |                     |   | 51 |
| 1.653                               |                     |                     |                     |   | 52 |
| 4.797                               | = C53/B53           | = D53/B53           | = E53/B53           |   | 53 |
| 0                                   |                     |                     |                     |   | 54 |
| 3.067                               |                     |                     |                     |   | 55 |
| 1.889                               | = C56/B56           | = D56/B56           | = E56/B56           |   | 56 |
| 1.02                                | = C57/B57           | = D57/B57           | = E57/B57           |   | 57 |
| 3.914                               | = C58/B58           | = D58/B58           | = E58/B58           |   | 58 |
| 7.148                               |                     |                     |                     |   | 59 |
| 1.75                                | = C60/B60           | = D60/B60           | = E60/B60           |   | 60 |
| = SUM(E41:E60)-E59                  |                     |                     |                     |   | 61 |
| = E61/B61                           | = AVERAGE (F42:F60) | = AVERAGE (G42:G60) | = AVERAGE (H42:H60) |   | 62 |

(continued)

Table 3 (continued)

| A           | B                       | C                                     | D                                     |
|-------------|-------------------------|---------------------------------------|---------------------------------------|
| Amino Acid  | Theoretical Composition | = C40                                 | = D40                                 |
| CyA         | = B7                    | = IF(C41/[F\$62])=0, " ", [C41/F\$62] | = IF(D41/[G\$62])=0, " ", [D41/G\$62] |
| Asx         | = B8                    | = IF(C42/[F\$62])=0, " ", [C42/F\$62] | = IF(D42/[G\$62])=0, " ", [D42/G\$62] |
| Thr         | = B9                    | = IF(C43/[F\$62])=0, " ", [C43/F\$62] | = IF(D43/[G\$62])=0, " ", [D43/G\$62] |
| Ser         | = B10                   | = IF(C44/[F\$62])=0, " ", [C44/F\$62] | = IF(D44/[G\$62])=0, " ", [D44/G\$62] |
| Glx         | = B11                   | = IF(C45/[F\$62])=0, " ", [C45/F\$62] | = IF(D45/[G\$62])=0, " ", [D45/G\$62] |
| Pro + CySH  | = B12                   | = IF(C46/[F\$62])=0, " ", [C46/F\$62] | = IF(D46/[G\$62])=0, " ", [D46/G\$62] |
| Gly         | = B13                   | = IF(C47/[F\$62])=0, " ", [C47/F\$62] | = IF(D47/[G\$62])=0, " ", [D47/G\$62] |
| Ala         | = B14                   | = IF(C48/[F\$62])=0, " ", [C48/F\$62] | = IF(D48/[G\$62])=0, " ", [D48/G\$62] |
| 1/2 Cys-Cys | = B15                   | = IF(C49/[F\$62])=0, " ", [C49/F\$62] | = IF(D49/[G\$62])=0, " ", [D49/G\$62] |
| Val         | = B16                   | = IF(C50/[F\$62])=0, " ", [C50/F\$62] | = IF(D50/[G\$62])=0, " ", [D50/G\$62] |
| Met         | = B17                   | = IF(C51/[F\$62])=0, " ", [C51/F\$62] | = IF(D51/[G\$62])=0, " ", [D51/G\$62] |
| Ile         | = B18                   | = IF(C52/[F\$62])=0, " ", [C52/F\$62] | = IF(D52/[G\$62])=0, " ", [D52/G\$62] |
| Leu         | = B19                   | = IF(C53/[F\$62])=0, " ", [C53/F\$62] | = IF(D53/[G\$62])=0, " ", [D53/G\$62] |
| Nle         | = B20                   | = IF(C54/[F\$62])=0, " ", [C54/F\$62] | = IF(D54/[G\$62])=0, " ", [D54/G\$62] |
| Tyr         | = B21                   | = IF(C55/[F\$62])=0, " ", [C55/F\$62] | = IF(D55/[G\$62])=0, " ", [D55/G\$62] |
| Phe         | = B22                   | = IF(C56/[F\$62])=0, " ", [C56/F\$62] | = IF(D56/[G\$62])=0, " ", [D56/G\$62] |
| His         | = B23                   | = IF(C57/[F\$62])=0, " ", [C57/F\$62] | = IF(D57/[G\$62])=0, " ", [D57/G\$62] |
| Lys         | = B24                   | = IF(C58/[F\$62])=0, " ", [C58/F\$62] | = IF(D58/[G\$62])=0, " ", [D58/G\$62] |
| NH4         | 0                       | = IF(C59/[F\$62])=0, " ", [C59/F\$62] | = IF(D59/[G\$62])=0, " ", [D59/G\$62] |
| Arg         | = B25                   | = IF(C60/[F\$62])=0, " ", [C60/F\$62] | = IF(D60/[G\$62])=0, " ", [D60/G\$62] |

2. After removal of the acid, reconstitute the samples with Li-S buffer, then analyze using lithium citrate buffers with a lithium-exchange column.
3. Use the analysis conditions given in **Table 4**. If needed, the separation between Nle and Tyr (which elutes after Nle) can be increased by lowering the column temperature.
4. Set the detector to the most sensitive scale (0.1 AUFS) (*see Note 17*). A chromatogram is given in **Fig. 3**.

### 3.4. Hydroxylysine Analysis

1. Hydrolyze samples containing 50–100 µg of protein (*see Note 18*) for 24 h as described in **Subheading 3.1**.
2. Remove the acid, then reconstitute the samples with Li-S buffer, and analyze using the modified program given in **Table 5** (*see Note 19*). The standard chromatogram is given in **Fig. 4** (*see Note 20*).

### 3.5. Carboxypeptidase Analysis

Applications involving single or combinations of carboxypeptidases to assign C-terminal protein sequences have been adequately described elsewhere (**18**).

1. Add norleucine to samples prior to the addition of carboxypeptidases at equimolar ratios (e.g., 10 nmol Nle for a sample containing 10 nmol of polypeptide).
2. Take aliquots at various time-points and place in Eppendorf tubes containing an equal volume of 2 N glacial acetic acid.
3. Heat for 2 min at 100°C on a boiling water bath to halt the digestion and precipitate the protein.

| E                                   |    |
|-------------------------------------|----|
| = E40                               | 66 |
| = IF(E41/[HS62]=0, " ", [E41/HS62]) | 67 |
| = IF(E42/[HS62]=0, " ", [E42/HS62]) | 68 |
| = IF(E43/[HS62]=0, " ", [E43/HS62]) | 69 |
| = IF(E44/[HS62]=0, " ", [E44/HS62]) | 70 |
| = IF(E45/[HS62]=0, " ", [E45/HS62]) | 71 |
| = IF(E46/[HS62]=0, " ", [E46/HS62]) | 72 |
| = IF(E47/[HS62]=0, " ", [E47/HS62]) | 73 |
| = IF(E48/[HS62]=0, " ", [E48/HS62]) | 74 |
| = IF(E49/[HS62]=0, " ", [E49/HS62]) | 75 |
| = IF(E50/[HS62]=0, " ", [E50/HS62]) | 76 |
| = IF(E51/[HS62]=0, " ", [E51/HS62]) | 77 |
| = IF(E52/[HS62]=0, " ", [E52/HS62]) | 78 |
| = IF(E53/[HS62]=0, " ", [E53/HS62]) | 79 |
| = IF(E54/[HS62]=0, " ", [E54/HS62]) | 80 |
| = IF(E55/[HS62]=0, " ", [E55/HS62]) | 81 |
| = IF(E56/[HS62]=0, " ", [E56/HS62]) | 82 |
| = IF(E57/[HS62]=0, " ", [E57/HS62]) | 83 |
| = IF(E58/[HS62]=0, " ", [E58/HS62]) | 84 |
| = IF(E59/[HS62]=0, " ", [E59/HS62]) | 85 |
| = IF(E60/[HS62]=0, " ", [E60/HS62]) | 86 |

4. After cooling on wet ice, centrifuge the samples and transfer the supernatant to another Eppendorf tube.
5. Dry by rotary evaporation using a Savant SpeedVac.
6. Reconstitute the samples with Li-S buffer.
7. Follow the operating conditions given in **Table 6**. The initial 40-min segment of the chromatogram obtained using this modified program is shown in **Fig. 5**.

### 3.6. Amino Sugar Analysis (see Note 21)

1. Divide samples containing 2–20  $\mu\text{g}$  of protein or 0.5–5 nmol of peptide fractions into two identical aliquots.
2. Hydrolyze one aliquot for 24 h at 110°C as described in **Subheading 3.1**.
3. Hydrolyze the other aliquot for only 2 h at 110°C (*see Note 22*).
4. After removal of the acid, reconstitute the 2-h hydrolysates with Na-S buffer and analyze using a modified program given in **Table 7** (*see Note 23*).
5. Analyze the 24-h hydrolysates using the standard method (**Table 1**) (**Fig. 6**) for quantitation of the protein/peptide to permit molar GlcNAc and/or GalNAc determinations. Tryptophan standards should also be analyzed to ensure that Trp does not coelute with GalNH<sub>2</sub>; if necessary, this resolution can be improved by lowering the column temperature (*see Note 24*).

### 3.7. Cysteine Alkylation Monitoring

1. Hydrolyze desalted samples containing 2–20  $\mu\text{g}$  of S-carboxy-methylated or S-carboxyamidomethylated proteins for 24 h as described in **Subheading 3.1**.
2. After removal of the acid, reconstitute the samples with Na-S buffer, and analyze using the standard program (**Table 1**) (*see Note 25*). Representative chromato-

| AAA<br>Macro            |                             | 3            |              |              |           |
|-------------------------|-----------------------------|--------------|--------------|--------------|-----------|
| Protein Template        |                             |              |              |              |           |
| Name:                   | Researcher                  |              | Protein Name |              | Protein X |
| Extension:              | 0                           |              | Mail Stop:   |              | 0         |
| Amino Acid              | Theoretical Composition     | data file #1 | data file #2 | data file #3 | Averages  |
| CyA                     | 0                           | 0.1          | 0.1          | 0.1          | 0.1       |
| Asx                     | 35                          | 34.7         | 34.8         | 34.8         | 34.7      |
| Thr                     | 37                          | 34.8         | 34.9         | 34.9         | 34.9      |
| Ser                     | 59                          | 52.0         | 52.0         | 52.0         | 52.0      |
| Glx                     | 41                          | 39.7         | 39.8         | 39.7         | 39.7      |
| Pro + Cys SH            | 24                          | 29.9         | 29.0         | 30.3         | 29.7      |
| Gly                     | 33                          | 32.7         | 32.6         | 32.6         | 32.7      |
| Ala                     | 30                          | 30.7         | 30.7         | 30.7         | 30.7      |
| 1/2 Cys-Cys             | 10                          | 8.7          | 9.4          | 8.1          | 8.7       |
| Val                     | 35                          | 33.3         | 33.3         | 33.3         | 33.3      |
| Met                     | 3                           | 2.6          | 2.5          | 2.7          | 2.6       |
| Ile                     | 12                          | 11.4         | 11.4         | 11.3         | 11.4      |
| Leu                     | 32                          | 33.0         | 33.0         | 32.9         | 33.0      |
| Nle                     | 0                           |              |              |              |           |
| Tyr                     | 22                          | 21.0         | 21.0         | 21.0         | 21.0      |
| Phe                     | 13                          | 13.0         | 13.0         | 12.9         | 13.0      |
| His                     | 7                           | 7.0          | 7.0          | 7.0          | 7.0       |
| Lys                     | 27                          | 26.9         | 26.8         | 26.8         | 26.8      |
| Arg                     | 12                          | 11.9         | 11.9         | 12.0         | 11.9      |
|                         |                             |              |              |              |           |
|                         |                             |              |              |              |           |
| Molecular Mass (g/mole) | 47503.01                    |              |              |              |           |
|                         | ul in Ampoule               |              |              |              |           |
|                         | smpl recon(ul)              |              |              |              |           |
|                         | dilution factor             |              |              |              |           |
| column load (ul)        | 50                          |              |              |              |           |
|                         | nMols Protein               | 0.148        | 0.146        | 0.146        |           |
|                         | Protein Concentration mg/mL | 1.054        | 1.039        | 1.040        | 1.044     |

Fig. 2. Summary sheet using the AAA macro. Average compositions are given in the right-hand column, and the average mg/mL value is provided in the lower right hand box. CyA refers to cysteic acid, which is present when samples are oxidized intentionally.

grams for the standard mixture containing carboxymethylcysteine (CMCys) and for an *S*-carboxymethylated recombinant antibody sample are given in **Fig. 7A,B**, respectively.

3. Monitor CMCys and half-cystine residue/mol values to determine the extent of cysteine alkylation (*see* **Notes 26** and **27**).

**Table 4**  
**Norleucine Analysis**

| Time (min) | Event                   | Conditions         |
|------------|-------------------------|--------------------|
| 0.0        | Sample injection        | Li-A buffer, 38°C  |
| 44.0       | Buffer change           | Li-A to Li-D       |
| 73.0       | Buffer change           | Li-D to Li-R       |
| 74.0       | Reagent pump            | Ninhydrin to water |
| 77.0       | Buffer change           | Li-R to Li-A       |
| 80.0       | Reagent pump            | Water to ninhydrin |
| 90.0       | Recyle (start next run) |                    |

Buffer pump: 20 mL/h.

Reagent pump: 10 mL/h.

#### 4. Notes

1. Precolumn derivatization with RP-HPLC separation is used for amino acid analysis; a popular version is the Waters AccQTag system (21). These precolumn methods may not be suitable for detection of trace levels of minor amino acids (the needle-in-a-haystack problem) because the peak resolutions are diminished when the sample loads are increased, whereas resolution is maintained with higher loads using the cation-exchange systems. In addition, precolumn accuracy may be limited if derivatization is incomplete, a problem that does not occur with postcolumn derivatization systems. Similarly, cation-exchange systems are more tolerant of salts and residual HCl than the precolumn systems.
2. Hydrolysis ampoules are wrapped in heavy duty foil and pyrolyzed by heating for 24 h at 400°C in a muffle furnace before use.
3. Production of the Beckman 6300 analyzers described in this chapter has been halted, but a similar system can be fashioned using components offered by Pickering Labs (Mountain View, CA) (19). Pickering also supplies amino acid analysis buffers, reagents, and columns for the Beckman 6300, but care must be taken not to combine Pickering's Trione ninhydrin reagent with mobile phases that contain alcohols (such as Beckman's Na-A and Na-B) as this combination may clog the analyzer's reactor.
4. Hitachi also offers a cation-exchange amino acid analysis instrument.
5. Dionex has recently introduced an anion-exchange system that detects underivatized amino acids using pulsed amperometric detection (20) (see also Chapter 7, this volume), but we have no experience with this system.
6. A wash bottle containing 1 M sodium bicarbonate is kept nearby wherever HCl ampoules are opened to neutralize spills.
7. The prepared standards should be stored refrigerated in aliquots using screw-top Eppendorf tubes equipped with a rubber gasket to prevent evaporation. Tryptophan tends to degrade over time in acid conditions, so fresh Trp standards should be prepared when needed; commercial preparations containing Trp may not be reliable.

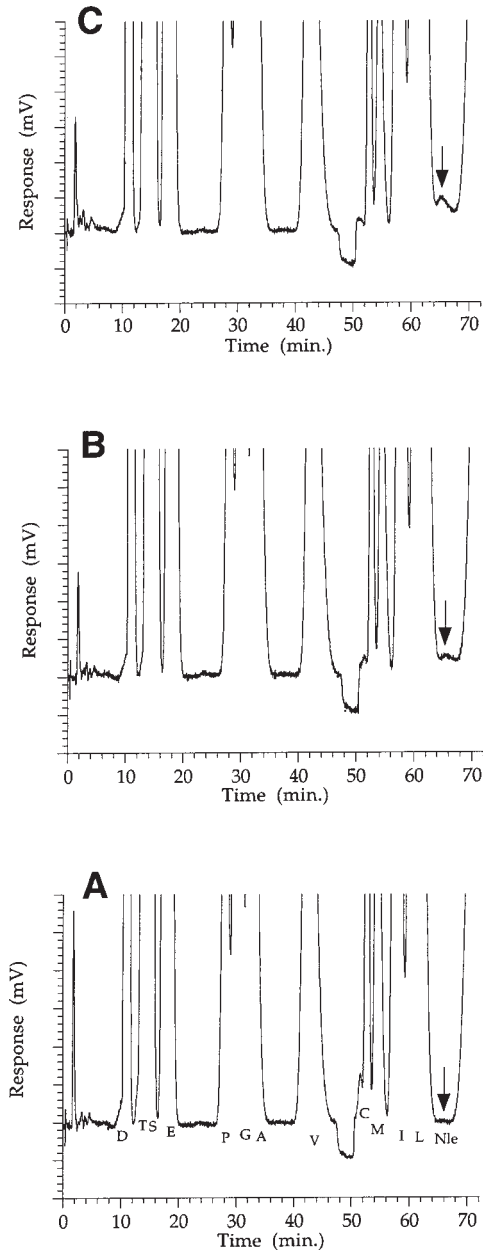


Fig. 3. Analysis for norleucine incorporation at Met positions. Aliquots from 40  $\mu$ g of a recombinant protein are given, with the arrow indicating the Nle peak after additions of (A) 0 pmol Nle (B) 200 pmol Nle, or (C) 400 pmol Nle. Operating parameters are given in **Table 4**.

**Table 5**  
**Hydroxylysine Analysis**

| Time (min) | Event                    | Conditions              |
|------------|--------------------------|-------------------------|
| 0.0        | Sample injection         | Li-A buffer, 38°C       |
| 12.0       | Temperature change       | 38°C to 50°C over 8 min |
| 42.0       | Buffer change            | Li-A to Li-B            |
| 60.0       | Temperature change       | 50°C to 71°C over 8 min |
| 70.0       | Buffer change            | Li-B to Li-C            |
| 110.0      | Reagent pump             | Ninhydrin to water      |
| 111.0      | Buffer change            | Li-C to Li-R            |
| 114.0      | Buffer change            | Li-R to Li-A            |
| 125.0      | Reagent pump             | water to ninhydrin      |
| 135.0      | Recycle (start next run) |                         |

Buffer pump: 20 mL/h.

Reagent pump: 10 mL/h.

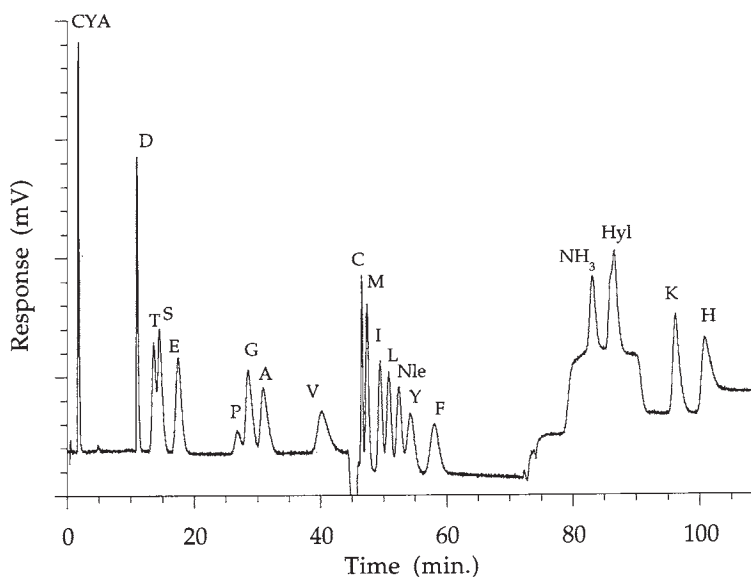


Fig. 4. Analysis for hydroxylysine. A standard mixture containing 1 nmol of each component was loaded. Hyl appears as a poorly-resolved peak pair. Operating parameters are given in **Table 5**.

- Protein/peptide quantitation can be compromised by multiple sample transfers. When accuracy is essential, samples should be transferred directly from the primary container to the hydrolysis ampoule.

**Table 6**  
**Analysis of Carboxypeptidase Supernatants**

| Time (min) | Event                   | Conditions              |
|------------|-------------------------|-------------------------|
| 0.0        | Sample injection        | Li-A buffer, 38°C       |
| 12.0       | Temperature change      | 38°C to 50°C over 8 min |
| 43.0       | Buffer change           | Li-A to Li-B            |
| 60.0       | Temperature change      | 50°C to 73°C over 8 min |
| 60.0       | Buffer change           | Li-B to Li-C            |
| 130.0      | Reagent pump            | Ninhydrin to water      |
| 132.0      | Buffer change           | Li-C to Li-R            |
| 134.0      | Buffer change           | Li-R to Li-A            |
| 140.0      | Temperature change      | 73°C to 38°C            |
| 140.0      | Reagent pump            | Water to ninhydrin      |
| 155.0      | Recyle (start next run) |                         |

Buffer pump: 20 mL/h.

Reagent pump: 10 mL/h.

**Table 7**  
**Amino Sugar Analysis**

| Time (min) | Event                   | Conditions         |
|------------|-------------------------|--------------------|
| 0.0        | Sample injection        | Na-F buffer, 66°C  |
| 55.0       | Reagent pump            | Ninhydrin to water |
| 57.0       | Buffer change           | Na-F to Na-R       |
| 58.5       | Buffer change           | Na-R to Na-F       |
| 60.0       | Reagent pump            | Water to ninhydrin |
| 74.0       | Recyle (start next run) |                    |

Buffer pump: 20 mL/h.

Reagent pump: 10 mL/h.

9. Alternative hydrolysis systems have been proposed, including the Waters PicoTag batch hydrolysis system, in which the 6 *N* HCl is placed outside the sample tubes in a chamber that can be evacuated, closed, and heated. Phenol must also be added to prevent destruction of tyrosine. This system has the advantage that direct contact with the acid is avoided, eliminating a potential source of contamination, but in our experience the poor hydrolysis of Ile-Ile, Ile-Val, and Val-Val bonds with vapor-phase hydrolysis makes this technique unsuitable.
10. Hydrolysis at 155°C for 60 min has also been proposed, but we seldom use this procedure because Thr and Ser values are greatly reduced. Also, because we typically batch samples together, a 24-h hydrolysis is often more convenient from an operational standpoint.
11. A Perkin Elmer Model 200 autosampler has replaced the original coil system, with a fixed 50- $\mu$ L volume used for standards and samples.



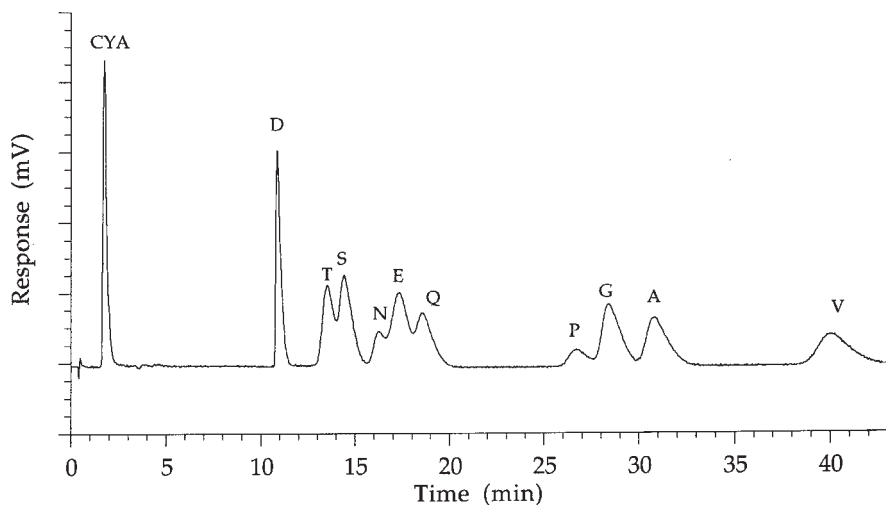


Fig. 5. Analysis of carboxypeptidase digestion samples (expanded view). A standard mixture containing 1 nmol of each component was loaded. For clarity, only the early region of the chromatogram is provided to show the elution positions of Asn and Gln; the complete chromatogram is essentially the same as **Fig. 8**. Operating parameters are given in **Table 6**.

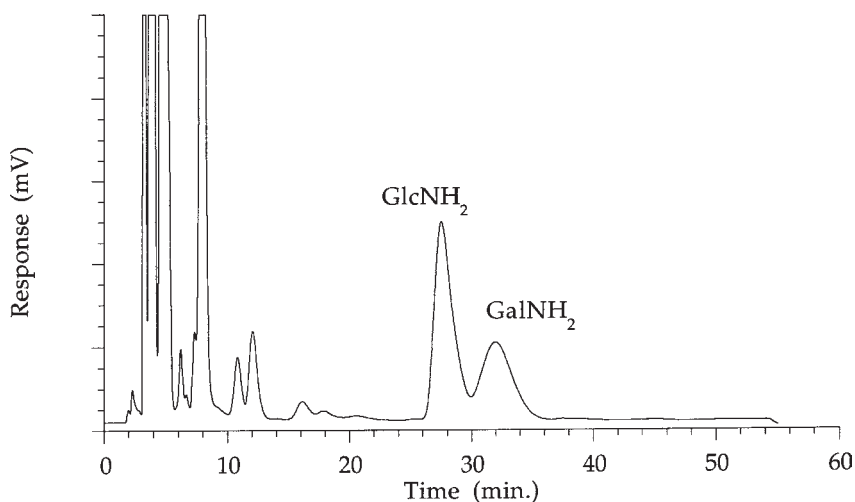


Fig. 6. Amino sugar analysis. A hydrolysate from 1 nmol of a recombinant glycoprotein was loaded. Operating parameters are given in **Table 7**.

12. In the tables, the “reagent pump” event refers to changing the solution added postcolumn from a ninhydrin-containing reagent to water (or vice versa); this is done to avoid having NaOH (Na-R) or LiOH (Li-R) mix with the ninhydrin reagent.

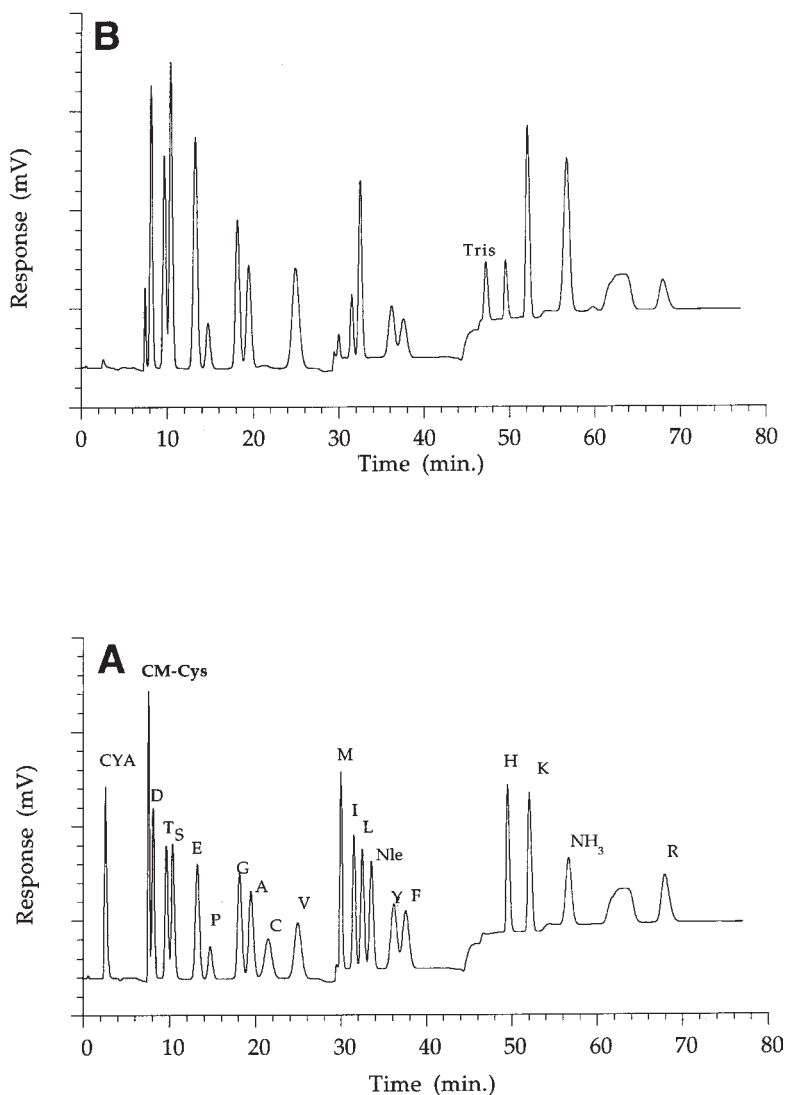


Fig. 7. Carboxymethylcysteine analysis. (A) Standard mixture containing 2 nmol of each component. (B) Analysis of 5  $\mu$ g of a recombinant antibody after reduction and S-carboxymethylation. Operating parameters are given in **Table 1**. The peak eluting at approx 47 min is Tris buffer.

- Key amino acids that typically provide quantitative recoveries (e.g., Asx, Glx, Ala, Leu, Phe, His, Lys, Arg) are used to determine the total nmol of protein or peptide (in the example provided for data file #1 in **Table 3**, 5.132 nmol of Asx are divided by 35 residues of Asx expected per mol of protein to produce a nmol

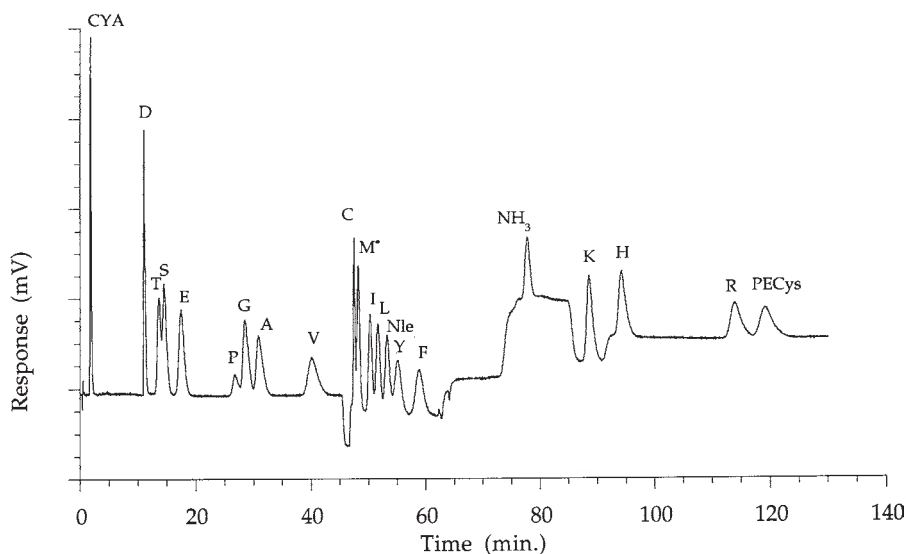


Fig. 8. Pyridylethylcysteine analysis. Analysis of a standard mixture containing 1 nmol of each component. Operating parameters are given in **Table 6**.

protein value); use of these selected amino acids avoids the low recoveries experienced for the acid-labile amino acids, especially Trp, Cys, Thr, and Ser (*I*).

14. The amino acid nmol values are divided by the total protein/peptide nmol value to produce residues/mol values that are averaged in the right-hand column (in the example provided in **Table 3** and **Fig. 2**, the nmol values in rows 42–60 of columns C, D, and E are divided by the average protein value of 0.148, 0.146, and 0.146, respectively, then averaged). The nmol protein/peptide values in **Fig. 2** are also multiplied by the molecular mass to provide the total  $\mu\text{g}$  injected on the analyzer, which is then converted to a  $\mu\text{g}/\mu\text{L}$  value (same as a  $\text{mg}/\text{mL}$  value) by dividing by the volume loaded in the ampoule (e.g., 20  $\mu\text{L}$  in **Fig. 2**), then correcting for the reconstitution volume and  $\mu\text{L}$  injected (e.g., multiplied by 150/50 in **Fig. 2**); these concentrations are then averaged as shown in the lower right-hand box of **Fig. 2**.
15. When the protein/peptide composition is not known, or if the sample contains a mixture of proteins, then quantitation is performed by summing the nanograms contributed by each amino acid residue. For example, instead of calibrating to a 2 nmol standard value for alanine, the data system calibrates to 142 ng (equivalent to 2 nmol of alanine using the residue mass). The values for all amino acids are summed. This method generates values that are usually 5–10% lower than their true quantity because of poor recoveries of acid-labile amino acids, but in our experience these values are likely to be more sensitive and reliable than colorimetric methods.

16. Norleucine quantitation is difficult at levels below 1% Nle-for-Met replacement. Nle added at several levels to samples can establish the lowest level of quantitation, which for us is typically about 75 pmol. In our experience, the incorporation of Nle occurs proportionately at every Met position; therefore, the mol Nle per mol protein value can be divided by the number of methionines to provide percent Nle-for-Met replacement values. Nle replacement can sometimes be observed by electrospray mass spectrometry of intact proteins (18 Dalton lower mass), provided that no other sources of heterogeneity are present (V. Ling, unpublished data).
17. Most amino acids will be present in great abundance, saturating the detector, but the glycine and proline peaks are usually still on-scale, and thus can be used for mol Nle per mol protein quantitation. Chromatograms for 40  $\mu$ g hydrolysates of a recombinant protein spiked with 0, 200, or 400 pmol Nle are given in **Fig. 3A, B, and C**, respectively.
18. Sensitive detection of hydroxylysine (Hyl) is difficult because of the fact that most noncollagenous molecules are at most partially modified at just one -Lys-Gly- position, thus the overall percentage of modified Lys is very low.
19. In the standard amino acid analysis method (**Table 1**), Hyl coelutes with histidine, so it would not be observed in an intact protein that contains His, and it might be misinterpreted in a peptide fraction. The long delay for the second buffer change is needed to increase the resolution of Hyl from ammonia. Hyl appears as a partially resolved doublet peak due to racemization of the  $\delta$  carbon during hydrolysis; therefore, the peak areas are summed.
20. Once it has been determined that Hyl is present, peptide maps may be used to assign the site provided the investigator is aware that Hyl-Gly bonds are fairly resistant to trypsin and endoproteinase Lys-C digestion (**22**).
21. Amino sugar analysis does not provide complete monosaccharide determinations of the types obtained by techniques such as HPAEC-PAD or GC-MS, but it does have the advantages that no additional equipment is required, and the results are routinely quantitative. This approach is most useful when assaying proteins for the mucin-type O-linked oligosaccharides that contain GalNAc at their reducing termini. O-linked structures that lack GalNAc are rare, but have been found in EGF-like domains of several glycoproteins (**23**). Some N-linked structures contain GalNAc, particularly in proteins from human embryonic kidney (293) cells (**24**) or melanoma cells (**25**), but these N-linked structures can be released using PNGaseF to allow discrimination between N-linked and O-linked GalNAc residues.
22. After 2 h of hydrolysis, GlcNAc is hydrolyzed to GlcNH<sub>2</sub>, whereas GalNAc is hydrolyzed to GalNH<sub>2</sub>, and both are released quantitatively.
23. This program starts with the second buffer used in the standard analysis, so most amino acids elute near the beginning of the chromatogram. A chromatogram from 1 nmol of a hydrolysate of a recombinant glycoprotein containing both N-linked and O-linked sites is given in **Fig. 6**.

24. Proteins that are highly glycosylated will have some residual amino sugars that will appear as a broad peak that elutes in the Ile-Leu-Nle region of the standard chromatogram. Increasing the hydrolysis time to 72 h will eliminate this peak.
25. Methionine residues can also be unintentionally S-alkylated, but this can be detected by the presence of trace levels of homoserine, a hydrolysis product of S-carboxymethylmethionine that elutes between Ser and Glx.
26. Samples that have been alkylated using 4-vinylpyridine need to be analyzed using the lithium citrate program that is used for the carboxypeptidase digestion samples (**Table 6**). Pyridylethylcysteine (PECys) is very basic, and elutes after Arg (**Fig. 8**).
27. When monitoring Cys alkylation conditions, attention should be paid to methionine recoveries, as the conditions (such as trace metals or residual O<sub>2</sub>) that affect Met recoveries will also affect CMCys recoveries. In addition, methionine sulfoxide can coelute with CMCys; therefore, samples should be analyzed promptly after acid removal.

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## Purification of Proteins Using UltraMacro Spin Columns or ProSorb Sample Preparation Cartridges for Amino Acid Analysis

Li Zhang and Nancy Denslow

### 1. Introduction

Amino acid analysis of proteins is one of the best and most accurate methodologies to quantify proteins. Ideally, samples should be pure prior to hydrolysis, however, they are often not only at low concentration, but also in buffers, salts, and/or detergents. Sample contaminants contribute to background noise that hampers amino acid analysis, variably affecting recoveries of individual amino acids (**1**). The necessity to obtain samples in high concentration and free of contaminants is a problem for protein quantitation and structural characterization (*see Note 1*). Many of the existing methods for removing buffers and salts from proteins, such as precipitation methods or reverse-phase high-performance liquid chromatography (RP-HPLC) result in variable and unacceptably high losses of protein (*see Note 2*), making an assessment of the true concentration difficult. In this chapter, we describe two methods commonly used to recover proteins from buffers: a gel filtration method, Ultra microspin columns, and adsorption to a polyvinylidene difluoride (PVDF) membrane in the ProSorb sample preparation cartridge from Applied Biosystems. Both methods work well, with somewhat better recoveries for the gel filtration method.

### 2. Materials

#### 2.1. Equipment

1. UltraMicro spin (gel filtration) columns (Amika Corp, Columbia, MD) are available in two sizes (*see Note 3*):

|                       | Sample vol     | Bed vol     |
|-----------------------|----------------|-------------|
| Ultramicrospin column | 5–25 mL        | 100 $\mu$ L |
| Macrospin column      | 50–150 $\mu$ L | 500 $\mu$ L |

2. ProSorb sample preparation cartridge (Applied Biosystems, Foster City, CA).
3. Immobilon-PSQ transfer membrane (Millipore Corp., Bedford, MA).
4. Sample hydrolysis tubes (Corning Inc., Corning, NY) (*see Note 4*).
5. Reaction Vials (Waters Associates, Milford, MA).

## 2.2. Reagents

1. HPLC-grade water (*see Note 5*).
2. Trifluoroacetic acid (TFA).
3. Methanol.
4. Hydrochloric acid (HCl).
5. Phenol.

## 2.3. Reagents

1. 20% methanol, 0.1 *N* HCl.
2. 6 *N* HCl + 0.1% phenol.
3. 0.1% TFA.

## 3. Methods

### 3.1. Preparation of the Macrospin column (sample 50–150 $\mu$ L)

1. The macrospin columns should be prepared basically as suggested by the manufacturer (2), except that we have introduced several more washes to better equilibrate the columns. In brief, tap the column gently to recover the gel material at the bottom of the column. Hydrate the gel with 500  $\mu$ L of HPLC-grade water for 15 min at room temperature (*see Note 6*) and then centrifuge the column for 4 min at about 2000*g* to equilibrate the column.
2. Wash the column 3 times with 500  $\mu$ L of water, to ensure full equilibration.
3. Blot the exterior of the column dry to remove all moisture and place it in a new collecting tube. For the macrospin columns, add 50–150  $\mu$ L of sample to the top, being careful to place the sample in the center of the column.
4. Centrifuge the tube for 4 min at 2000*g*, and wash the column with at least 200  $\mu$ L of water (*see Note 7*).
5. Remove the purified sample and speed-vacuum to dryness before hydrolysis (*see Note 8*).

### 3.2. Ultra Microspin Columns (sample 5–25 $\mu$ L)

1. As for the macrocolumns, tap the dry gel to the bottom of the column. Wet with 50  $\mu$ L of water and centrifuge it for 3 min, at 1000*g* to equilibrate the column. For best results, wash the column three times with 200  $\mu$ L of water. To centrifuge these columns, place them directly into a microcentrifuge or use an empty 1.7-mL microfuge tube as an adapter.



**Table 1**  
**Maximum Detergent Concentrations<sup>a</sup>**

| Detergent              | Maximum concentration (%)<br>without interference to binding<br>ProSorb PVDF membranes (v/v or w/v) |
|------------------------|-----------------------------------------------------------------------------------------------------|
| Triton X-100 (reduced) | 0.01                                                                                                |
| Tween-20               | 0.01                                                                                                |
| SDS                    | 0.02                                                                                                |
| Octyl glucoside        | 0.25                                                                                                |
| Brij 35                | 0.02                                                                                                |

<sup>a</sup>Data from PE Biosystems (1996) (7).

2. After the column has been equilibrated, blot dry to remove any moisture that may be adhering to the exterior of the column.
3. Add the 5–25  $\mu\text{L}$  sample to the top of the spin column, being careful to place the sample in the center of the column.
4. Place the column in a new collecting tube, spin the tube for 3 min at 1000g. After centrifugation, the purified sample will be in the collecting tube and will be ready for further use.

### 3.3. Preparation of the ProSorb Sample Cartridge

1. Make sure sample is in 0.1% TFA for optimum results (*see Note 9*). If the sample volume is less than 100  $\mu\text{L}$ , dilute sample to 100  $\mu\text{L}$  with 0.1% TFA. If the sample has detergent, dilute such that the detergent concentration is below the values shown in **Table 1** (*see Notes 10 and 11*).
2. Place ProSorb device in an appropriate test tube rack.
3. Wet the PVDF membrane in the reservoir with 10  $\mu\text{L}$  of methanol (*see Note 12*).
4. Add sample to reservoir (100 to 400  $\mu\text{L}$  volume), and allow the sample to pass through the membrane (*see Note 13 and 14*).
5. Ensure that the sample reservoir has intimate contact with the absorbent and fluid transfer begins.
6. Add additional sample aliquots, if necessary (change absorbent after each 750  $\mu\text{L}$ ).
7. Wash the membrane with 100  $\mu\text{L}$  of 0.1% TFA two times.
8. Remove sample reservoir and air-dry the PVDF membrane.
9. Punch out PVDF membrane, then continue with the next processing steps.

### 3.4. Hydrolysis of Sample in Solution

There are several methods that can be used to hydrolyze the sample including vapor phase hydrolysis (3), hydrolysis in the tube (4), and microwave hydrolysis (5). All work about equally well. A vapor phase method that works very well is described below.

1. Place 6 × 50 mm sample tubes in a bulge modified 40-mL screw-cap vial containing 200 μL of 6 N HCl plus 1% phenol. Seal the vial with a modified minaret slide valve.
2. Evacuate briefly (20 s) then flush with nitrogen for 5 s. Repeat this alternating process three times. Close the slide valve on the fourth vacuum step. Heat 24 h at 120°C.
3. Add sample dilution buffer and transfer the dilution to a spin filter tube, centrifuge at 2000g for 4 min and store at 4°C until ready for AAA.

### 3.5. Hydrolysis for PVDF Membrane

There have been several methods described to improve the recovery of a sample hydrolyzed on a PVDF membrane (**4**) (*see Note 15*). We routinely use Immobilon-PSQ transfer membrane (Millipore Corp.).

1. Push the PVDF membrane into the bottom of a hydrolysis tube.
2. Hydrolyze as normal. After the acid has been dried from the hydrolyzed sample, add 10 μg of methanol. Extract two times with 50 μL of 20% MeOH, 0.1 N HCl, and transfer to a labeled 0.5-mL Eppendorf tube.
3. Put the Eppendorf tube into a dry PICOTAG vessel and dry it down.
4. Dilute with sample dilution buffer and filter using a spin column. Sample is ready for AAA.

### 3.6. Amino Acid Analysis

Amino acid analysis can be performed either by precolumn derivatization with phenylisothiocyanate on the Applied Biosystem 420 or by postcolumn derivation with Ninhydrin after ion-exchange separation on the Beckman 6300 high performance analyzer.

## 4. Notes

1. A common source of amino acid contamination is fingerprints. Be sure to wear gloves to keep this type of contamination away from the samples. Another source of contamination is dust.
2. Every time a sample is handled, a portion is lost. Minimize sample handling as much as possible.
3. It is important to choose the appropriate size of microspin column for your sample volume. For sample volumes of 5–25 μL, the smaller ultramicrospin columns are most appropriate. They have a packed bed volume of about 100 μL. Larger volumes, 50–150 μL, should be applied to the larger macrospin columns, which have a packed-bed volume of 500 μL.
4. All glassware used for hydrolysis should be thoroughly cleaned and pyrolyzed at 500°C.
5. Always use the highest quality of reagents (analytical grade) and water. We recommend MilliQ water.

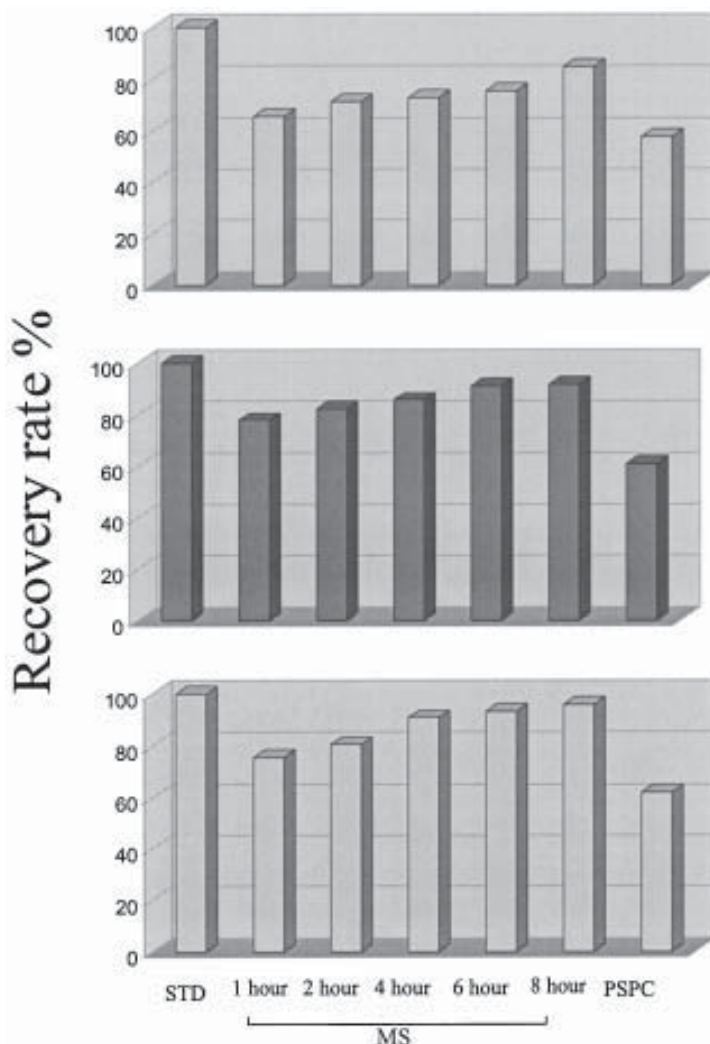


Fig. 1. Comparison of protein recoveries of desalted samples. CAII, TIM, or BSA were prepared in either H<sub>2</sub>O (std) or in a buffer containing 50 mM Tris, pH 8.0, and 0.2% SDS and desalted through either a Macrospin Column (MSC) or a ProSorb Sample Preparation Cartridge (PSPC) as indicated. Samples were hydrolyzed and analyzed on a Beckman 6300 Amino Acid Analyzer. Each bar represents average recovery from three replicates.

6. The percentage of sample recovery increases with increased hydration time for the spin column. When columns are hydrated for 4 h or longer, it is feasible to recover more than 90% of the sample (**Fig. 1**).

7. It is wise to monitor the recovery of the protein by absorbance of the column eluent at 280 nm. Collect sample until all of the material is eluted.
8. When samples containing a high concentration of protein or peptide are available in reasonable volume, there is no need to purify the sample further, simply dilute the sample with HPLC-grade water to a salt concentration that will not interfere with the analysis.
9. Acetonitrile (ACN) concentrations greater than 15% interfere with sample binding to PVDF. We recommend, therefore, that HPLC fractions be diluted with distilled water or 0.1% TFA to reduce the concentration of ACN to less than 15%, prior to using the ProSorb cartridge.
10. Often proteins submitted for amino acid analysis contain detergents used in their preparation. Detergents can dramatically affect the binding of proteins to PVDF membranes. **Table 1** summarizes the concentrations of detergents that are tolerable in a sample prior to binding to the membrane. Samples containing higher than these concentrations should be diluted to reduce the detergent and then added to the membrane in batches.
11. Many samples contain significantly more detergent than the amounts listed in **Table 1**. One way to deal with higher amounts of detergent is to add a small amount of methanol to the sample before loading it into the ProSorb cartridge. This will weaken sample interaction with the detergent.
12. ProSorb cartridges should be prepared as directed by the manufacturer, using MilliQ water for all wash steps to reduce persistent background levels of contaminating amino acids.
13. Be careful with the manipulation of samples at every step. Be aware that proteins and peptides can irreversibly bind to glass and/or plastic tubes. If your sample is small, you can minimize losses in the cartridge by adding 100  $\mu\text{L}$  of buffer to the prewetted membrane prior to the addition of sample. The sample can be added directly to this buffer.
14. Samples or washes larger than 400  $\mu\text{L}$  are accommodated by loading multiple aliquots into the reservoir. The PVDF membrane should not be allowed to dry between aliquots.
15. Based on a recent ABRF study on amino acid analysis (8), samples bound to PVDF membranes tend to have higher error rates (20%) than those analyzed free in solution (12%). These error rates are averages for the study.

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## Amino Acid Analysis by High-Performance Liquid Chromatography after Derivatization with 1-Fluoro-2,4-dinitrophenyl-5-L-alanine Amide (Marfey's Reagent)

Sunil Kochhar, Barbara Mouratou, and Philipp Christen

### 1. Introduction

Precolumn derivatization of amino acids followed by their resolution by reverse-phase high-performance liquid chromatography (RP-HPLC) is now the preferred method for quantitative amino acid analysis. The derivatization step introduces covalently bound chromophores necessary not only for interactions with the apolar stationary phase for high resolution, but also for photometric or fluorometric detection.

Marfey's reagent, 1-fluoro-2,4-dinitrophenyl-5-L-alanine amide or (S)-2-[(5-fluoro-2,4-dinitrophenyl)-amino]propanamide, can be used to separate and to determine enantiomeric amino acids (*I*). The reagent reacts stoichiometrically with the amino group of enantiomeric amino acids to produce stable diastereomeric derivatives that can readily be separated by RP-HPLC (**Fig. 1**). The dinitrophenyl alanine amide moiety strongly absorbs at 340 nm ( $\epsilon = 30,000 \text{ M}^{-1} \text{ cm}^{-1}$ ), allowing detection in the subnmol range.

Precolumn derivatization with the reagent is also used to quantify the 19 commonly analyzed L-amino acids (*2*). Major advantages of Marfey's reagent over other precolumn derivatizations are: (1) possibility to carry out chromatography on any multipurpose HPLC instrument without column heating; (2) detection at 340 nm is insensitive to most solvent impurities; (3) simultaneous detection of proline in a single chromatographic run; and (4) stable amino acid derivatives. For the occasional user, the simple methodology provides an attractive and inexpensive alternative to the dedicated amino acid analyzer. The

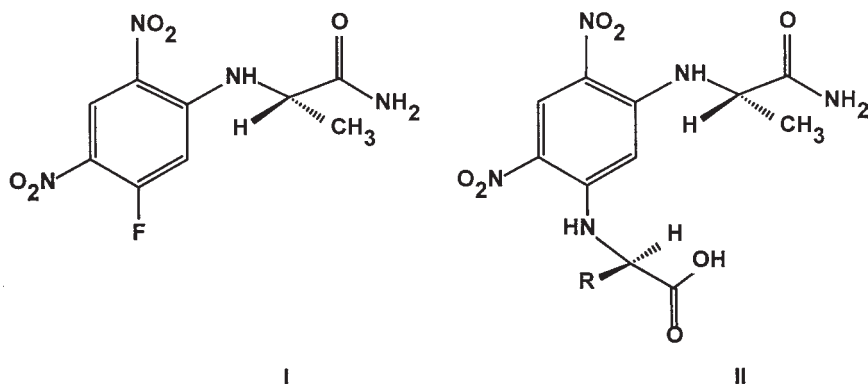


Fig. 1. Marfey's reagent (I) and L,L-diastereomer derivative from L-amino acid and the reagent (II).

precolumn derivatization with Marfey's reagent has found applications in many diverse areas of biochemical research (3–14) including determination of substrates and products in enzymic reactions of amino acids (see Note 1).

## 2. Materials

### 2.1. Vapor-Phase Protein Hydrolysis

1. Pyrex glass vials (25–50 × 5 mm) from Corning.
2. Screw-cap glass vials.
3. 6 N HCl from Pierce.

### 2.2. Derivatization Reaction

1. Amino acid standard solution H from Pierce.
2. L-Amino acid standard kit LAA21 from Sigma.
3. Marfey's reagent from Pierce. **Caution:** Marfey's reagent is a derivative of 1-fluoro-2,4-dinitrobenzene, a suspected carcinogen. Recommended precautions should be followed in its handling (15).
4. HPLC/spectroscopic grade triethylamine, methanol, acetone, and dimethyl sulfoxide (DMSO) from Fluka.

### 2.3. Chromatographic Analysis

1. Solvent delivery system: Any typical HPLC system available from a number of manufacturers can be used for resolution of derivatized amino acids. We have recently used HPLC systems from Bio-Rad (HPLC 800 equipped with an autoinjector AST 100), Hewlett Packard 1050 system and Waters system.
2. Column: A silica-based C<sub>8</sub> reverse-phase column, e.g., LiChrospher 100 RP 8 (250 × 4.6 mm; 5 μm from Macherey-Nagel), Aquapore RP 300 (220 × 4.6 mm;

- 7  $\mu\text{m}$  from Perkin-Elmer), Nucleosil 100- $\text{C}_8$  (250  $\times$  4.6 mm; 5  $\mu\text{m}$  from Macherey-Nagel), or Vydac  $\text{C}_8$  (250  $\times$  4.6 mm; 5  $\mu\text{m}$  from The Sep/a/rations group).
3. Detector: A UV/vis HPLC detector equipped with a flow cell of lightpath 0.5–1 cm and a total volume of 3–10  $\mu\text{L}$  (e.g., Bio-Rad 1790 UV/VIS monitor, Hewlett Packard Photodiode Array 1050, Waters Photodiode Array 996). The derivatized amino acids are detected at 340 nm.
  4. Peak Integration: Standard PC-based HPLC software with data analysis program can be employed to integrate and quantify the amino acid peaks (e.g., ValueChrom from Bio-Rad, Chemstation from Hewlett Packard or Millennium from Waters).
  5. Solvents: The solvents should be prepared with HPLC-grade water and degassed.
    - a. Solvent A: 13 mM trifluoroacetic acid (TFA) plus 4% (v/v) tetrahydrofuran in water.
    - b. Solvent B: Acetonitrile (50% v/v) in solvent A.

### 3. Methods

#### 3.1. Vapor-Phase Protein Hydrolysis

1. Transfer 50–100 pmol of protein sample or 20  $\mu\text{L}$  of amino acid standard solution H containing 2.5  $\mu\text{mol}/\text{mL}$  each of 17 amino acids into glass vials, and dry under reduced pressure in a SpeedVac concentrator.
2. Place the sample vials in a screw-cap glass vial containing 200  $\mu\text{L}$  of 6 N HCl.
3. Flush the vial with argon for 5–15 min, and cap it air tight.
4. Incubate the glass vial at 110°C for 24 h or 150°C for 2 h in a dry-block heater.
5. After hydrolysis, remove the glass vials from the heater, cool to room temperature, and open it slowly.
6. Remove the insert vials, wipe their outside clean with a soft tissue paper, and dry under reduced pressure.

#### 3.2. Derivatization Reaction

1. Add 50  $\mu\text{L}$  of triethylamine/methanol/water (1:1:2) to the dried sample vials, mix vigorously by vortexing, and dry them under reduced pressure (*see Note 2*).
2. Prepare derivatization reagent solution (18.4 mM) by dissolving 5 mg of Marfey's reagent in 1 mL of acetone.
3. Dissolve the dried amino acid mixture or the hydrolyzed protein sample in 100  $\mu\text{L}$  of 25% (v/v) triethylamine, add 100  $\mu\text{L}$  of the Marfey's reagent solution, and mix by vortexing (*see Note 3*).
4. Incubate the reaction vial at 40°C for 60 min with gentle shaking protected from light (*see Note 4*).
5. After incubation, stop the reaction by adding 20  $\mu\text{L}$  of 2 N HCl (*see Note 5*). Dry the reaction mixture under reduced pressure.
6. Store the dried samples at –20°C in dark until used (*see Note 6*).
7. Dissolve the dry sample in 1 mL of 50% (v/v) DMSO.



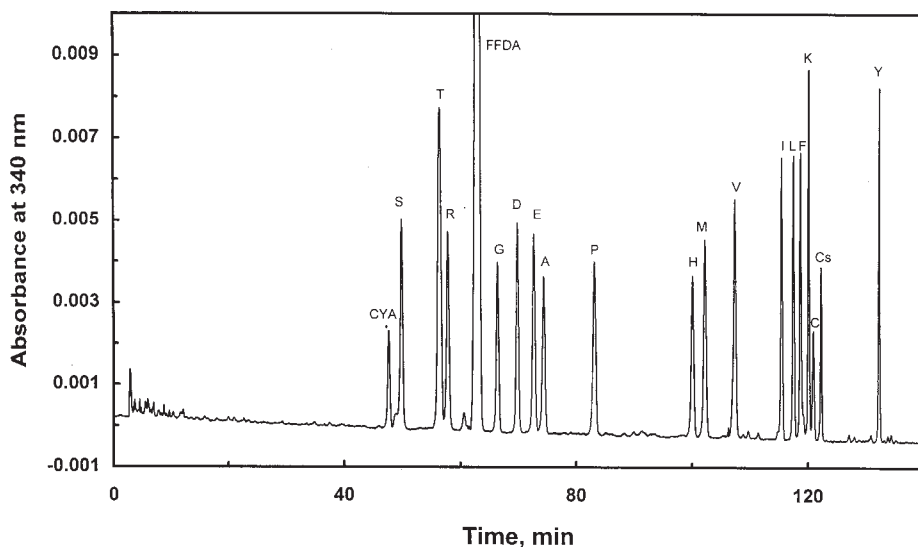


Fig. 2. Separation of 2,4-dinitrophenyl-5-L-alanine amide derivatives of L-amino acids by HPLC. A 20- $\mu$ L aliquot from the amino acid standard mixture (standard H from Pierce Chemical Company) was derivatized with Marfey's reagent. Chromatographic conditions: Column, LiChrospher 100 RP 8 (250  $\times$  4.6 mm; 5  $\mu$ m from Macherey-Nagel); sample, 100 pmol diastereomeric derivatives of 18 L-amino acids; solvent A, 13 mM trifluoroacetic acid plus 4% (v/v) tetrahydrofuran in water; solvent B, 50% (v/v) acetonitrile in solvent A; flow rate, 1 mL/min; detection at 340 nm; elution with a linear gradient of solvent B in solvent A (**Table 1**). The amino acid derivatives are denoted by single-letter code, cysteic acid as CYA, and cystine as Cs. FFDA indicates the reagent peak; the peaks without denotations are reagent-related, as shown in an independent chromatographic run of the reagent alone.

### 3.3. Chromatographic Analysis

1. Prime the HPLC system according to the manufacturer's instructions with solvent A and solvent B.
2. Equilibrate the column and detector with 90% solvent A and 10% solvent B.
3. Bring samples for analysis to room temperature; dilute, if necessary, with solvent A, and inject 20  $\mu$ L (50–1000 pmol) onto the column (*see Note 7*).
4. Elute with gradient as described in **Table 1** (*see Note 8*). Resolution of 2,4-dinitrophenyl-5-L-alanine amide derivatives of 18 commonly occurring L-amino acids and of cysteic acid is achieved within 120 min (**Fig. 2**) (*see Note 9*).
5. Determine the response factor for each amino acid from the average peak area of standard amino acid chromatograms at 100, 250, 500, and 1000 pmol amounts.
6. When HPLC is completed, wash the column and fill the pumps with 20% (v/v) degassed methanol for storage of the system. Before reuse, purge with 100% methanol.

**Table 1**  
**HPLC Program for Resolution of Amino Acids Derivatized with Marfey's Reagent**

| Time (min) | % Solvent A | % Solvent B | Input              |
|------------|-------------|-------------|--------------------|
| 0          | 90          | 10          |                    |
| 15         | 90          | 10          | Detector auto zero |
| 15.1       | 90          | 10          | Inject             |
| 115        | 50          | 50          |                    |
| 165        | 0           | 100         |                    |
| 170        | 0           | 100         |                    |

#### 4. Notes

1. For analysis of amino acids as substrates or products in an enzymic reaction, deproteinization is required. Add 4 M perchloric acid to a final concentration of 1 M and incubate on ice for at least 15 min. Excess perchloric acid is precipitated as KClO<sub>4</sub> by adding an equal volume of ice-cold 2 M KOH. After centrifugation for 15 min at 4°C, the supernatant is collected, dried and used for derivatization.
2. Complete removal of HCl is absolutely essential for quantitative reaction between amino acids and Marfey's reagent.
3. The molar ratio of the Marfey's reagent to the total amino acids should not be more than 3:1.
4. During derivatization, the color of the reaction mixture turns from yellow to orange-red.
5. On acidification, the color of the reaction mixture turns to yellow.
6. The dried amino acids are stable for over one month when stored at -20°C in the dark. In 50% (v/v) DMSO, derivatives are stable for 72 h at 4°C and for over 6 wk at -20°C.
7. To obtain good reproducibility, an HPLC autoinjector is highly recommended.
8. Silica-based C<sub>8</sub> columns from different commercial sources produce base-line resolution of the 19 amino acid derivatives; nevertheless, the gradient slope of solvent B should be optimized for each column. *S*-Carboxymethyl-L-cysteine and tryptophan, if included, are also separated in a single chromatographic run (*see ref. 2*).
9. The identity of each peak is established by adding a threefold molar excess of the amino acid in question. A reagent blank should be run with each batch of the Marfey's reagent to identify reagent-related peaks. Excess reagent interferes with base-line separation of the arginine and glycine peaks. Lysine and tyrosine are separated as disubstituted derivatives.

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## Anion Exchange Chromatography and Integrated Amperometric Detection of Amino Acids

Petr Jandik, Christopher Pohl, Victor Barreto,  
and Nebojsa Avdalovic

### 1. Introduction

During the 1980s and 1990s, electrochemical detection has gained acceptance as a method of choice for some important biomolecules. Initially, the technique was utilized with glassy carbon electrodes and in the mode of constant potential amperometry. The first widely used applications were for catecholamines and their derivatives (1). Today, most laboratories around the world still rely on electrochemical detection when it comes to analyzing catecholamines in biological fluids. Although that approach has been highly successful, the next important advance came in the form of pulsed amperometric detection (PAD). The introduction of voltage pulses broadens the scope of electrochemical detection by regenerating the electrode in time for each new detection measurement. Gradual or sudden fouling of the electrode surface by the products of electrode reaction and the corresponding decreased detection response is eliminated with properly designed PAD techniques. The first important application of PAD was for carbohydrates following their separation by anion exchange (2,3).

The analysis of amino acids based on anion exchange chromatography and integrated pulsed amperometric detection (IPAD) has evolved from Johnson and Polta's (4) original approach reported in 1983. The first high-performance liquid chromatography (HPLC) system based on that methodology was introduced in 1999 (5).

#### 1.1. Anion Exchange of Amino Acids

Alkaline pH is a requirement for the IPAD detection of amino acids on gold electrodes. Alkaline pH of anion exchange eluents thus makes IPAD possible.

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**Table 1**  
**Amino Acids and Other Compounds Analyzed by AAA Direct**

| Compound                   | Type                      | Compound             | Type                           |
|----------------------------|---------------------------|----------------------|--------------------------------|
| 1 Arginine                 | Amino acid                | 29 Glycine           | Amino acid                     |
| 2 Mannitol                 | <i>Alditol</i>            | 30 Ribose            | <i>Monosaccharide</i>          |
| 3 Fucose                   | <i>Monosaccharide</i>     | 31 Valine            | Amino acid                     |
| 4 Hydroxy-Lysine           | Hydroxy-amino acid        | 32 Hydroxy-Proline   | Hydroxy-amino acid             |
| 5 Gly-Lys                  | <i>Peptide</i>            | 33 Gly-OHPro         | <i>Peptide</i>                 |
| 6 Ornithine                | <i>Amino acid</i>         | 34 Serine            | Amino acid                     |
| 7 MetSulfoxide             | <i>Ox. Product of Met</i> | 35 Sucrose           | <i>Disaccharide</i>            |
| 8 Lysine                   | Amino acid                | 36 Gly-Ser           | <i>Peptide</i>                 |
| 9 Gly-Pro                  | <i>Peptide</i>            | 37 Proline           | Amino acid                     |
| 10 Rhamnose                | <i>Monosaccharide</i>     | 38 Isomaltose        | <i>Disaccharide</i>            |
| 11 Glucosamine             | Amino-sugar               | 39 IsoLeucine        | Amino acid                     |
| 12 Galactosamine           | Amino-sugar               | 40 Leucine           | Amino acid                     |
| 13 Arabinose               | <i>Monosaccharide</i>     | 41 Methionine        | Amino acid                     |
| 14 Glutamine               | Amino acid                | 42 NorLeucine        | Amino acid                     |
| 15 Citrulline              | Amino acid                | 43 Taurine           | Amino sulfonic acid            |
| 16 Mannose                 | <i>Monosaccharide</i>     | 44 Histidine         | Amino acid                     |
| 17 $\alpha$ -Amino-butyric | Amino acid                | 45 Phospho-Arginine  | Phospho-amino acid             |
| 18 Asparagine              | Amino acid                | 46 Diaminopimelic    | Amino acid                     |
| 19 Glucose                 | <i>Monosaccharide</i>     | 47 Phenylalanine     | Amino acid                     |
| 20 Galactose               | <i>Monosaccharide</i>     | 48 Glutamate         | Amino acid                     |
| 21 Xylose                  | <i>Monosaccharide</i>     | 49 Aspartate         | Amino acid                     |
| 22 (Gly)3                  | <i>Peptide</i>            | 50 Cystine           | Amino acid                     |
| 23 Alanine                 | Amino acid                | 51 Phospho-Serine    | Phospho-amino acid             |
| 24 Gly-Gly                 | <i>Peptide</i>            | 52 Phospho-Threonine | Phospho-amino acid             |
| 25 Threonine               | Amino acid                | 53 Tyrosine          | Amino acid                     |
| 26 (Gly)4                  | <i>Peptide</i>            | 54 Phospho-Tyrosine  | Phospho-amino acid             |
| 27 $\gamma$ -Amino-butyric | Amino acid                | 55 Cysteic Acid      | <i>Ox. Prod. of Cys/(Cys)2</i> |
| 28 Methionine Sulfone      | <i>Ox. Product of Met</i> | 56 Tryptophan        | Amino acid                     |

The italics denote compounds usually separated by modified gradients. The actual elution order for these compounds may vary depending on the gradient method selected. **Note:** Separation of methionine oxidation products requires a different column temperature.

**Table 2**  
**Gradient Conditions for Analyzing Amino Acids in Hydrolysates**

| Time (min) | %E1 | %E2 | %E3 | Curve | Comments<br>(Flow rate: 0.25 mL/min)           |
|------------|-----|-----|-----|-------|------------------------------------------------|
| Init       | 76  | 24  | 0   |       | Autosampler fills the sample loop              |
| 0.00       | 76  | 24  | 0   |       | Valve from Load to Inject                      |
| 2          | 76  | 24  | 0   |       | Begin OH gradient, valve from Inject to Load   |
| 8          | 64  | 36  | 0   | 8     |                                                |
| 11         | 64  | 36  | 0   |       | Begin acetate gradient                         |
| 18         | 40  | 20  | 40  | 8     |                                                |
| 21         | 44  | 16  | 40  | 5     |                                                |
| 23         | 14  | 16  | 70  | 8     |                                                |
| 42         | 14  | 16  | 70  |       |                                                |
| 42.1       | 20  | 80  | 0   | 5     | Column wash with hydroxide                     |
| 44.1       | 20  | 80  | 0   |       |                                                |
| 44.2       | 76  | 24  | 0   | 5     | Start of reequilibration to initial conditions |
| 75         | 76  | 24  | 0   |       |                                                |

As expected, anion exchange causes the amino acids to elute in a reverse order to that observed on the more commonly used cation exchangers (6). **Table 1** lists a selection of amino acids and other compounds that can be separated by the anion exchange method. All the separations are achieved by ternary gradients employing three simple eluents (E1: water, E2: 0.25 M NaOH, and E3: 1.0 M NaAc). The three most frequently utilized gradient programs are listed in **Tables 2–4**. The corresponding chromatograms of standard mixtures are shown in **Figs. 1–3**.

### 1.2. Principle of Integrated Pulsed Amperometric Detection

With IPAD, one can achieve sensitivities comparable to or better than those obtained by detecting fluorescence of various amino acid derivatives. The most important advantage of IPAD is that it does not require any form of derivatization to make the analytes detectable. As illustrated in **Figs. 2 and 3**, IPAD also detects other classes of compounds (numerous examples are also shown in **Table 1**).

The routine use of IPAD for amino acid analysis is relatively new. However, many reports on IPAD of amino acids have been published during the past 15 yr (5,7–12), and a comprehensive book (10) dedicated to the technique has recently become available.

**Table 3**  
**Group Gradient: Gradient Conditions for Simultaneous Separations of Amino Acids and Carbohydrates**

| Time (min) | %E1 | %E2 | %E3 | Curve | Comments                                       |
|------------|-----|-----|-----|-------|------------------------------------------------|
| Init       | 94  | 6   | 0   |       | Autosampler fills the sample loop              |
| 0.00       | 94  | 6   | 0   |       | Valve from Load to Inject                      |
| 2          | 94  | 6   | 0   |       | Begin OH gradient, valve from Inject to Load   |
| 12         | 68  | 32  | 0   | 8     |                                                |
| 16         | 68  | 32  | 0   |       | Begin acetate gradient                         |
| 24         | 36  | 24  | 40  | 8     |                                                |
| 40         | 36  | 24  | 40  |       |                                                |
| 40.1       | 20  | 80  | 0   | 5     | Column wash with hydroxide                     |
| 42.1       | 20  | 80  | 0   |       |                                                |
| 42.2       | 94  | 6   | 0   | 5     | Start of reequilibration to initial conditions |
| 62         | 94  | 6   | 0   |       |                                                |

**Table 4**  
**Glucose Gradient: Gradient Conditions for Simultaneous Separations of Amino Acids and Glucose**

| Time (min) | %E1 | %E2 | %E3 | Curve | Comments                                       |
|------------|-----|-----|-----|-------|------------------------------------------------|
| Init       | 84  | 16  | 0   |       | Autosampler fills the sample loop              |
| 0.00       | 84  | 16  | 0   |       | Valve from Load to Inject                      |
| 2          | 84  | 16  | 0   |       | Begin OH gradient, valve from Inject to Load   |
| 12         | 68  | 32  | 0   | 8     |                                                |
| 16         | 68  | 32  | 0   |       | Begin acetate gradient                         |
| 24         | 36  | 24  | 40  | 8     |                                                |
| 40         | 36  | 24  | 40  |       |                                                |
| 40.1       | 20  | 80  | 0   | 5     | Column wash with hydroxide                     |
| 42.1       | 20  | 80  | 0   |       |                                                |
| 42.2       | 84  | 16  | 0   | 5     | Start of reequilibration to initial conditions |
| 62         | 84  | 16  | 0   |       |                                                |

Like most other contemporary electrochemical techniques, IPAD is carried out in a three-electrode detection cell (**Fig. 4A**). The gold working electrode is subjected to several carefully chosen potential steps (E1 to E6 in **Fig. 4B**). The

1. Arginine
2. Hydroxylysine
3. Lysine
4. Glutamine
5. Asparagine
6. Alanine
7. Threonine
8. Glycine
9. Valine
10. Hydroxyproline
11. Serine
12. Proline
13. Isoleucine
14. Leucine
15. Methionine
16. Norleucine
17. Histidine
18. Phenylalanine
19. Glutamate
20. Aspartate
21. Cystine
22. Tyrosine

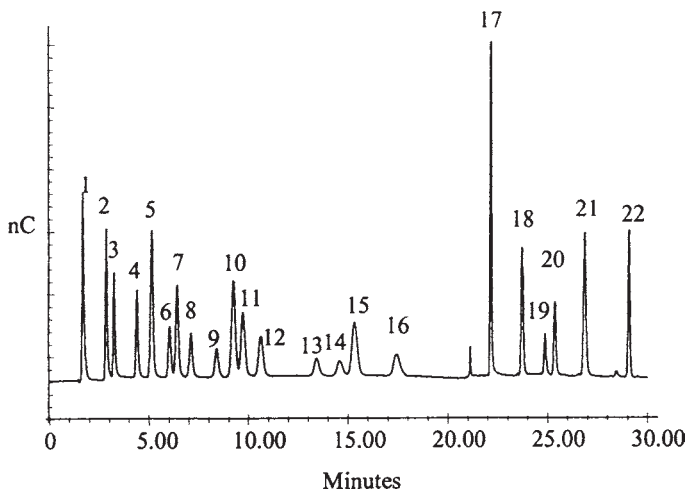


Fig. 1. Separation of amino acids using the Standard Gradient Conditions in **Table 2**.

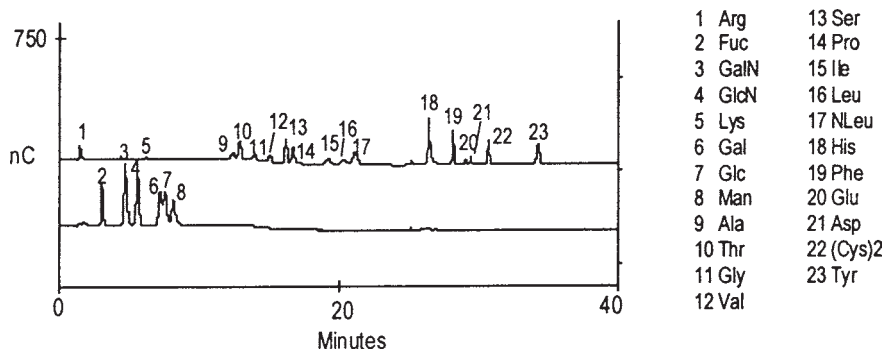


Fig. 2. Simultaneous separations of amino acids, amino sugars, and glucose using the gradient from **Table 4**.

potentials are adjusted between the gold surface and a silver/silver chloride reference electrode (more recently in reference to a glass/silver/silver chloride combination electrode (5), see **Table 5**). Some chemistries that occur at the gold electrode surface are described in **Fig. 4B**. Additionally, oxidation and sometimes also adsorption and oxidative desorption are observed in the presence of the analytes in **Table 1**. Electrons transferred to the gold surface during



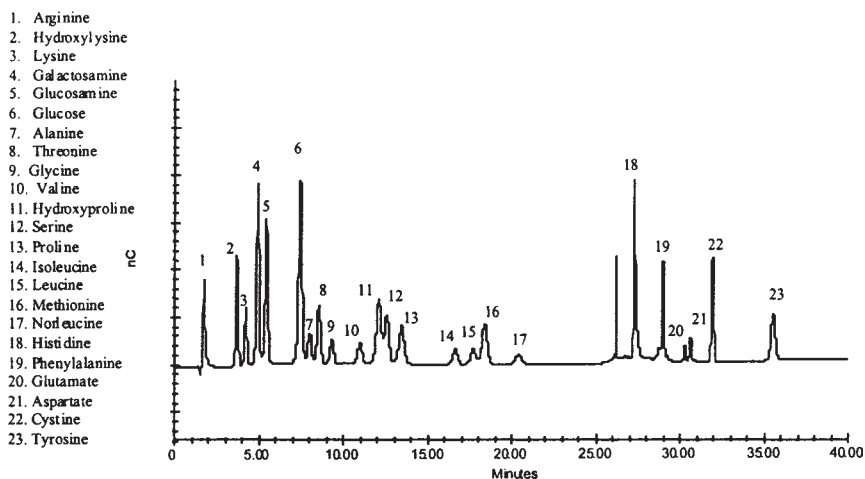


Fig. 3. Optimized gradient (*see Table 3*) allowing separations of amino acids, amino sugars, and sugars in one run.

analyte oxidation give rise to an electric current between the gold electrode and counterelectrode. The use of two separate electrodes (reference and counterelectrode) for voltage application and current sensing was introduced in the early days of electrochemistry. It helps to avoid significant electric currents running through the reference electrode, thereby keeping its potential stable over a long period. If the main portion of the current is allowed to pass through the reference electrode, it causes the silver chloride layer to dissolve from the silver wire and consequently modifies the value of the reference potential. The long-term stability of the reference potential is essential for good reproducibility of the detection response.

The current resulting from the analyte oxidation at the gold electrode is measured by integration over the time period indicated in **Fig. 4B**. The detection signal thus obtained is in nanocoulombs (coulomb = amperes second) rather than in amperes.

## 2. Materials

### 2.1. Equipment

1. Equipment for vacuum hydrolysis: Most common hydrolysis methods can be performed in borosilicate glass tubes equipped with Teflon plugs (P/N 29550, Pierce, Rockford, IL). Tubes are placed into aluminum heating block (P/N 18808, Pierce) and heated up by a Heating Module (P/N 18870 [120V] Pierce).
2. Equipment for evaporative centrifugation: Hermetically sealing centrifuge (SVC 100, Savant, Farmingdale, NY), refrigerated vapor trap (RVT 400, Savant), vacuum gage MVG-5 connected to DV-24 gauge tube (both from Savant), chemi-

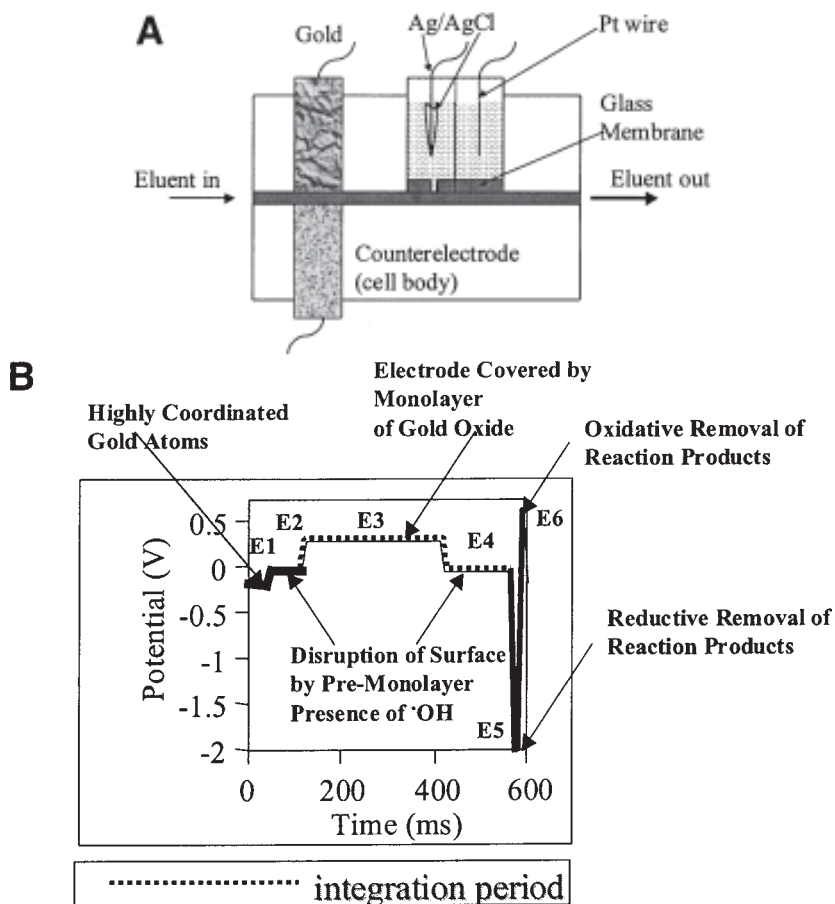


Fig. 4. (A) Three-electrode detection cell allowing a choice between silver/silver chloride and pH referencing. See ref. 5 for more details. (B) The surface of the gold working electrode at different potentials of the detection waveform.

cal vapor trap (SCT 120, Savant) containing acid adsorbent cartridge (DC120A) and a vacuum pump (e.g., P/N 7893-F10, Thomas Scientific, Swedesboro, NJ) are interconnected by thick-walled, half-inch ID, Tygon tubing (e.g., P/N 9561-E28, Thomas Scientific).

- HPLC hardware: AAA Direct System (Dionex Corporation, Sunnyvale, CA) is optimized for anion exchange/IPAD of amino acids. It includes the AS50 PEEK autosampler, a Thermal Compartment, GP50 PEEK microbore pump with Vacuum Degas Option, and an ED40 electrochemical detector with AAA-Certified Gold Amperometry Cell. Mobile phase components are kept under helium or nitrogen at all times. For fully automatic operation, the PeakNet Control Windows 98 Workstation should also be selected.

**Table 5**  
**Detection Waveform for Amino Acids, Amino Sugars,**  
**and Sugars**

| Time (ms) | Potential (V) <sup>a</sup> vs pH | Integration |
|-----------|----------------------------------|-------------|
| 0         | 0.13                             |             |
| 40        | 0.13                             |             |
| 50        | 0.28                             |             |
| 110       | 0.28                             | Begin       |
| 120       | 0.61                             |             |
| 410       | 0.61                             |             |
| 420       | 0.28                             |             |
| 560       | 0.28                             | End         |
| 570       | -1.67                            |             |
| 580       | -1.67                            |             |
| 590       | 0.93                             |             |
| 600       | 0.13                             |             |

<sup>a</sup>All potentials as in **Fig. 4B** +0.33V.

4. Anion-exchange column (*see Note 1*): Dionex distributes the high-pH-stable anion-exchange columns optimized for the separation of amino acids and other compounds under the name “AminoPac PA10” (*see Note 1*). They are typically used as a combination of guard column (2-mm id, 50-mm long) and main column (2-mm id, 250 mm long).

## 2.2. Reagents (*see Note 2*)

1. Oxidation and/or hydrolysis: formic acid (0128-01, J.T. Baker, Phillipsburg, NJ), 30% hydrogen peroxide (0128-01, J.T. Baker), 6 *N* HCl (24309, Pierce), 4.0 *M* MSA (25555, Pierce), 4.2 *M* NaOH (30,657-6, Sigma, St. Louis, MO), 13 *M* TFA (53102, Pierce).
2. Chromatographic eluents: 18 megohm water from a Milli-Q (Millipore, MA), 50% sodium hydroxide (UN1824, Certified Grade, Fisher, Pittsburgh, PA), anhydrous sodium acetate (71179, Microselect Grade, Fluka, Milwaukee, MI).
3. Diluents and standards: norleucine (N1398, Sigma), sodium azide (S 8032, Sigma), Standard Reference Material Amino Acids (SRM 2389 NIST, Gaithersburg, MD), cysteic acid (C7630, Sigma), methionine sulfone (M0751, Sigma), methionine sulfoxide (M1001, Sigma), Mixture of Six Carbohydrates (P/N 43162, Dionex).

## 2.3. Preparation of Eluents and Standards (*see Notes 3–6*)

1. Eluent E1: Deionized water — Filter the pure deionized water through 0.2- $\mu$ m nylon filters, then transfer into a DX-500 eluent bottle. Use, for example, steril-

- ized, sterile-packed, 1-L-funnel, vacuum-filtration units from Nalge (28194–514, VWR, West Chester, PA). We have found these ideal for filtration of all eluents. Seal the filtered water immediately.
2. Eluent E2: 250 mM sodium hydroxide — The first step in preparing sodium hydroxide eluent is to filter a water aliquot (typically 1.0 L) using the sterilized Nalgene filtration unit as aforementioned. Hermetically seal the filtered water immediately after filtration while preparing a disposable glass pipet (10.0-mL sterile, serological pipets, Fisher) and a pipet filler. Using a pipet filler, draw an aliquot of 50% sodium hydroxide into the pipet. Most serological 10.0-mL pipets can be filled to the 13.1-mL volume required for 1.0 L of 250 mM sodium hydroxide. Unseal the filtered water and insert the full pipet approx 1 in. below the water surface, and release the sodium hydroxide. If done properly and without stirring, most of the concentrated sodium hydroxide remains at the lower half of the container and the carbon dioxide adsorption rate is much lower than with a 250 mM sodium hydroxide solution. Seal the container immediately after the sodium hydroxide transfer is complete. Remember to immediately put the screw-cap back on the 50% hydroxide bottle. Mix the contents of the tightly sealed container holding the 250 mM hydroxide. Unscrew the cap of the eluent bottle E2 attached to the DX-500. Allow the helium or nitrogen gas to blow out of the cap. Unseal the bottle holding 250 mM hydroxide, and immediately start the transfer into the eluent bottle E2. Try to minimize the carbon dioxide absorption by holding the gas orifice of the bottle cap as close as possible to the 250 mM hydroxide during the transfer. Follow aseptic procedures. With the inert gas still blowing, put the cap back on the eluent bottle. Allow the pressure to build inside the bottle and reopen the cap briefly several times to allow trapped air to be gradually replaced by the inert gas.
  3. Eluent E3: 1.0 M sodium acetate — Dissolve 82.04 g of anhydrous sodium acetate in approx 750 mL of pure deionized water. Seal the container during the dissolution step. Make up to 1.0 L with water and filter through a 0.2- $\mu$ m Nylon filter. We recommend using sterile Nalgene vacuum filtration units for this step (*see* above). Transfer the filtered sodium acetate eluent into the eluent bottle E3 of the DX-500, using the same procedure as for the sodium hydroxide.
  4. Diluent solutions containing sodium azide: Prepare 1 or 2 L of an azide diluent solution containing 20 mg  $\text{NaN}_3/\text{L}$ . Use the solution without norleucine whenever an internal standard is not required. Store the pure azide solution in a closed container at room temperature.
  5. Diluent solution containing norleucine internal standard: Prepare a 4.0 mM stock solution of norleucine (524.8 mg/L) in 0.1 M HCl. Dilute 500x with a deionized water solution containing 20 mg azide per liter. The resulting diluent solution is stable for months if stored in a refrigerator. Use it to prepare final dilutions from standard stock solution and to redissolve hydrolysate samples after evaporation to dryness.
  6. Amino acid standards: Use aliquots of Standard Reference Material 2389 diluted 500x or 250x with the diluent (*see Subheading 2.3., step 5*) to obtain 5  $\mu$ M or 10

$\mu\text{M}$  standard solutions. The standard solutions thus prepared remain stable for months if stored in a refrigerator. The trace of sodium azide introduced with the diluent solution stabilizes standards up to 48 h at room temperature.

7. Standard mixture of oxidation products: Prepare a 5 mM stock solution of oxidation standards by dissolving 82.6 mg of methionine sulfoxide, 90.6 mg of methionine sulfone, and 84.6 mg of cysteic acid in 100.0 mL of 0.1 M hydrochloric acid. Transfer 100  $\mu\text{L}$  of the stock solution to a 100 mL volumetric flask and fill up to volume with the diluent from **Subheading 2.3., step 5**.
8. Carbohydrate standard: Add 1.0 mL of sodium azide diluent (**Subheading 2.3., step 4**) to the Dionex carbohydrate standard to obtain a 100- $\mu\text{M}$  stock solution of each of the six components. Vortex to support the dissolution of the dry residue. Using the sodium azide diluent again, dilute an aliquot of the stock solution to obtain a 2.0  $\mu\text{M}$  concentration of all four sugars and two amino sugars.

### 3. Methods

#### 3.1. Oxidation of Methionine and Cystine/Cysteine Prior to Hydrolysis

1. Prepare the performic acid reagent freshly from formic acid and hydrogen peroxide before each application. Mix 900  $\mu\text{L}$  of the formic acid and 100  $\mu\text{L}$  30% hydrogen peroxide in a microcentrifuge tube. Vortex the mixture.
2. Allow the mixture to adjust to room temperature.
3. Pipet an aliquot containing about 1  $\mu\text{g}$  of the sample into another microcentrifuge vial and remove the liquid by evaporative centrifugation.
4. Add 250  $\mu\text{L}$  of the chilled performic acid solution to the dry sample. Seal the microcentrifuge vial containing the reaction mixture, and keep it at room temperature for 4 h.
5. Carry out evaporative centrifugation to remove the rest of the performic acid solution from the sample.
6. Proceed with hydrolysis (go to **Subheading 3.2., step 2**. Addition of 6 M HCl).

#### 3.2. Hydrolysis (see Notes 7 and 8)

1. Pipet a volume of protein or peptide solution into a clean microcentrifuge tube. Carry out evaporative centrifugation to remove the liquid phase.
2. To a dry sample containing no more than 200  $\mu\text{g}$  of protein or peptide, add 200  $\mu\text{L}$  of the hydrolysis agent (6 M HCl, 2 N TFA, 4.0 M MSA, or 4.2 M NaOH). If you are processing a performic-acid-oxidated sample, add 200  $\mu\text{L}$  of 6 M HCl to the dry sample from **Subheading 3.1., step 5**.
3. Speed up the dissolution of the solid residue in the hydrolysis agent by vortexing. Transfer the liquid contents into a vacuum hydrolysis tube (*see Note 9*).
4. Insert the Teflon plug, and screw it down leaving only a narrow passageway open. Attach the side arm of the tube to the vacuum tubing connected to a three-way stopcock. Connect the stopcock to a vacuum pump and to a source of inert gas.
5. Alternate between vacuum and inert gas to remove all traces of oxygen from the space above the hydrolysis mixture.

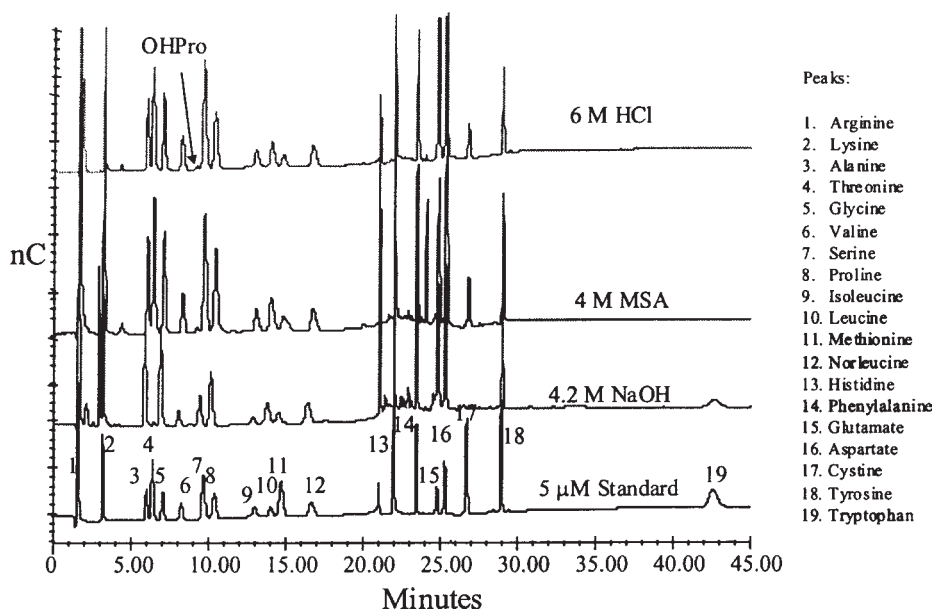


Fig. 5. Soybean powder: Chromatographic results from three different hydrolytic procedures.

6. Screw down the Teflon plug to seal the vacuum inside the tube.
7. Place the sealed tube into a preheated Reacti-Therm Module, and hydrolyze (*see Note 7* for temperature setting and timing of the respective hydrolytic protocols).
8. Remove the tube from the heating module and allow it to cool before breaking the vacuum inside the tube.
9. Transfer the hydrolysate into a clean microcentrifuge tube.
10. Carry out evaporative centrifugation (*see Note 10*).
11. Redissolve the dry contents of the microcentrifuge tube in a suitable volume of azide solution with/without internal standard (typically 1x to 4x of the volume during the hydrolysis). Dry samples from **step 10** are stable for up to 1 yr if stored in a freezer. The redissolved samples can be stored for up to 1 mo in a refrigerator. At room temperature, the azide-stabilized samples remain unchanged for at least 2 d. Proceed with chromatographic separation.

### 3.3. Chromatographic Separations of HCl, MSA, and NaOH Hydrolysates (*see Note 11; Fig. 5*)

The Standard Gradient in **Table 2** (*see Note 12*) offers a useful separation of hydrolysate amino acids and all other compounds listed in regular print in **Table 1**. The italics indicate the need for a modified gradient. All modified gradients, however, utilize the same three eluents (E1, E2, and E3; *see Subheading 1.1.*)

and are carried out at a constant flow rate of 0.250 mL/min. Selectivity changes are achieved by varying the initial gradient conditions or by changing the separation temperature (*see* **Notes 13, 14**).

1. Verify proper operation of the chromatographic system as described in the Dionex System Manual (background signal at initial gradient conditions < 80 nC, gradient rise < 30 nC, *see* Dionex Document No. 031481, **Notes 15–17**).
2. Transfer the samples from **Subheading 3.2., step 11** into 1.5- or 0.5-mL autosampler vials (*see* **Notes 18–20**). Place vials into the autosampler, note their positions.
3. Fill a 1.5-mL vial with 8  $\mu$ M solution of hydrolysate standard. Fill another 1.5-mL vial with water. Place vials into the autosampler, note their positions.
4. Write a PeakNet Schedule (*see* PeakNet Software User's Guide, Dionex Document No. 034914) specifying 25.0- $\mu$ L full-loop injections and assigning the gradient pump method from **Table 2** together with the detection method from **Table 5** to each of the samples and standards. The sequence of injections should start with at least one injection of water, followed by one or two injections of the standard. Ideally, there should be at least two standard injections at the beginning, in the middle, and at the end of each sequence of injection.
5. Load the PeakNet Schedule from **Subheading 3.3., step 4** into the PeakNet Run Window and start the automatic run.

### **3.4. Chromatographic Analysis of Oxidation Products of Cysteine and Methionine**

1. Equilibrate the HPLC system using the initial conditions of standard gradient in **Table 2** with the column thermostat set at 35°C (*see* **Notes 13 and 21**). The background signal should be <100 nC.
2. Fill a 1.5-mL vial with water. Place vial into autosampler. Note its position.
3. Fill a 1.5-mL vial with 8 mM solution of hydrolysate standard. Place vial into autosampler. Note its position.
4. Fill a 1.5-mL vial with 5  $\mu$ M solution of oxydation products standard. Place vial into autosampler. Note its position.
5. Transfer the samples from **Subheading 3.2., step 11** into 0.5- or 1.5-mL autosampler vials (**Note 20**). Note their positions.
6. Write a PeakNet Schedule (*see* PeakNet Software User's Guide, Dionex Document No. 034914) specifying 25.0- $\mu$ L full-loop injections and assigning the gradient method from **Table 2** together with the detection method from **Table 5** to each of the samples and standards (verify the 35°C column temperature setting). The sequence of injections should start with at least one injection of water, followed by one or two injections of each of the two standards. Ideally, there should be two injections of each of the two standards at the beginning, in the middle and at the end of each sequence of injection.
7. Load the PeakNet Schedule from **Subheading 3.4., step 6** into the PeakNet Run Window, and start the automatic run (**Fig. 6**).

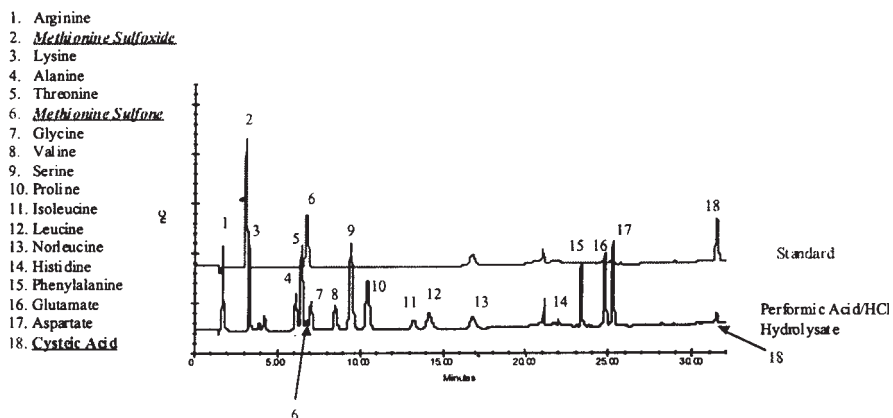


Fig. 6. Bovine Colostrum Hydrolysate: Analysis of oxidation products of cystine/cysteine and methionine. The separation of a mixture consisting of methionine-sulfoxide, methioninesulfone, and cysteic acid is included for easier identification of oxidation products. The methioninesulfoxide is not detected in the sample hydrolysed by HCl. To separate methioninesulfone from the neighboring threonine and glycine, the column temperature has to be maintained at 35°C. Since tyrosine is destroyed during the oxidative step, cysteic acid, the product of cysteine/cystine oxidation, elutes as an isolated peak at approx 32 min.

### 3.5. Chromatographic Analysis of Sugars, Amino Sugars, and Amino Acids in Hydrolysates Using a Single Gradient Method

The gradient method in **Table 3** makes possible a simultaneous separation of sugars, amino sugars and amino acids, *see* the chromatograms in **Fig. 7**.

1. Hydrolyze three aliquotes of a glycoprotein using the procedures in **Note 7A–C** and the hydrolytic protocol **Subheading 3.2., steps 1–11**.
2. Verify proper operation of the chromatographic system as described in the Dionex System Manual (background signal at initial gradient conditions < 80 nC, gradient rise < 30 nC, *see* Dionex Document No. 031481).
3. Fill a 1.5-mL vial with water. Place vial into autosampler. Note its position.
4. Fill a 1.5-mL vial with 8 mM solution of hydrolysate standard. Place vial into autosampler. Note its position.
5. Fill a 1.5-mL vial with 8 mM solution of hydrolysate standard (from **Subheading 2.3., step 6**). Place vial into autosampler. Note its position.
6. Fill a 1.5-mL vial with 2 mM solution of carbohydrate standard (from **Subheading 2.3., step 8**). Place vial into autosampler. Note its position.
7. Transfer the samples from step **Subheading 3.2., step 11** of each of the three hydrolytic protocols in **Note 7A–C** into 0.5-mL autosampler vials (**Note 20**).
8. Place sample vials into autosampler. Note their position.



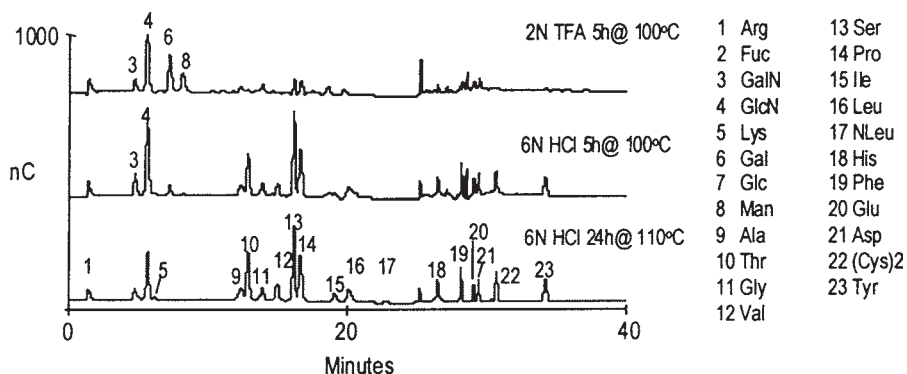


Fig. 7. Identical amounts (200  $\mu\text{g}$ ) of fetuin were hydrolyzed by three different procedures. The volumes of hydrolyzing agents were 200  $\mu\text{L}$  2 N TFA and 400  $\mu\text{L}$  6 M HCl. The hydrolysates were dried down by evaporative centrifugation and redissolved in 1.0 mL of diluent (**Subheading 2.3.5.**) All three hydrolysates could be analyzed by the same gradient method (**Table 3**).

9. Write a PeakNet Schedule (*see* PeakNet Software User's Guide, Dionex Document No. 034914) specifying 25.0- $\mu\text{L}$  full-loop injections and assigning the gradient method from **Table 3** together with the detection method from **Table 5** to each of the samples and standards. The sequence of injections should start with at least one injection of water, followed by one or two injections of each of the two standards. Ideally, there should be two injections of each of the two standards at the beginning, in the middle, and at the end of each sequence of injection.
10. Load the PeakNet Schedule from **Subheading 3.5., step 9** into the PeakNet Run Window and start the automatic run.

### 3.6. Data Processing Following Chromatographic Separation of Hydrolysate Samples (Notes 22–25)

1. At the conclusion of the sequence defined in step **Subheading 3.3., step 5**, **Subheading 3.4., step 7**, or **Subheading 3.5., step 10**, inspect the standard chromatograms first. In a properly operated system, the reproducibility of peak areas is < 5% RSD and reproducibility of retention times is < 1% RSD. If your results do not conform with the %RSD values specified for the standard injections, optimize integration and repeat post-run data processing (*see* Verification of Reproducibility, Dionex LPN. 01156-01).
2. Create a Component Table listing amino acids (or amino acids, sugars, amino sugars and oxidation products, if applicable) of interest, retention times, and mean values of peak areas for the concentrations of the injected standard (*see* PeakNet Software User's Guide, Dionex Document No. 034914). Using the optimized peak integration from **Subheading 3.6., step 1**, apply post-run processing to the hydrolysate samples to calculate amino acid concentrations (or sugar, amino

sugar, amino acid, and oxidation products concentrations, if applicable) in the hydrolysate samples.

### 3.7. Off-Line Analysis of Cell-Culture Samples (see Note 26)

The analysis of amino acids and sugars in cell cultures is part of a broader strategy aimed at maximizing the product titer (9,16). Without analytical control, glucose concentration has to be maintained at excessively high levels, whereas the concentration of glutamine, another important nutrient, may decrease to the levels too low for the process to be efficient. The strong rise in acidity stemming from excessive lactic acid formation is neutralized by adding sodium hydroxide to the cell culture at regular intervals.

With the help of chromatographic analysis, an improved feeding strategy can be designed that makes it possible to replenish the glucose levels according to actual demand and keeps it at much lower levels. Lower levels of glucose result in less frequent additions of sodium hydroxide and a slower increase of ionic strength. The level of glutamine can also be kept relatively low and within very narrow boundaries. The production of ammonia is minimized and depletion of glutamine avoided.

1. Verify proper operation of the chromatographic system as described in the Dionex System Manual (background signal at initial gradient conditions < 80 nC, gradient rise < 30 nC, *see* Dionex Document No. 031481).
2. Equilibrate the HPLC system using gradient conditions in **Table 4** and the detection method from **Table 6**.
3. Run at least three 25- $\mu$ L injections of standard (use NIST standard from **Subheading 2.3., step 6**) and complete the PeakNet Component Table (*see* PeakNet Software User's Guide, Dionex Document No. 034914).
4. Take samples (1–5 mL) of fermentation at timed intervals.
5. Using a sterilized syringe, pass an aliquot of the sample through a sterile 0.45- $\mu$ m filter.
6. Dilute an aliquot of each filtered sample 250x using the diluent from **Subheading 2.3., step 4**.
7. Inject 25  $\mu$ L of the dilute sample into the chromatographic system.
8. Initiate a run using a pump method based on **Table 4** gradient (*see* **Note 27**) and **Table 6**-detection method (*see* **Note 28**). Make the Component Table from **Subheading 3.6., step 2**, also a part of the PeakNet method, to obtain automatic calculation of amino acid concentrations in the fermentation broth (**Fig. 8**).

## 4. Notes

1. The AminoPac PA10 columns are packed with 9- $\mu$ m-diameter microporous resin beads, consisting of ethylvinylbenzene crosslinked with 55% divinylbenzene. The MicroBead layer agglomerated on the surface of these beads consists of 200-nm VBC-based (vinylbenzylchloride) latex particles functionalized with a precise

**Table 6**  
**Modified for the Nonselective and Selective Detection**

| Time (ms) | Potential (V) <sup>a</sup><br>vs pH | Integration<br>to detect<br>Amino acids, sugars,<br>and amino sugars | OH-selective<br>integration |
|-----------|-------------------------------------|----------------------------------------------------------------------|-----------------------------|
| 0         | 0.13                                |                                                                      |                             |
| 40        | 0.13                                |                                                                      |                             |
| 50        | 0.28                                |                                                                      |                             |
| 110       | 0.28                                |                                                                      | Begin                       |
| 210       | 0.28                                | Begin                                                                | End                         |
| 220       | 0.61                                |                                                                      |                             |
| 460       | 0.61                                |                                                                      |                             |
| 470       | 0.28                                |                                                                      |                             |
| 560       | 0.28                                | End                                                                  |                             |
| 570       | -1.67                               |                                                                      |                             |
| 580       | -1.67                               |                                                                      |                             |
| 590       | 0.93                                |                                                                      |                             |
| 600       | 0.13                                |                                                                      |                             |

<sup>a</sup>See the waveform graphs in **Fig. 10**.

mix of quaternary/tertiary ammonium groups. This mix of functionalized groups has a controlled ratio, resulting in excellent selectivity for the separation of amino acids. The small particle size of the anion-exchange layer exhibits adequate mass transfer characteristics enabling high chromatographic peak efficiency. The highly crosslinked core of the AminoPac PA10 packing makes it possible to use organic solvents to facilitate column cleanup.

2. Use chemicals exactly as specified by catalog number and supplier. Consult with Dionex should one or more of the specified reagents be not available in your geographical area.
3. Minimize any extraneous contamination of eluents. For example, a trace of an ion-pairing agent introduced into the eluent from a “shared” filtration apparatus will cause interference with some of the amino acid peaks. Dedicate glassware, pipets, filtration apparatus for exclusive use in preparation of AAA eluents only. Wear disposable powder-free gloves whenever preparing or refilling eluents.
4. Minimize the level of carbonate introduced into the eluents during preparation.
5. Avoid bacterial contamination of eluent bottles and tubing. The bacterial contamination is minimized by wearing disposable, powder-free gloves, keeping containers closed whenever possible, and by ultrafiltration (filter pore size < 0.2 μm). Use ultrafiltration as indicated in the instructions for preparing each of the three mobile phases. Microorganisms, if present in the system, produce amino acids, causing elevated background levels and spurious peaks.

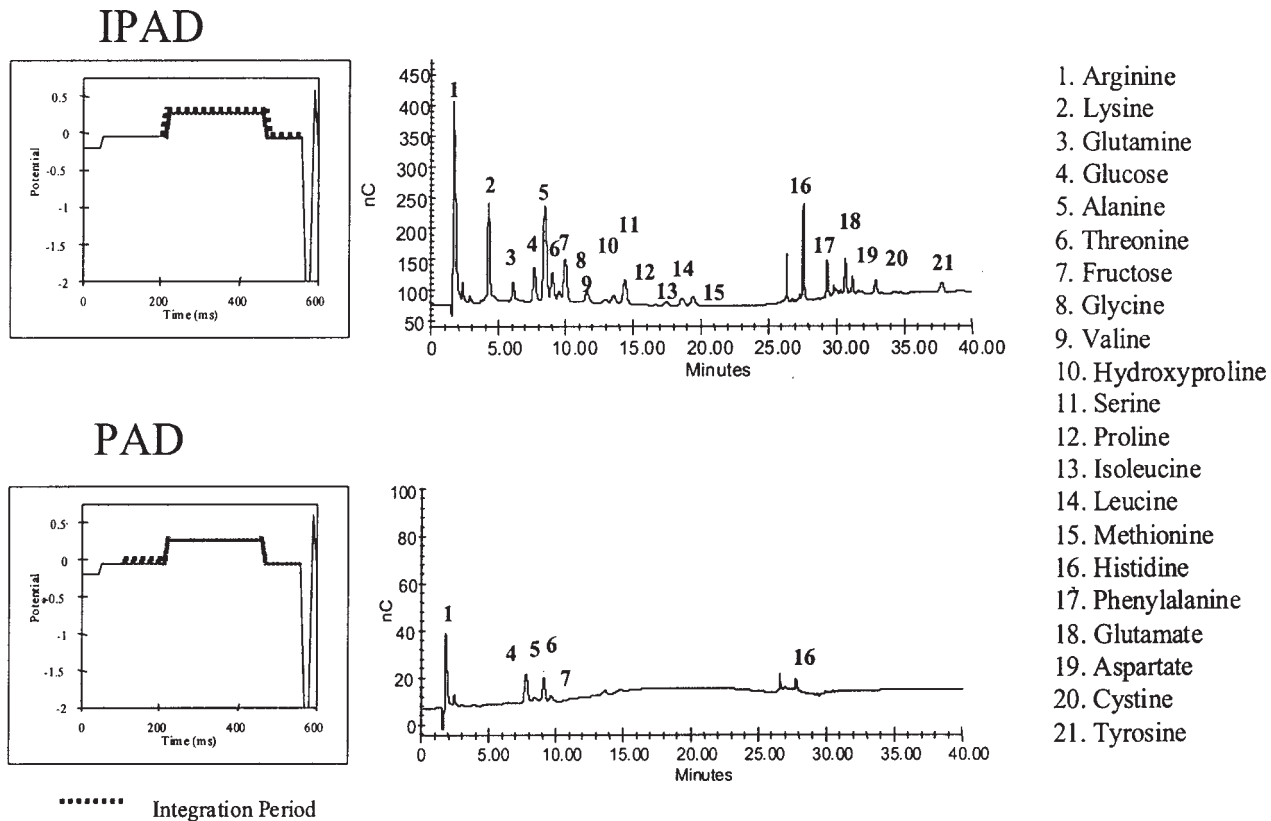


Fig. 8. Selective IPAD detection applied to a 1:250 diluted cell culture sample. The lower trace obtained with the hydroxy specific detection mode (*see Note 28*) helps to identify an unknown peak (Peak #7) as fructose. The identification of fructose in the upper chromatogram is difficult since it coelutes closely with glycine.

6. Remember, atmospheric carbon dioxide adsorbs even into pure water, albeit at much lower levels than in alkaline solutions. Minimize the time the surface of the water comes in contact with the atmosphere.
7. Follow the generally accepted hydrolysis procedures. The present method works with the protocols (**13–15**) listed below (**Fig. 5**):
  - a. Monosaccharide analysis: 2 M TFA (31  $\mu$ L of 13 N TFA plus 169  $\mu$ L of water, 4–5 h, 100°C.
  - b. Amino sugar analysis: 6 M HCl 4–5 h, 100°C.
  - c. Amino acid analysis: 6 M HCl 18–24 h, 110°C.
  - d. Amino acid analysis: 4.0 M MSA 24 h, 110°C.
  - e. Tryptophan analysis: 4.2 M NaOH 24 h, 110°C.
8. The HCl hydrolysates usually show less interference compared to other protocols. Hydrolysis by MSA is preferred by many workers for analyzing plant related materials or animal tissue. Whenever using MSA hydrolysis for cellulose (plants) or carbohydrate (processed meats) containing materials, it is advisable to employ a gradient resolving glucose from alanine (**Subheading 3.5., step 9**). Unlike the ninyhydrin/cation approach, the IPAD technique is very compatible with NaOH hydrolysates. Neither evaporation nor neutralization is required. We have found that an accurate analysis of tryptophan is easier with NaOH hydrolysis, especially for plants and animal tissues (**Fig. 5**).
9. Always check blanks before reusing the hydrolysis tubes. Chemical cleaning (chromic acid, commercial cleaning formulations) does not always work. Even the most reliable procedure (heating the tubes to 500°C in a muffle furnace) may have to be repeated to obtain blanks free of carryover impurities.
10. HCl hydrolysates containing high enough concentrations of amino acids (0.1–2  $\mu$ g/200  $\mu$ L) do not require an evaporative centrifugation step. They can be injected after a 100- or 1000-fold dilution with the norleucine/azide diluent (**Subheading 2.3., step 5**). All hydrolysates in sodium hydroxide can be injected directly. Neither a neutralization nor evaporation is required. All MSA hydrolysates should be diluted at least 1000-fold.
11. Sample pH adjustments are not required with the present chromatographic method.
12. The standard gradient runs and all AminoPac PA10 chromatographic runs can be subdivided into four parts:
  - a. First, amino acids are eluted by the hydroxide eluent only. This is started by an isocratic composition and finished off by a gradient. In the **Fig. 1** chromatogram, this part is represented by a group of peaks between arginine and norleucine.
  - b. To elute the second group of peaks (histidine through tryptophan) requires a more strongly eluting species, the acetate anion. The sharp spike (not numbered) between norleucine and histidine peak is always present and indicates the point where the conversion of the anion exchanger from hydroxide to acetate is complete. The spike itself consists of hydroxide ions and illustrates the focusing effect of the acetate anion on all the other ions still retained on the column at the end of the first part of the chromatographic run.

The second part of the run is complete when the acetate concentration has dropped to zero, and for 2 min the hydroxide concentration is increased to 200 mM.

- c. This third step accelerates the conversion of the anion exchange groups back to the hydroxide form, it also effects the removal of traces of carbonate brought on the column as an impurity in E1, E2, or E3.
  - d. In the fourth step, the column is readjusted by pumping of initial gradient composition of sodium hydroxide.
13. The separation temperature must be controlled within  $\pm 1^\circ\text{C}$ . The prescribed temperature is  $30^\circ\text{C}$  for most of the existing methods. The comparison of four chromatograms obtained between  $30$  and  $45^\circ\text{C}$  (**Fig. 9**) shows that hydroxyproline, serine, and methionine are most affected by temperature changes. Only a small effect is discernible on the amino sugars and threonine. Also, all the temperature-induced shifts are within the “hydroxide part” of the chromatogram. The position of the peaks eluted by acetate is virtually unchanged.
  14. The position of serine relative to proline (**Fig. 9**) is a good indication of whether or not the calibration of the column thermostat is correct. At lower temperatures, that peak closely coelutes with proline and at a temperature that may be too high it moves closer to valine.
  15. Do not polish the Dionex AAA Gold electrode. Polishing the gold electrodes with  $1\ \mu\text{m}$  alumina is a standard procedure in connection with the analysis of carbohydrates by PAD. Since IPAD is carried out at higher potentials than PAD, the microscopic roughness resulting from polishing leads to excessive background levels. Alternative procedures such as electropolishing and chemical cleaning are described in the Dionex AAA Direct System Manual.
  16. Do not allow the pH and Ag/AgCl combination electrode to dry out. The electrode dries out easily and is irreversibly damaged if left in a disconnected detector cell or if stored dry. Follow the storage instructions in the Dionex ED40 manual.
  17. The ED40 reference electrode should be checked against “known good” reference electrodes in regular intervals. A known good reference electrode is, for example, another unused reference electrode of the same type that was properly stored in a  $3\ \text{M}$  KCl solution. Discard any reference electrode if its potential differs by more than 50 mV from the known good reference. Use only a high impedance voltmeter (e.g., a laboratory pH meter) to check reference potentials.
  18. Sample tray cooling is available with the Dionex AS50 autosamplers.
  19. The azide-containing diluent solution stabilizes samples and standards for at least three days. Overnight and over-the-weekend runs are possible even without a sample tray cooling.
  20. Use low-volume autosampler vials ( $V = 300\ \mu\text{L}$ , PN 500114, Sun International, Wilmington, DE) for limited volumes of undiluted samples ( $< 0.5\ \text{mL}$ ). Use standard autosampler vials made of glass ( $V = 1.5\ \text{mL}$ , Dionex 055427) for larger volumes ( $> 0.5\ \text{mL}$ ).

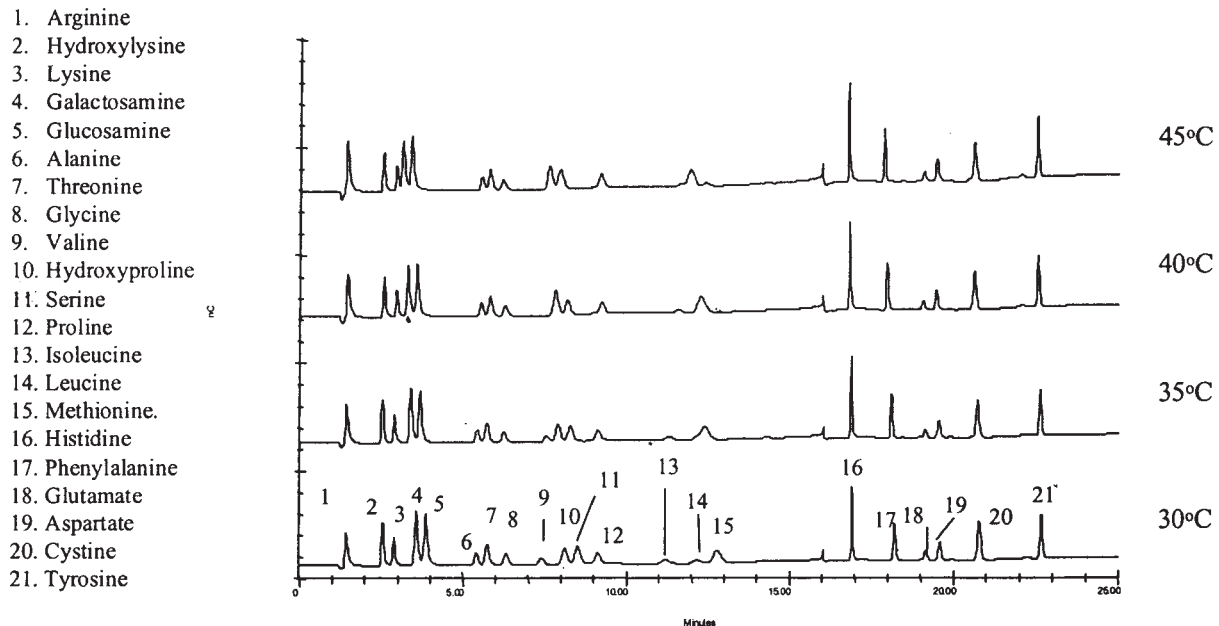


Fig. 9. Influence of column temperature on the separation of amino acids. Note the changes of relative positions of methionine vs leucine and hydroxyproline and serine.

21. The column temperature of 35°C has been found useful for separating amino acids in the presence of methionine sulfone and cysteic acid.
22. Evaluation of linearity in the range of 1 to 1000 pmol (**5**) showed the following results: Calibration plots are linear across three orders of magnitude for Ala, Thr, Gly, Val, Ser, Pro, Ile, Leu, Met, Phe, Glu, Asp, Cys, and Tyr. Calibration plots are linear over 2.25 orders of magnitude for Arg, Lys, Asn, Gln, and His.
23. The following limits of detection (3x noise in picomoles) were determined by injecting 10 pmol of each compound under the conditions of **Fig. 2**: Arg 0.15, Lys 0.21, Gln 0.48, Asn 0.35, Ala 0.72, Thr 0.40, Gly 0.62, Val 1.48, Ser 0.49, Pro 1.29, Ile 2.84, Leu 4.42, Met 1.12, His 0.11, Phe 0.36, Glu 0.78, Asp 0.51 (Cys) 2 0.26, Tyr 0.44.
24. If a mixture of cysteine and cystine is injected, only a single peak is observed. Single component standards of cysteine or cystine each yield a peak at the identical retention time.
25. Analytical results obtained by the IPAD method are in a good agreement with the ninhydrin/cation-exchange technique for a large number of protein and peptides. Two reports comparing the results for ribonuclease A and collagen are available in the literature (**5,12**).
26. The possibility of monitoring amino acids and glucose with an On-Line Chromatographic Analyzer (Dionex DX-800) has been demonstrated. Fully automatic sampling is achieved using a tangential membrane filtration device as a sterile barrier. All connecting tubing between the fermentor and the filter and that leading from the filter to the On-Line analyzer has to be carefully sterilized prior to use.
27. Glucose and amino acids can be quantified reliably by using the gradient method in **Table 4**. We have found it possible to achieve an automatic peak detection of alanine by the PeakNet software up to the glucose/alanine molar ratio of 200 to 1. Should the sample contain an even higher excess of glucose and/or additional carbohydrate components, it may become necessary to use the Group Separation Gradient from **Table 3**. Using that gradient, we have verified experimentally that automatic peak detection and quantitation of alanine are possible up to a 1000-fold molar excess of glucose.
28. IPAD detection of amino acids and carbohydrates relies on the presence of suitable catalysts on the gold electrode surface. While the detection of hydroxide groups in carbohydrates requires the presence of hydroxide radicals, a monolayer of gold oxide is necessary to make the oxidation of amino groups of amino acids possible. In most applications, both types of catalytic surfaces are sequentially generated during each cycle of the IPAD detection waveform (*see Fig. 4B*). A typical IPAD method detects amino acids by integrating over the entire length of step E3. The sugars are detected by adding the entire length of step E4 to the period of current integration. The codetection of carbohydrates is, to a degree, a side benefit; the main purpose of the “dual-step” integration is to eliminate any contribution of gold oxide formation to the detection signal. Oxidation and reduction currents cancel each other out during the E3 and E4 integration (**11**).



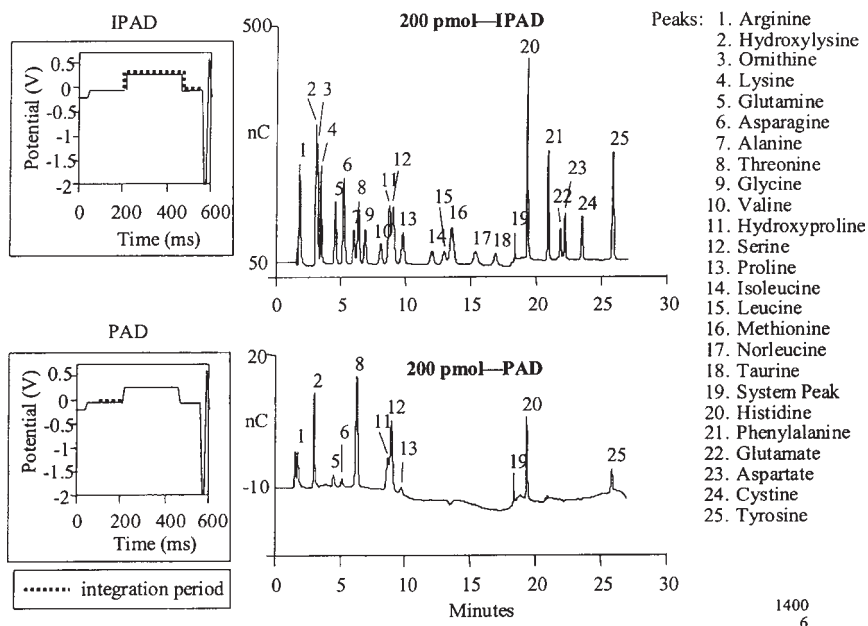


Fig. 10. Changes of selectivity in standard chromatograms with two different waveforms. **Note:** the coeluting hydroxylysine is resolved from ornithine by the “hydroxy-selective waveform” in the lower chromatogram.

If the detection waveform is modified to expand the length of step E2 as shown in **Fig. 10**, it is possible to design a current-integration method that is sensitive for carbohydrates only. This relatively minor modification of the detection conditions enables us to choose between the simultaneous detection of amino acids and carbohydrates (**Fig. 10**, upper trace) or the more selective detection of only those molecules carrying hydroxy functional groups (**Fig. 10**, lower trace). The latter include not only sugar alcohols, amino sugars, and mono- and disaccharides, but also hydroxyamino acids.

The hydroxy-specific current-integration also detects the basic amino acids (arginine, lysine, glutamine, asparagine, and histidine, **Fig. 10**, lower trace) if they are present at high enough concentrations.

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## **Ion-Pair Chromatography for Identification of Picomolar-Order Protein on a PVDF Membrane**

**Noriko Shindo, Tsutomu Fujimura, Saiko Kazuno,  
and Kimie Murayama**

### **1. Introduction**

There are several highly sensitive methods for identifying proteins: N-terminal sequencing, peptide mapping, and Western blotting, which are based on partial information about the structure of the protein. The amino acid composition of a protein is reflected in the primary structure of the whole protein, and this information is highly conserved.

Recently, we have reported a highly sensitive method of amino acid analysis — 6-aminoquinolyl-carbamyl (AQC)-amino acid analysis — using ion-pair chromatography (1) and used it to identify multidrug resistant protein 1 (MDR1) of mouse leukemia P388 cells on a polyvinylidene fluoride (PVDF) membrane with the aid of World Wide Web (WWW)-accessible tools (2).

MDR1 could not be confirmed by immunostaining with a commercially available murine monoclonal antibody (MAbC219) because of low specificity for this protein. Membrane proteins from the cells were separated by 1-D sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), and blotted onto a polyvinylidene difluoride (PVDF) membrane. The 160-kDa protein band was excised and hydrolyzed in a vapor phase, then subjected to *in situ* AQC-derivatization without extraction of the amino acids. The amino acid composition of the protein was obtained using a highly sensitive AQC-amino acid analyzer employing ion-pair chromatography, and the protein was identified on the basis of its amino acid composition using the AACompIdent tool in ExPASy and PROPSEARCH in EMBL via the WWW. The methods employed are described in this chapter.

The most important consideration for such highly sensitive analysis is to minimize or decrease the extent of contamination from the environment, and sample loss. This was done by sample manipulation in a hand-made clean hood with gloves as far as possible, and *in situ* AQC-derivatization without extraction of amino acids from the PVDF membrane.

## 2. Materials

### 2.1. Reagents

**Table 1** shows the reagents and consumables employed.

### 2.2. Apparatus

1. Clean hood: This was made from a metal frame ( $60 \times 60 \times 60$  cm), the top, back, and sides of which were covered with clean transparent vinyl sheets and taped. The front sheet was hung to allow manipulation, and on its outer surface, a glass plate ( $10 \times 20$  cm) was attached with a glued tape to facilitate a clear view (*see Note 3*). Before and after the experiment, the inside of the hood was sterilized with ethanol for disinfection (*see Note 4*).
2. Hydrolysis vessel: JASCO Co. (<http://www.jasco.co.jp/>) code no. R154 (*see Fig. 1*). Another commercially available hydrolysis vessel is described by Wilkins et al. (3).
3. Shimadzu high-performance liquid chromatograph LC-10A system: Column oven CTO-10AC; Auto sample injector SIL-10AXL with sample cooler; Fluorescence detector RF-10AXL; LC-Workstation Class-LC10.
4. Other apparatus and tools are shown in **Table 2**.

## 3. Methods

**Fig. 2** shows a flow chart of the manipulation under the clean hood, gloves, and arm covers.

### 3.1. SDS-PAGE and PVDF Membrane Blotting

1. Prepare a plasma-rich membrane fraction of leukemia cells ( $1 \times 10^8$  cells) by sonication ( $2 \times 10$  s), followed by centrifugation ( $100,000g$  for 1 h at  $4^\circ\text{C}$ ), and suspend in 1 mL of Dulbecco's phosphate-buffered saline (PBS) by ultrasonication ( $2 \times 10$  s).
2. Determine the protein concentration using BCA protein assay reagent (product no. 23225, Pierce, Rockford, IL).
3. Dissolve one aliquot (100  $\mu\text{g}$  protein) in an equal volume of sample buffer, and load onto a 1D SDS-PAGE gel.
4. Use 10  $\mu\text{g}$  of bovine serum albumin (BSA) as a calibration protein for determining the composition of the amino acids in this experiment (*see Note 5*).
5. Electrophorese at 5 mA for 15 h at room temperature.
6. Perform the transfer onto the PVDF membrane using the tank blotting method and 10 mM CAPS buffer (*see Notes 6 and 7*). Wash the membrane with MeOH

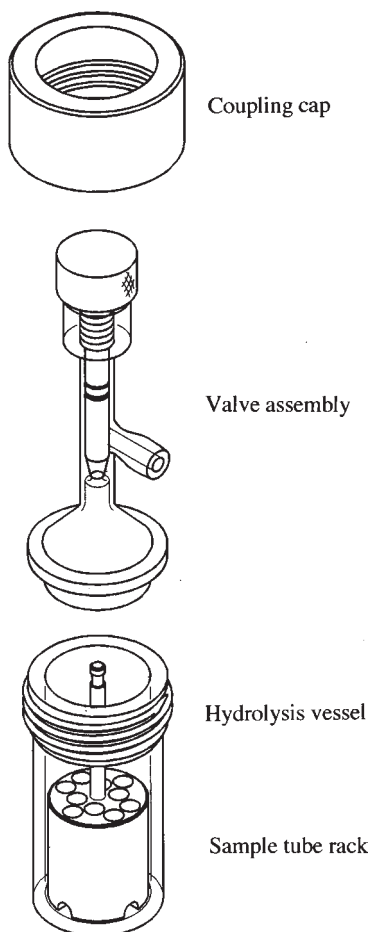


Fig. 1. The design of the hydrolysis vessel. It is composed of a glass container, a valve assembly, a coupling cap, and a Teflon sample tube rack.

before use, and then soak in the transfer buffer for a few minutes. Equilibrate the gel in the buffer for 5 min.

7. Carry out the electrotransfer at 4°C for 1.5 h at 50 V.
8. Stain the membrane with 0.1% Coomassie brilliant blue R-350 for 5 min.
9. Destain quickly five times in 50% MeOH (each 30 s) in order to prevent protein loss, and the dry membrane in a refrigerator.
10. Determine the molecular weight (MW) using the image analyzer.

### 3.2. Acid Hydrolysis

1. Cut out the protein band on the PVDF membrane with a knife or a surgical blade, and cut this into 2 or 3 pieces, if necessary (*see Note 8*). To prevent the band from spreading, do not use a wet blade. Also, cut out a piece of blank membrane from

**Table 1**  
**Reagents**

| Method                 | Solution                                                                                   | Specification               | Maker                                               | Product no. |
|------------------------|--------------------------------------------------------------------------------------------|-----------------------------|-----------------------------------------------------|-------------|
| <b>General Use</b>     |                                                                                            |                             |                                                     |             |
|                        | fresh Milli-Q Water                                                                        | Analytical grade            | PE Biosystems, CA, USA                              |             |
|                        | Acetonitrile (MeCN)                                                                        | HPLC grade                  | Wako Pure Chemical Industries Ltd.,<br>Osaka, Japan | 015-08633   |
|                        | Methanol (MeOH)                                                                            | HPLC grade                  | Wako Pure Chemical Industries                       | 052-03343   |
|                        | Ethanol for disinfection                                                                   | 76.9–81.4%                  | Wako Pure Chemical Industries                       | 328-00035   |
| <b>SDS-PAGE</b>        |                                                                                            |                             |                                                     |             |
| 1D-PAGE                | 4% for stacking gel, 6.2% for separation gel                                               | 13.5 × 15 × 0.1 cm          |                                                     |             |
| Buffer                 | 125 mM Tris-HCl, pH 6.8                                                                    |                             |                                                     |             |
| Sample buffer          | The above buffer containing 4% SDS, 10%<br>2-mercaptoethanol, and 0.0025% BPB              |                             |                                                     |             |
| <b>Electroblotting</b> |                                                                                            |                             |                                                     |             |
| Transfer buffer        | 10 mM CAPS buffer (3-[Cyclohexylamino]-<br>1-propanesulfonic acid) in 10% MeOH,<br>pH 11.0 |                             | Dojindo Laboratories,<br>Kumamoto, Japan            | 343-00484   |
| Staining solution      | 0.1% CBB (Coomassie brilliant blue [CBB]<br>R-350) in 30% MeOH, 10% acetic acid            | PhastGel BlueR              | Amersham Pharmacia Biotech,<br>Uppsala, Sweden      | 17-0518-01  |
| Destaining solution    | 50% MeOH                                                                                   |                             |                                                     |             |
| <b>Hydrolysis</b>      |                                                                                            |                             |                                                     |             |
| Acid solution          | 6 N HCl containing 2% thioglycolic acid<br>or 2.5% phenol                                  |                             |                                                     | 17-0518-01  |
|                        | Hydrochloric acid (HCl)                                                                    | Amino acid analysis grade   | Wako Pure Chemical Industries Ltd.                  | 086-03925   |
|                        | Phenol, crystal                                                                            | for Nucleic acid extraction | Wako Pure Chemical Industries Ltd.                  | 168-1272    |
|                        | Thioglycolic acid                                                                          | Special grade               | Wako Pure Chemical Industries Ltd.                  | 204-01023   |

**AQC-Derivatization**

|             |                                                                                       |                         |                              |        |
|-------------|---------------------------------------------------------------------------------------|-------------------------|------------------------------|--------|
| AHC Reagent | 10 mM AHC [6-Aminoquinolyl-N-hydroxysuccinimidyl carbamate] in MeCN ( <b>Note 1</b> ) | AccQ-Fluor™ Reagent Kit | Waters Co., Millford, MA     | 52880  |
| Solution    | 0.2 M Borate buffer pH8.8 ( <b>Note 1</b> )                                           | for Electrophoresis     | Nacalai Tesque, Kyoto, Japan | 052-18 |

**Ion-Pair****Chromatography**

|                     |                                                                                                                                                            |                             |                                     |           |
|---------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------|-------------------------------------|-----------|
| Column              | Cosmosil 5C8-MS                                                                                                                                            | 4.6 mm ID × 150 mm          | Nacalai Tesque, Kyoto, Japan        | 381-55    |
| Elution buffer A    | 30 mM Phosphate buffer pH 7.2 containing 5 mM TBA-Cl<br>Tetrabutylammonium chloride (TBA-Cl)                                                               | for ion-pair chromatography | Fluka Chemie AG, Buchs, Switzerland | 86852     |
| Elution buffer B    | The solution pH 7.2, which are mixed 260 mL of MeCN, 65 mL of 30 mM phosphate buffer and 175 mL of 30 mM KH <sub>2</sub> PO <sub>4</sub> ( <b>Note 2</b> ) |                             |                                     |           |
| Amino acid solution | Wako amino acid standard solution                                                                                                                          | type H (2.5 μmol/mL, each)  | Wako Pure Chemical Industries Ltd.  | 019-08393 |

**Table 2**  
**Apparatus and Tools**

| Name                                                    | Specification                                                        | Maker                                        | Product no. |
|---------------------------------------------------------|----------------------------------------------------------------------|----------------------------------------------|-------------|
| <b>General Use</b>                                      |                                                                      |                                              |             |
| Flush Mixer                                             |                                                                      |                                              |             |
| Oven                                                    | set at 300°C for pyrolyzation of glass wares                         |                                              |             |
| Microcapillary tips                                     | 83-mm length (0.5–200 µL)                                            | Sorenson™ BioScience, Inc., UT               | 13791       |
| Gloves                                                  | powder free latex gloves, disposable                                 | Microflex Medical Corp., Malaysia            | EV2050      |
| Disposable arm cover                                    |                                                                      | Iuchiseieido, Osaka, Japan                   | 88-1073-01  |
| <b>Electroblotting</b>                                  |                                                                      |                                              |             |
| Minitransblotting apparatus                             | or equivalent apparatus                                              | Nippon Eido, Co., Ltd., Tokyo, Japan         | NC10115     |
| Polyvinylidene fluoride (PVDF) membrane                 | ProBlott                                                             | PE Biosystems, CA                            |             |
| Image Analyzer                                          | analysis program: RFLP                                               | Scanalytics, MA                              |             |
| <b>Hydrolysis</b>                                       |                                                                      |                                              |             |
| Hydrolysis vessel                                       | (see Fig. 1)                                                         | JASCO Co., Hachioji, Japan                   | R154        |
| Hydrolysis tube and silicon plug or an inner glass tube | 5.2 mm ID × 46.2 mm                                                  | JASCO Co., Hachioji, Japan                   | R158, R159  |
| Aluminum block                                          |                                                                      | JASCO Co., Hachioji, Japan                   | R157        |
| Hydrolysis furnace                                      | set at 110°C.                                                        |                                              |             |
| Glass plate                                             | for excising bands. 10 cm × 10 cm.                                   |                                              |             |
| Dry seal vacuum desiccator                              | grease-free                                                          | Wheaton, Millville, NJ                       | 365884      |
| 0.45 µm Filter cartridge                                | Ekicrodisc 13                                                        | Gelman Science Japan, Tokyo, Japan           | E131        |
| Vacuum pump 1                                           | Pumping speed 6L/min<br>(suction of solution, with a glass trap)     | Sinku Kiko, Yokohama, Japan                  | G-5         |
| Glass trap                                              | (see Fig. 2)                                                         |                                              |             |
| Vacuum pump 2                                           | Pumping speed more than 100 L/min<br>equipped with a solid-NaOH trap | Sato Vacuum Machinery Ind. Co., Ltd. Japan   | DW-180      |
| Surgical blade or knife                                 | for excising protein band.                                           | Feather Safety Razor Co., Ltd., Osaka, Japan |             |
| <b>AQC-Derivatization</b>                               |                                                                      |                                              |             |
| Ultrasonic bath                                         | add fresh Milli-Q when use.                                          | Branson Ultrasonic Co., CT                   | B-12        |
| Heating bath                                            | set at 55°C.                                                         | Riko-Kagaku Sangyo Co., Ltd., Japan          | MH-10D      |



- an area on which sample proteins have not run on. Place the standard, sample, and blank membrane fragments in separate glass tubes.
2. Wash the membrane pieces with 200  $\mu\text{L}$  of 50% MeCN (or MeOH) on a mixer. Immediately suck up the washing solution with an aspirator, e.g., a handmade aspirator using pump 1. Excessive exposure to MeCN leads to sample loss (*see Note 9*).
  3. While the membranes are wet, wash them with 200  $\mu\text{L}$  of 0.01 *N* HCl, and then aspirate the solution. This washing step is done 2 or 3 times, and finally the solution takes place in 50% MeCN.
  4. Dry the membranes in a vacuum desiccator for 10 min, and then slowly introduce the air into the desiccator through a 0.45- $\mu\text{m}$  filter cartridge.
  5. Add 6 *N* HCl (containing 2.5% phenol or 2% thioglycolic acid) to the hydrolysis vessels.
  6. Place the tube in a Teflon sample tube rack, which is then placed into the vessel. Tightly close the cap and the valve of the vessel.
  7. Move the vessel out of the hood, evacuate for 2 min, and place in a preheated aluminum block (110°C).
  8. Carry out the hydrolysis at 110°C overnight.
  9. Connect the valve of the hot vessel from the heating block to pump 2, and evacuate the residual 6 *N* HCl (roughly 5 min).

### 3.3. In Situ Derivatization of AQC-Amino Acids

1. Break vacuum of the vessel through a 0.45- $\mu\text{m}$  filter cartridge.
2. Put the sample tube rack in a vacuum desiccator, and the membrane are completely dried with pump 1 for 10 min (*see Note 10*).
3. Moisten the membranes by dropping 10  $\mu\text{L}$  of neat MeCN directly onto them, followed by 10  $\mu\text{L}$  of 0.01 *N* HCl, and 10  $\mu\text{L}$  of fresh Milli-Q water. AQC-derivatization of the standard amino acid mixture is done by adding 10  $\mu\text{L}$  of standard solution in 0.01 *N* HCl instead of 0.01 *N* HCl for the membrane sample.
4. Add 50  $\mu\text{L}$  of 0.2 *M* borate buffer and 20  $\mu\text{L}$  of 10 *mM* AHC reagent, and leave the solution to stand for 1 min. Remove the sample rack from the hood and heat at 55°C for 10 min (*see Note 11*).
5. Transfer the derivatized solution to an autosampler plastic tube before HPLC analysis to avoid adsorption of amino acids onto the membrane and the wall of the glass tube.
6. Dilute 10  $\mu\text{L}$  of the solution 10-fold with 0.2 *M* borate buffer.

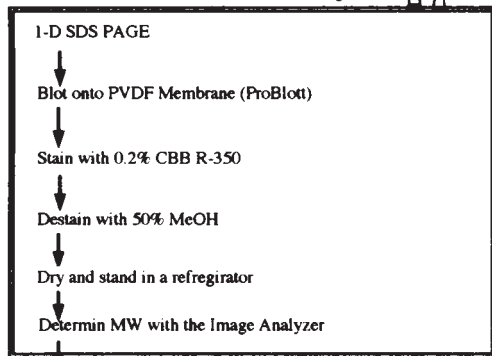
### 3.4. AQC-Amino Acid Analysis

1. AQC-amino acids are separated by ion-pair chromatography using TBA-Cl.
2. The elution program is shown in **Table 3** (*see Notes 12 and 13*). The column temperature is kept at 30°C.
3. The elution profile of standard AQC-amino acids (1 pmol) is shown in **Fig. 3**.
4. Prepare the standard AQC-amino acid solution to derivatize 10  $\mu\text{L}$  of 10 pmol/ $\mu\text{L}$  amino acid standard mixture, and store at -40°C (*see Note 14*). Dilute the AQC-

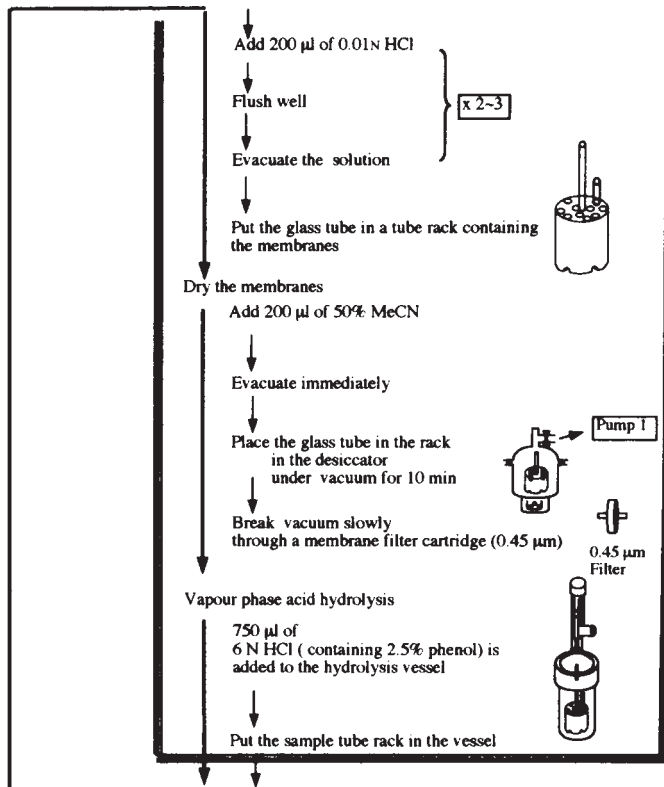
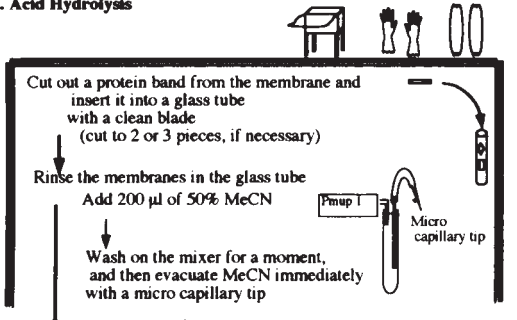


(see Note 3,4)

**3.1. SDS-PAGE and PVDF Membrane Blotting**



**3.2. Acid Hydrolysis**



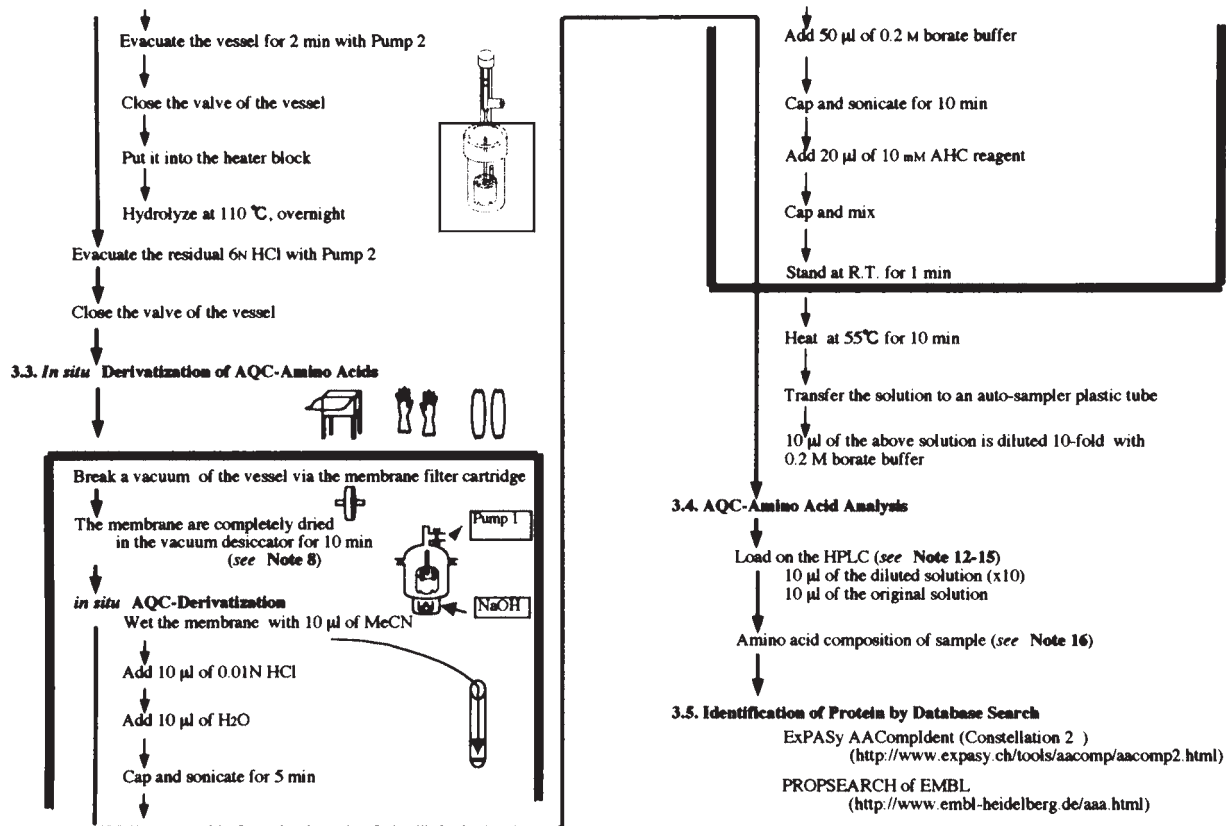


Fig. 2. Flowchart of manipulation under the hood with gloves and arm covers.

**Table 3**  
**Elution Program**

| Time (min) | Concentration of buffer B (%) |
|------------|-------------------------------|
| 0.00       | 11.5                          |
| 3.00       | 11.5                          |
| 3.01       | 16.3                          |
| 63.00      | 62.5                          |
| 63.01      | 100.0                         |
| 71.00      | 100.0                         |
| 71.01      | 11.5                          |
| 91.00      | 11.5                          |

Buffer A: 30 mM phosphate buffer (pH 7.2), 5 mM TBA  
 Buffer B: 52% MeCN, 48% of 30 mM phosphate buffer  
 Flow rate: 0.5 mL/min  
 Column Temperature: 30C, isothermal  
 Detection: Ex 250 nm, Em 395 nm

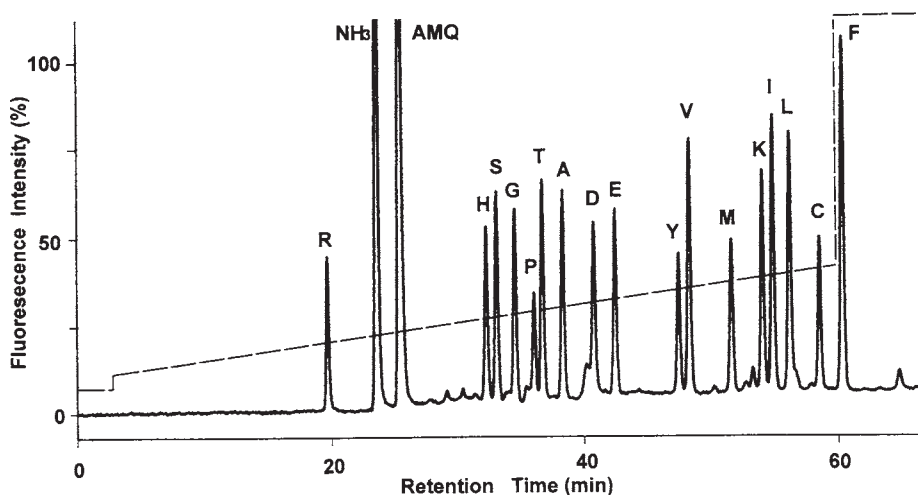


Fig. 3. Chromatogram of 1 pmol of the AQC-amino acid standard. AQC- amino acids are shown as one-letter code.

standard solution and sample 10-fold with 0.2 M borate buffer (pH 8.8), and inject 10- $\mu$ L aliquots into the column (see **Note 15**).

- Figure 4** shows the 1-D SDS PAGE pattern for P388/S and P388/DOX cells (**A**) and the AQC-amino acid chromatogram for the 160-kDa protein band of P388/DOX cells, showing drug resistance for doxorubicin (**B**).

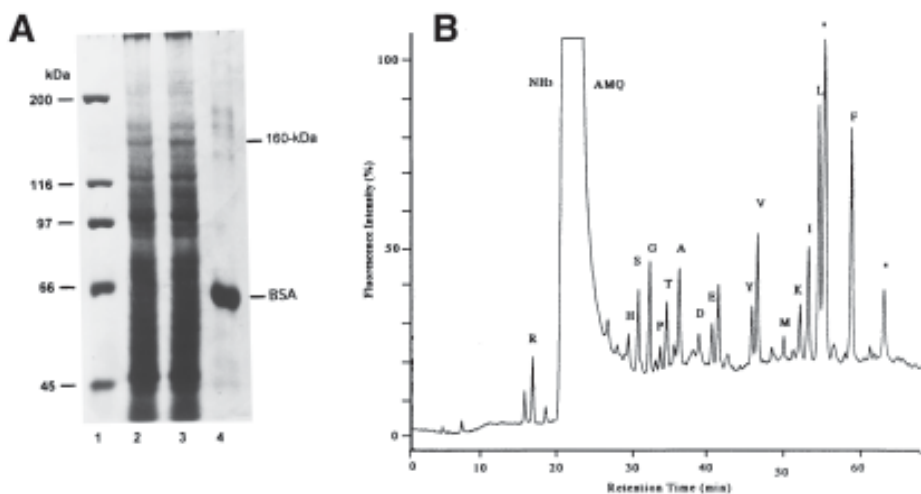


Fig. 4. (A) 1-D SDS-PAGE of P388/S and P388/DOX cells. Lane 1: Molecular weight standards, Lane 2: P388/S (control), Lane 3: P388/DOX, and Lane 4: BSA as a calibration standard. (B) AQC-amino acid chromatogram for the 160-kDa protein band of P388/DOX cells. \*: Reagent-related peak.

6. Confirm the peak identification and peak area of each amino acid.
7. Calculate the concentration of the 16 amino acids against the standard amino acid mixture, and then export the data to a Microsoft Excel program.
8. Calculate the amino acid composition as the picomolar composition (%) in table form for the subsequent database search.
9. **Table 4** was prepared for the database search using ExpASY. It contained the amount of amino acid (pmol) and the amino acid compositions of P388/S, P388/DOX, and BSA that was used to correct the errors resulting from the hydrolysis (see **Note 16**).
10. Monitor the contamination of the amino acids using the membrane blank chromatogram.

### 3.5. Identification of Protein Using WWW Tools

1. Protein identification is done using its amino acid composition and MW and/or pI. These values are obtained from image analysis of the CBB-stained membrane. Protein identification is done using AACompIdent of the ExpASY molecular biology server (Constellation 2, for 16 amino acids) (<http://www.expasy.ch/tools/aacomp/aacomp2.html>) or PROPSEARCH of EMBL (<http://www.embl-heidelberg.de/aaa.html>).
2. The database search with PROPSEARCH is done using the composition corrected by the recovery of calibrated protein. The MW was determined to be 160-kDa from the blotted 1-D SDS-PAGE membrane using an image analyzer.

**Table 4**  
**Amino Acid Compositions of 160-kDa Protein in P388/S and P388/DOX Cells**

| Amino Acid | Standard (BSA) |                     | P388/S        |                     | P388/DOX      |                     |
|------------|----------------|---------------------|---------------|---------------------|---------------|---------------------|
|            | Experimental   |                     | Experimental  |                     | Experimental  |                     |
|            | Amount (pmol)  | Composition (pmol%) | Amount (pmol) | Composition (pmol%) | Amount (pmol) | Composition (pmol%) |
| Arg        | 4.22           | 5.16                | 0.27          | 9.04                | 0.17          | 5.37%               |
| His        | 2.66           | 3.26                | 0.05          | 1.68                | 0.06          | 1.91%               |
| Ser        | 4.26           | 5.21                | 0.21          | 7.10                | 0.22          | 7.03%               |
| Gly        | 3.36           | 4.11                | 0.27          | 8.90                | 0.30          | 9.71%               |
| Pro        | 3.74           | 4.57                | 0.09          | 3.10                | 0.09          | 2.90%               |
| Thr        | 5.05           | 6.18                | 0.17          | 5.70                | 0.14          | 4.69%               |
| Ala        | 6.91           | 8.45                | 0.20          | 6.66                | 0.19          | 6.11%               |
| Asp        | 4.37           | 5.35                | 0.07          | 2.51                | 0.11          | 3.50%               |
| Glu        | 6.79           | 8.31                | 0.18          | 6.10                | 0.18          | 5.84%               |
| Tyr        | 5.13           | 6.27                | 0.17          | 5.65                | 0.17          | 5.65%               |
| Val        | 6.78           | 8.29                | 0.26          | 8.61                | 0.30          | 9.85%               |
| Met        | 0.60           | 0.74                | 0.08          | 2.83                | 0.06          | 1.88%               |
| Lys        | 5.31           | 6.50                | 0.11          | 3.58                | 0.14          | 4.46%               |
| Ile        | 2.01           | 2.46                | 0.16          | 5.49                | 0.24          | 7.72%               |
| Leu        | 12.93          | 15.82               | 0.42          | 14.07               | 0.44          | 14.35%              |
| Phe        | 7.63           | 9.33                | 0.27          | 8.99                | 0.28          | 9.02%               |
| Total      | 81.74          | 100.00              | 2.98          | 100.00              | 3.08          | 100.00%             |

3. The results from database search with ExPASy are shown in **Fig. 5**. This shows the expression of MDR 1 in P388/DOX cells (score 14).

#### 4. Notes

1. AccQ-Fluor™ Diluent and Borate Buffer in the kit were not used.
2. Elution buffer B in the original method (80% MeCN) was modified, and the elution program was changed to obtain the same separation as before, because of column maintenance.
3. A similar apparatus is commercially available. Our first clean hood used a computer rack as a frame.
4. It is important to prevent outside air from entering the hood. Therefore, it may be useful to spray the outside of the hood with ethanol for disinfection before use.
5. Usually, one of the SDS-PAGE MW standard bands is used as a calibration protein, but some of the prestained MW standards are not useful because they are largely modified from the original proteins.
6. The sponge pad, two filter papers, PVDF membrane, gel, two filter papers, and the sponge pad are wetted in the 10 mM CAPS buffer and place one after another

## A P388/S

The SWISS-PROT entries having pI and Mw values in the specified range for the species MOUSE and the specified keyword:

| Rank | Score | Protein    | pI   | Mw     | Description                      |
|------|-------|------------|------|--------|----------------------------------|
| 1    | 34    | PHYD_ARATH | 5.81 | 129302 | PHYTOCHROME D.                   |
| 2    | 34    | PHYB_ARATH | 5.62 | 129331 | PHYTOCHROME B.                   |
| 3    | 35    | CYA9_MOUSE | 6.85 | 150954 | ADENYLATE CYCLASE, TYPE IX (EC   |
| 4    | 35    | KPB1_MOUSE | 5.55 | 138793 | PHOSPHORYLASE B KINASE ALPHA     |
| 5    | 36    | NOS2_MOUSE | 7.76 | 130575 | NITRIC OXIDE SYNTHASE, INDUCIBLE |
| 6    | 36    | XDH_MOUSE  | 7.62 | 146518 | XANTHINE DEHYDROGENASE (EC 1.1   |
| 7    | 37    | MDR1_MOUSE | 8.51 | 140994 | MULTIDRUG RESISTANCE PROTEIN     |
| 8    | 37    | RPB2_ARATH | 7.90 | 135062 | DNA-DIRECTED RNA POLYMERASE      |
| 9    | 38    | CYA6_MOUSE | 8.16 | 130319 | ADENYLATE CYCLASE, TYPE VI (EC   |
| 10   | 38    | MSH6_MOUSE | 6.61 | 151076 | DNA MISMATCH REPAIR PROTEIN      |

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This message was generated on ExPASy.

## B P388/DOX

The SWISS-PROT entries having pI and Mw values in the specified range for the species MOUSE and the specified keyword:

| Rank | Score | Protein    | pI   | Mw     | Description                    |
|------|-------|------------|------|--------|--------------------------------|
| 1    | 14    | MDR1_MOUSE | 8.51 | 140994 | MULTIDRUG RESISTANCE PROTEIN   |
| 2    | 19    | MDR3_MOUSE | 8.94 | 140755 | MULTIDRUG RESISTANCE PROTEIN   |
| 3    | 24    | CFTR_MOUSE | 8.93 | 167853 | CYSTIC FIBROSIS TRANSMEMBRAN   |
| 4    | 26    | MDR2_MOUSE | 8.88 | 140333 | MULTIDRUG RESISTANCE PROTEIN   |
| 5    | 28    | CO5_MOUSE  | 6.39 | 186817 | COMPLEMENT C5.                 |
| 6    | 29    | CYA9_MOUSE | 6.85 | 150954 | ADENYLATE CYCLASE, TYPE IX (EC |
| 7    | 29    | PHYD_ARATH | 5.81 | 129302 | PHYTOCHROME D.                 |
| 8    | 29    | CO3_MOUSE  | 6.39 | 184179 | COMPLEMENT C3.                 |
| 9    | 30    | A2MG_MOUSE | 6.12 | 134726 | ALPHA-2-MACROGLOBULIN 165 KD   |
| 10   | 31    | AT7A_MOUSE | 6.18 | 161916 | COPPER-TRANSPORTING ATPASE     |

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This message was generated on ExPASy.

Fig. 5. Output from the ExPASy program.

in the blotting cassette. Take care to remove any trapped air bubbles. Soak the cassette in the tank containing 10 mM CAPS buffer.

7. The membrane was handled with gloves to avoid contamination with skin protein.
8. The blade is washed three times immersing sequentially in three vials of 50% MeCN for each sample, to avoid cross-contamination of samples.
9. It is important for the membrane to be sufficiently wet. The next washing step is inadequate if the membrane is dry. The wet membrane sinks to the bottom of the tube containing 0.01 N HCl solution.
10. As the optimal pH for AQC-derivatization is 8.8, any residual HCl prevents the reaction (4).
11. The excess reagent is destroyed immediately after derivatization
12. Although the resolution for Lys and Ile changes with column aging, it can be improved by changing the column temperature (50°C from 40 min to 60 min).
13. Column maintenance is performed with the same elution program as that for analysis, except that the flow rate is 0.1 mL/min.
14. AQC-Met is easily oxidized in the diluted solution, and is converted to AQC-methionine sulfoxide, which is eluted between Ser and Gly.
15. The concentration of amino acids may differ by more than 10–20-fold. Also, the fluorescence detection loses its linearity, as a result of quenching by high concentration of fluorophore. Therefore, it is better to analyze two levels of sample concentration successively (for example, an intact sample solution [x 1] and a diluted sample [x 10]).
16. For PROSEARCH in EMBL, the corrected amino acid composition is used for the database search. That is calculated from the experimental amino acid composition divided by the theoretical one using ProtParm tool (<http://www.expasy.ch/tools/#primary>).

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## Capillary Gas Chromatographic Analysis of Protein and Nonprotein Amino Acids in Biological Samples

Hiroyuki Kataoka, Sayuri Matsumura, Shigeo Yamamoto,  
and Masami Makita

### 1. Introduction

Gas chromatography (GC) has been widely utilized for amino acid analysis because of its inherent advantages of high resolving power, high sensitivity, simplicity, and low cost. However, the main problem is the necessity to derivatize the amino acids into less polar and more volatile compounds. Therefore, not only the development of GC systems, but also the concomitant development of suitable derivatives are important factors to accomplish satisfactory GC analysis. An enormous number of approaches for the preparation of volatile derivatives suitable for quantitative analysis of protein and nonprotein amino acids has been reported.

The N-perfluoroacyl alkyl esters including the N-trifluoroacetyl (TFA) methyl esters (**1,2**), the N-TFA n-propyl esters (**3**), the N-TFA n-butyl esters (**4,5**), N-pentafluoropropionyl isopropyl esters (**6**), the N-heptafluorobutyryl isobutyl esters (**5,7–12**), and the N-pentafluorobenzoyl isobutyl esters (**13**) have been studied most extensively, some of which have been practically employed for the successful quantitation of amino acids in biological samples. However, it has been recently pointed out that some of these derivatives possess some negative aspects of the derivatization procedures, such as requiring anhydrous conditions, high reaction temperatures, degradation of the amides (glutamine and asparagine) caused by the HCl catalyst and low solubilities of some amino acids in the higher alcohol. The single derivatives including the trimethylsilyl derivatives (**14**) and N,(O)-*tert*-butyldimethylsilyl (*tert*-BDMS) derivatives

(15–20) have also been reported. Although these methods are nonoxidative, one-step procedure, and will not hydrolyze peptides, these derivatization reactions require anhydrous conditions, high temperatures, and long times. Furthermore, these derivatives give multiple peaks for some amino acids, and some derivatives with higher boiling points tend to decompose on long capillary column. On the other hand, we developed new volatile derivatives, N(O,S)-isobutoxycarbonyl (isoBOC) methyl esters, which were prepared by reaction with isobutyl chloroformate (isoBCF) in aqueous alkaline media, followed by esterification with diazomethane (21–24). The derivatives can be easily, rapidly, and quantitatively prepared without the use of heat and were stable to moisture, and thus appeared to be useful in practice. Subsequently, the modified method based on the preparation of N(O,S)-isoBOC *tert*-BDMS derivatives was reported (25). However, these methods require dual columns for the complete resolution of all amino acids and conversion of arginine to ornithine with arginase.

Recently, an alternative approach for the preparation of the N(O,S)-ethoxycarbonyl ethyl esters with ethyl chloroformate (ECF) was introduced (26,27). Although this method is more simple and rapid by one-step reaction in water-ethanol-pyridine and shows excellent resolution on a capillary column, it is not selective for amino acids because ECF also reacts with other compounds such as fatty acids (28), amines (29), and other organic acids (30) under the same reaction conditions, and derivatization of arginine was not achieved. Current methods for the determination of amino acids have also been described in detail (refs. 31–36).

Although our aforementioned method (21–24) is relatively time-consuming and requires two packed columns on GC separation, the derivatives could be easily and quantitatively prepared in aqueous media without the use of heat and were stable to moisture. Recently, we have achieved the improvement of our previous method in terms of speed and simplicity by using a capillary column and sonication technique. We have previously reported a simple and rapid method for the determination of protein amino acids based on the preparation of N(O,S)-isoBOC methyl ester derivatives of amino acids by using sonication technique and subsequent GC analysis with hydrogen flame ionization detection (FID) (37). By using the FID-GC method, we have demonstrated that the 22 protein amino acids could be quantitatively and reproducibly resolved as single peaks within 9 min, and the amino acid compositions of proteins (37) and protein amino acid contents in serum (38) could be rapidly and simply determined. Furthermore, we have demonstrated that the 21 protein amino acids and 33 nonprotein amino acids could be simultaneously resolved within 25 min (39) and selectively and sensitively detected by nitrogen-phosphorus selective detection (NPD) (40).

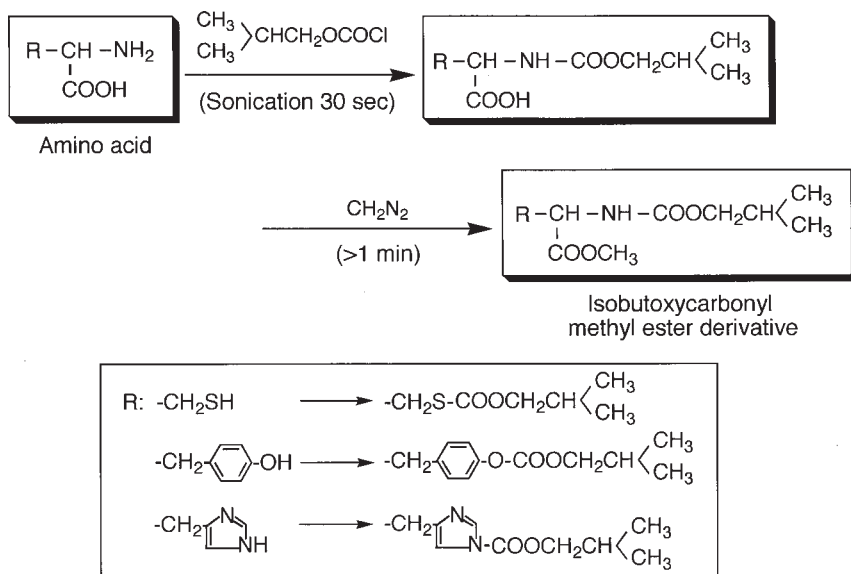


Fig. 1. Derivatization process of amino acids.

By using the NPD-GC method, the protein and nonprotein amino acids in small urine and serum samples (40,41) could be directly analyzed without prior clean-up and any interference from other substances. In this chapter, a method for the derivatization and FID- and NPD-GC analyzes of protein and nonprotein amino acids in biological samples is described on the basis of these results, and optimum conditions and typical problems encountered in the development and application of the method are discussed.

The derivatization process is shown in Fig. 1. N(O,S)-isoboc methyl esters of amino acids can be easily, rapidly, and quantitatively prepared by the reaction with isobcf by sonication in aqueous alkaline media, followed by esterification with diazomethane. In this derivatization, all amino, imino, imidazole, phenolic hydroxyl, and sulfhydryl groups are substituted with isoboc groups, whereas alcoholic hydroxyl groups and indole ring nitrogen, as well as amide groups are not. On the other hand, arginine, an exceptional amino acid in our method, is successfully analyzed as the derivative of ornithine after facile treatment with arginase (37). The chief advantage of this method is that the amino acids in aqueous solution can be easily converted into their N(O,S)-isoboc amino acids without the necessity of heating and without any step to exclude water being necessary prior to derivatization. Furthermore, the derivatives are stable to moisture and can be quantitatively and reproducibly resolved as single peaks using a capillary column.

## 2. Materials

### 2.1. Equipment

1. Hewlett-Packard 5890 Series II gas chromatograph equipped with a hydrogen flame ionization detector and a nitrogen-phosphorus detector, an electronic pressure control (EPC) system and a split/splitless capillary inlet system.
2. For the FID-GC analysis, a fused-silica capillary column of crosslinked DB-17 (50% phenyl-50% methylpolysiloxane, J & W, Folsom, CA: 15 m × 0.25-mm id, 0.25- $\mu$ m film thickness).
3. For the NPD-GC analysis, a fused-silica capillary column of crosslinked DB-17ht (50% phenyl-50% methylpolysiloxane, J & W, Folsom, CA: 20 m × 0.32-mm id, 0.15- $\mu$ m film thickness).
4. Model RD-41 centrifugal evaporator (Yamato Kagaku, Tokyo, Japan).
5. Pico-Tag workstation (Waters Associates, Milford, MA).
6. Model UT-104 Ultra sonic (39 kHz) cleaner (Sharp, Tokyo, Japan).
7. Pasteur capillary pipet (Iwaki glass No. IK-PAS-5P).

### 2.2. Reagents

1. Protein amino acids: glycine (Gly), L-alanine (Ala), L-valine (Val), L-leucine (Leu), L-isoleucine (Ile), L-threonine (Thr), L-serine (Ser), L-proline (Pro), L-aspartic acid (Asp), L-glutamic acid (Glu), L-methionine (Met), L-hydroxyproline (Hyp), L-phenylalanine (Phe), L-arginine (Arg), L-lysine (Lys), L-histidine (His), L-tyrosine (Tyr), L-tryptophan (Trp), and L-cystine (Cyt) were purchased from Ajinomoto (Tokyo, Japan); and L-cysteine (Cys), L-asparagine (Asn), and L-glutamine (Gln) from Nacalai Tesque (Kyoto, Japan).
2. Nonprotein amino acids: L- $\alpha$ -aminobutyric acid ( $\alpha$ -ABA),  $\alpha$ -aminoisobutyric acid ( $\alpha$ -AIBA), DL- $\beta$ -aminobutyric acid ( $\beta$ -ABA), DL- $\beta$ -aminoisobutyric acid ( $\beta$ -AIBA),  $\beta$ -alanine ( $\beta$ -Ala), DL-norvaline (NVal), DL-norleucine (NLeu),  $\gamma$ -aminobutyric acid (GABA), DL-homoserine (HSer), L-thioprolin (TPro), DL-homocysteine (HCys), L-ornithine (Orn) and DL-homocystine (HCyt) were purchased from Nacalai Tesque;  $\epsilon$ -amino-*n*-caproic acid ( $\epsilon$ -ACA), *p*-aminobenzoic acid (*p*-ABzA), and DL- $\alpha$ -aminopimelic acid ( $\alpha$ -APA) from Tokyo Kasei Kogyo (Tokyo, Japan); L-allo-isoleucine (AlIle), L-pipecolic acid (PCA),  $\delta$ -aminolevulinic acid ( $\delta$ -ALA), DL- $\alpha$ -aminoadipic acid ( $\alpha$ -AAA), DL-2,3-diaminopropionic acid (DAPA), DL-2,4-diaminobutyric acid (DABA), L-methionine sulphone (Met-S), DL- $\delta$ -hydroxylysine ( $\delta$ -HLys) and L-cystathionine (CTH) from Sigma (St. Louis, MO).
3. Internal standards: isonipecotic acid and 4-piperidinecarboxylic acid as internal standards (IS) were purchased from Tokyo Kasei Kogyo.
4. Protein samples: bovine serum albumin (Fraction V, 96–99%), hen ovalbumin (Type VII), calf thymus histone (Type II), and calf skin collagen (Type III, acid soluble) were purchased from Sigma.
5. Derivatizing reagents: isoBCF obtained from Tokyo Kasei Kogyo was used without further purification and stored at 4°C. N-Methyl-N-nitroso-*p*-toluenesulpho-

namide and diethyleneglycol mono-methyl ether for the generation of diazomethane (**42**) were obtained from Nacalai Tesque.

6. Other materials: dithioerythritol (DTE) was obtained from Nacalai Tesque. Peroxide-free diethyl ether was purchased from Dojindo Laboratories (Kumamoto, Japan). Distilled water was used after freshly purification with a Model Milli-Q Jr. water purifier (Millipore, Bedford, MA). All other chemicals were analytical grade.

### 2.3. Solutions

1. Standard amino acid solutions: three standard stock solutions (each 1 mg/mL), one containing the 22 protein amino acids, the second containing the 22 protein amino acids plus Orn, and the third containing the 21 protein amino acids (except for Arg) plus the 25 nonprotein amino acids; prepared in 0.05 M HCl.
2. Working standard solutions: 1 mg/mL (for FID-GC analysis) or 2 µg/mL (for NPD-GC analysis) of each amino acid were made up freshly as required by mixing of the stock solutions and then dilution with 0.05 M HCl. These working standard solutions are stable at 4°C for at least 2 wk.
3. Internal standard solutions: isonipecotic acid and 4-piperidinecarboxylic acid are dissolved in 0.05 M HCl at a concentration of 1 mg/mL as stock solution. These IS solutions are stable at 4°C for at least 2 wk.
4. Arginase solution: 5 mg of arginase (40 U/mg, Sigma) activated in 4.5 mL of 0.1 M ammonium acetate and 0.5 mL of 0.5 M manganese (II) sulfate at 37°C for 4 h. Centrifuge the solution for 1 min at 2000g, then separate the supernatant and store this solution at -20°C until used. This stored solution of activated arginase can be used without loss of activity for at least 15 d.
5. DTE solution: DTE is used as a 0.5 mM solution in distilled water.

## 3. Methods

### 3.1. Derivatization

Derivatization procedure of amino acids is shown in **Fig. 2**.

1. For the FID-GC analysis of protein amino acids, pipette an aliquot of the sample containing 0.2–50 µg of each amino acid and 20 µL of 0.1 mg/mL isonipecotic acid (IS) (if necessary) into a 10 mL Pyrex glass reaction tube with a PTFE-lined screw-cap.
2. For the NPD-GC analysis of protein and nonprotein amino acids, pipet an aliquot of the sample containing 0.02–2 µg of each amino acid and 0.1 mL of 2 µg/mL 4-piperidinecarboxylic acid (IS) (if necessary) into a reaction tube.
3. Add 50 µL of 0.5 mM DTE (*see Note 1*) and 0.25 mL of 10% Na<sub>2</sub>CO<sub>3</sub> to these solutions and make the total volume up to 1 mL with distilled water.
4. Immediately add 20 µL of isoBCF to the mixture and sonicate for 30 s at room temperature after shaking for a few s by hand (*see Notes 2–4*).
5. Extract the reaction mixture with 3 mL of peroxide-free diethyl ether, to remove the excess of reagent, with vigorous shaking for 5–10 s by hand (*see Note 5*).

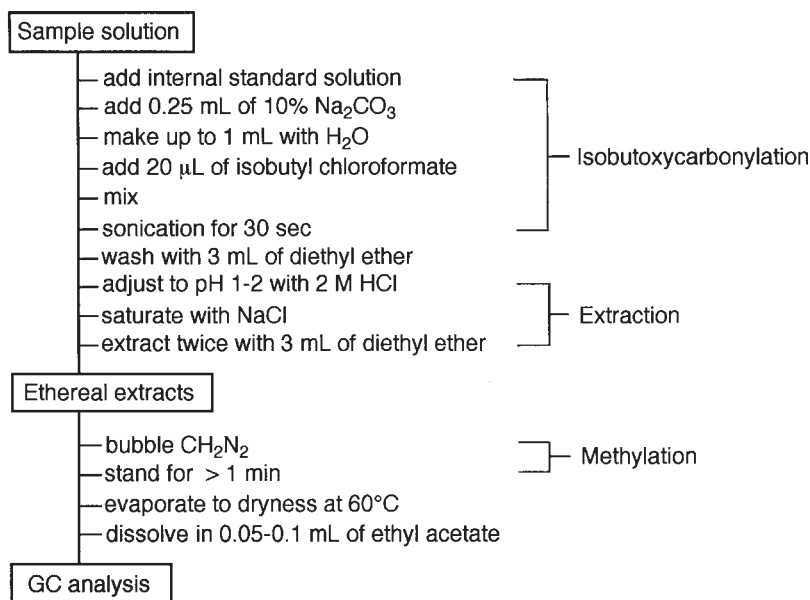


Fig. 2. Schematic flow diagram of the derivatization procedure.

6. Centrifuge at 2000g for 30 s, then discard the ethereal extract.
7. Acidify the aqueous layer to pH 1.0–2.0 with 2 M HCl and saturate with NaCl. Then extract twice with 3 mL of peroxide-free diethyl ether with vigorous shaking for 5–10 s by hand (*see Note 6*).
8. Again centrifuge at 2000g for 30 s. Transfer the ether layers into another tube by means of a Pasteur capillary pipet.
9. The pooled ethereal extracts are methylated by bubbling diazomethane, generated according to the microscale procedure (40) (*see Note 7*). As shown in **Fig. 3**, a stream of nitrogen is saturated with diethyl ether in the first side-arm test tube and then passed through a diazomethane generating solution containing N-methyl-N-nitroso-*p*-toluenesulphonamide, diethylene-glycol monomethyl ether and KOH. The generated diazomethane is carried into the sample tube via the nitrogen stream until a yellow tinge became visible. After standing for 1 min at room temperature, the solvents are removed by evaporation to dryness at 60°C under a stream of dry air. The residue on the walls of the tube is dissolved in 0.05–0.1 mL of ethyl acetate and then 1 µL of this solution is injected into the gas chromatograph by hot-needle-injection technique (*see Note 8*).
10. On the other hand, Arg is analyzed as the derivative of Orn after treatment with arginase (*see Note 9*). For conversion of Arg into Orn, pipet an aliquot of the solution containing 0.5–10 µg of Arg and 20 µL of 0.1 mg/mL IS (if necessary) into a reaction tube. To this solution, add 0.5 mL of 0.2 M Na<sub>2</sub>CO<sub>3</sub>-NaHCO<sub>3</sub> buffer (pH 9.5) and make the total volume up to 0.7 mL with distilled water. Add

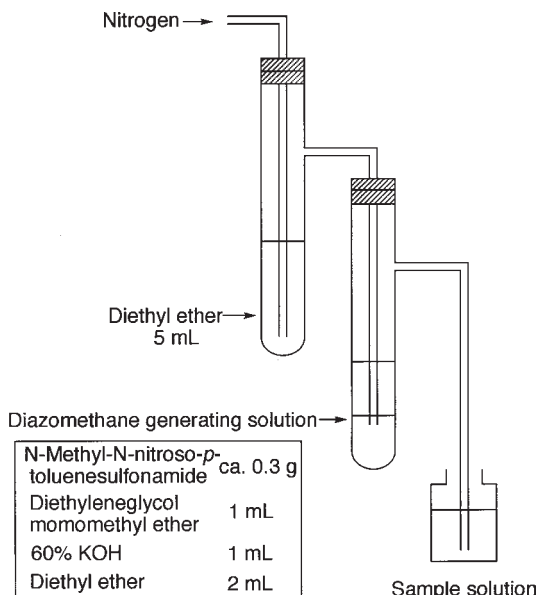


Fig. 3. Apparatus for microscale generation of diazomethane.

the activated arginase solution (50  $\mu$ L), incubate at 37°C for 10 min, and then derivatize as above (*see Note 10*).

### 3.2. Gas Chromatography

1. FID-GC analysis is carried out with a Hewlett-Packard 5890 Series II gas chromatograph equipped with a hydrogen flame ionization detector and a split injection system. A fused-silica capillary column of crosslinked DB-17 (15 m  $\times$  0.25-mm id, 0.25- $\mu$ m film thickness) is used. The operating conditions are as follows: column temperature program at 30°C/min from 140°C to 290°C and then hold for 5 min; injection and detector temperatures, 300°C; helium carrier gas and make-up gas flow rates, 1 mL/min and 30 mL/min, respectively; split ratio, 50:1.
2. NPD-GC analysis is carried out with a Hewlett-Packard 5890 Series II gas chromatograph equipped with a nitrogen-phosphorus detector (NPD), an electronic pressure control (EPC) system, and a split/splitless capillary inlet system. A fused-silica capillary column of crosslinked DB-17ht (20 m  $\times$  0.32-mm id, 0.15- $\mu$ m film thickness) is used. The operation conditions are as follows: column temperature program at 4°C/min from 120°C to 160°C, then program at 6°C/min from 160°C to 200°C and at 10°C/min from 200°C to 310°C; inlet helium carrier gas flow rate, control at 2 mL/min constant with EPC; split ratio, 10:1. Injection and detector temperatures and make-up gas flow rate are 320°C and 30 mL/min, respectively (*see Notes 11 and 12*).

3. A chromatographic blank run (run made with no sample injected) data should be subtracted from the sample run data to remove baseline drift (usually caused by column bleed) using a single-column compensation function. Baseline-corrected data is recorded on the chromatogram.
4. Measure the peak heights of amino acids and the IS, and calculate the peak height ratios of amino acids against the IS to construct calibration curves (*see* **Notes 13–19**).

### 3.3. Sample Preparation

#### 3.3.1. Hydrolysis of Protein Samples

1. Place an aliquot of the protein sample (10–100  $\mu\text{g}$ ) in a 5  $\times$  50 mm glass test tube, and dry in a centrifugal evaporator.
2. Hydrolyze the residue with 0.2 mL of 6 M HCl containing 1% phenol (*see* **Note 20**) in the vapor phase for 24 h at 110°C under vacuum in a Pico-Tag workstation (**43**) (*see* **Note 21**).
3. To the resulting hydrolysate, add 20  $\mu\text{L}$  of 0.1 mg/mL isonipectic acid (IS) and extract the mixture twice with 0.3 mL of distilled water.
4. Transfer the extracts to another reaction tube (10-mL Pyrex glass tube with a PTFE-lined screw-cap). The samples are ready for derivatization and FID-GC analysis (*see* **Note 22**).

#### 3.3.2. Preparation of Urine Samples

1. Collect early morning urine samples from healthy volunteers and process immediately or store at  $-20^{\circ}\text{C}$  until used.
2. Urine samples (10–50  $\mu\text{L}$ ) are directly used for the derivatization and NPD-GC analysis (*see* **Notes 23–25**).

#### 3.3.3. Preparation of Serum Samples

1. Collect venous blood sample from healthy volunteer in vacutainer tubes and centrifuge at 1600 g for 10 min.
2. Carefully collect the serum layer and process immediately.
3. Serum sample (50  $\mu\text{L}$ ) are directly used for the derivatization and NPD-GC analysis without deproteinization (*see* **Notes 23–25**).

## 4. Notes

1. In order to prevent the oxidation of sulfur amino acids during derivatization, DTE was added to reaction mixture (**44**).
2. Sonication technique, which can easily mix aqueous solution and oily isoBCF reagent by its vibration action, was useful for acceleration of the reaction.
3. The N(O,S)-isobutoxycarbonylation of amino acids with isoBCF was completed within 15 s by sonication at room temperature. The optimum concentration of  $\text{Na}_2\text{CO}_3$  was 2.5% as previously described (**21–24**) and the reaction yields for each amino acid were constant with  $>10$   $\mu\text{L}$  of isoBCF.
4. All amino, imino, imidazole, phenolic hydroxyl, and sulfhydryl groups are substituted with isoBOC groups, whereas alcoholic hydroxyl groups, guanidino group, and indole ring nitrogen, as well as amide groups are not.



5. The reaction mixture was washed with diethyl ether under alkaline condition in order to remove the excess reagent. This procedure also serves to exclude amines and phenols both of which often coexist with amino acids in biological samples, as they are derivatized to the corresponding N- or O-isoBOC derivatives, which are soluble in organic solvents under the same conditions as aforementioned (45–47).
6. The resulting N(O,S)-isoBOC amino acids in aqueous layer could be quantitatively and selectively extracted into diethyl ether after acidification to pH 1.0–2.0. In these extractions, peroxide-free diethyl ether should be employed as an extraction solvent to obviate decomposition of sulfur amino acids, and great care should be taken for this solvent when small amounts of these amino acids are to be analyzed by NPD-GC. Furthermore, the ether layers should be collected, taking care to avoid aqueous droplets. It was not necessary to completely draw the ether layer in each extraction.
7. The subsequent methylation of the ethereal extracts could be successfully carried out by bubbling diazomethane. This reaction should be performed in a well-ventilated hood because diazomethane is explosive and toxic.
8. The derivative preparation could be performed within 10 min and the N(O,S)-isoBOC methyl ester derivatives of protein and nonprotein amino acids were very stable under normal laboratory conditions, and no decomposition was observed during GC analysis.
9. Arg requires conversion into Orn with arginase prior to derivatization, because the guanidino group of Arg cannot be derivatized by this method and this amino acid therefore cannot be extracted into diethyl ether because of its polarity.
10. In this reaction, the conversion yield of Arg into Orn was above 93% at each level of Arg from 0.5–10 µg.
11. In a preliminary test for several megabore capillary columns, DB-17 and DB-1701 gave best separation for protein amino acids, although some pairs of amino acids were partially overlapped, e.g., Thr-Ser in DB-17 column, and Leu-Ile and Glu-Met in DB-1701 column (37). Therefore, we tested DB-17 narrow-bore columns at length of 5–20 m to obtain optimum analytical conditions with respect to the separation of amino acids and analysis time. Although the resolution of amino acids increased according to increasing length of column, the analysis time became long. Among the columns tested, 15-m column of DB-17 was proved to be best column for these purposes. As shown in Fig. 4, 22 protein amino acids could be completely resolved as single and symmetrical peaks within 9 min on a single capillary column (DB-17, 15 m × 0.25-mm id, 0.25-µm film thickness) (37). However, the separation of protein and nonprotein amino acids were incomplete on this column. Furthermore, high boiling point derivatives were eluted slowly with a broader peak width, and their sensitivities were reduced because of reduction in carrier gas flow rate at higher temperature.
12. In order to solve above problems, we tried to introduce a EPC system and a thin-film-coated and high temperature (max. at 340°C) column DB-17ht (20 m × 0.32-mm id, 0.15-µm film thickness). Of several GC conditions tested for this column, the three-ramp temperature programmes and EPC programs given in **Subhead-**

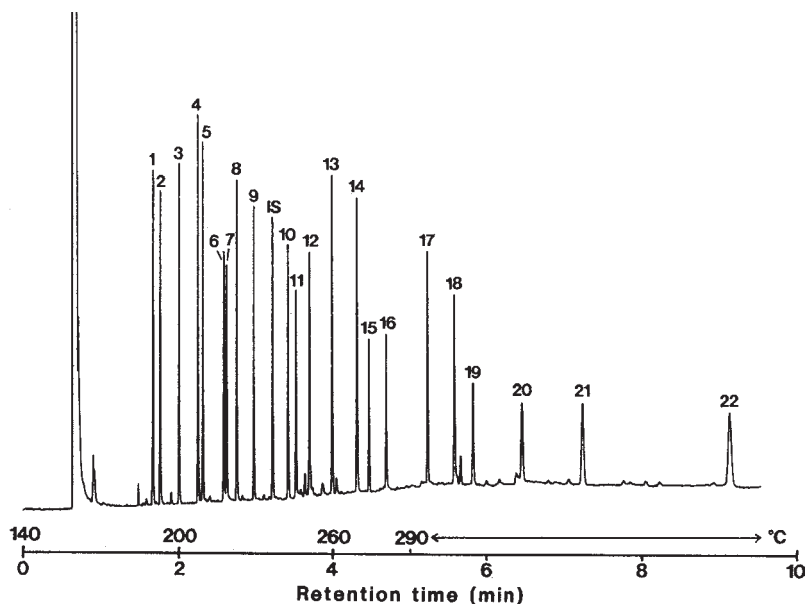


Fig. 4. Standard chromatogram of the N(O,S)-isobutoxycarbonyl methyl ester derivatives of protein amino acids (containing 25  $\mu\text{g}$  of asparagine, glutamine and cystine, and 5  $\mu\text{g}$  of other amino acids) by FID-GC. GC conditions: column, DB-17 (15 m  $\times$  0.25 mm ID, 0.25- $\mu\text{m}$  film thickness); column temperature, programmed at 30°C/min from 140°C to 290°C and then held for 5 min; injection and detector temperatures, 300°C; helium carrier gas and make-up gas flow rates, 1 mL/min and 30 mL/min, respectively; split ratio, 50:1. Peaks: 1 = alanine; 2 = glycine; 3 = valine; 4 = leucine; 5 = isoleucine; 6 = threonine; 7 = serine; 8 = proline; 9 = aspartic acid; 10 = glutamic acid; 11 = methionine; 12 = hydroxyproline; 13 = phenylalanine; 14 = asparagine; 15 = cysteine; 16 = glutamine; 17 = ornithine; 18 = lysine; 19 = histidine; 20 = tyrosine; 21 = tryptophan; 22 = cystine; IS = isonipectic acid (from **ref. 37**, with permission).

**ing 3.2.** were proved to give the most satisfactory separation of the protein and nonprotein amino acids. As shown in **Fig. 5**, the 21 protein amino acids and the 25 nonprotein amino acids could be well separated as single symmetrical peaks within 28 min on a DB-17ht capillary column, except for  $\delta$ -HLys, which showed two peaks caused by the allo form present in the standard (**41**).

13. The reproducibility of sample injection, expressed as percent variation from the mean peak height ratio against the IS, was determined from three independent injections of the standard derivative mixture. For syringe manipulation, a hot-needle-injection technique (the needle inserted into injection zone was allowed to heat up for 3 s prior to the sample injection) was used to prevent the sample discrimination caused by incomplete vaporization and flashback into other parts of the inlet. A split-injection system was proved to be reproducible and the rela-

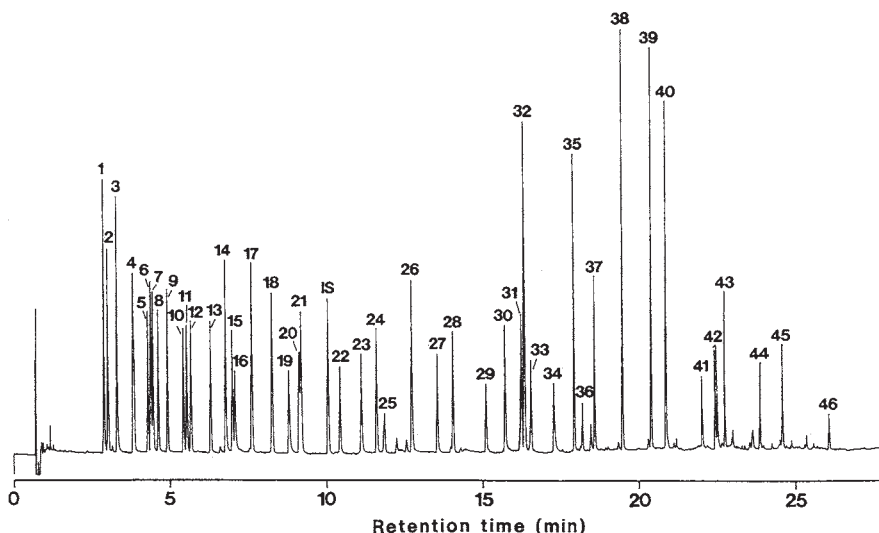


Fig. 5. Standard chromatogram obtained from the N(O,S)-isobutoxycarbonyl methyl ester derivatives of protein and nonprotein amino acids (containing 0.4  $\mu\text{g}$  of asparagine and glutamine, and 0.2  $\mu\text{g}$  of other amino acids) by NPD-GC. GC conditions: column, DB-17ht (20 m  $\times$  0.32 mm ID, 0.15- $\mu\text{m}$  film thickness); column temperature, programmed at 4°C/min from 120 to 160°C, then programmed at 6°C/min from 160 to 200°C and at 10°C/min from 200 to 310°C; injection and detector temperatures, 320°C; inlet helium carrier gas flow rate, controlled 2 mL/min constant with EPC; make-up gas flow rate, 30 mL/min; split ratio, 10:1. Peaks: 1 =  $\alpha$ -aminoisobutyric acid, 2 = alanine, 3 = glycine, 4 =  $\alpha$ -aminobutyric acid, 5 = valine, 6 =  $\beta$ -alanine, 7 =  $\beta$ -aminobutyric acid, 8 =  $\beta$ -aminoisobutyric acid, 9 = norvaline, 10 = leucine, 11 = *allo*-isoleucine, 12 = isoleucine, 13 = norleucine, 14 =  $\gamma$ -aminobutyric acid, 15 = threonine, 16 = serine, 17 = proline, 18 = pipercolic acid, 19 = homoserine, 20 =  $\delta$ -aminolevulinic acid, 21 = aspartic acid, 22 = thioproline, 23 =  $\epsilon$ -aminocaproic acid, 24 = glutamic acid, 25 = methionine, 26 = hydroxyproline, 27 =  $\alpha$ -aminoadipic acid, 28 = phenylalanine, 29 =  $\alpha$ -aminopimelic acid, 30 = asparagine, 31 = *p*-aminobenzoic acid, 32 = 2,3-diaminopropionic acid, 33 = cysteine, 34 = glutamine, 35 = 2,4-diaminobutyric acid, 36 = homocysteine, 37 = methionine sulphone, 38 = ornithine, 39 = lysine, 40 = histidine, 41 = tyrosine, 42 = *d*-hydroxylysine, 43 = tryptophan, 44 = cystathionine, 45 = cystine, 46 = homocystine, IS = 4-piperidinecarboxylic acid (from **ref. 41**, with permission).

tive standard deviations ranged from 0.06 to 2.9%. On the other hand, a splitless-injection system could introduced most of sample onto the column and increased sensitivity, but it was proved to reduce reproducibility because of poor refocusing (39). Particularly, the relative responses of high-boiling-point compounds, such as the derivatives of LTH, CTH, Cyt, and HCyt were remarkably reduced. Therefore, we adopted a hot-needle-injection technique in split mode as optimum GC injection technique.

14. The calibration curves for protein amino acids by FID-GC were conducted using isonipecotic acid, which showed a similar behavior to other amino acids during the derivatization and was well separated from other amino acids on a chromatogram as the IS. A linear relationship was obtained with correlation coefficient being above 0.998 in the range 1–50  $\mu\text{g}$  for Asn, Gln, and Cyt, 0.5–10  $\mu\text{g}$  for Arg and 0.2–10  $\mu\text{g}$  for other amino acids. The relative standard deviations for each amino acid in each point were 0.5–7.0% ( $n = 3$ ).
15. The minimum detectable amounts of protein amino acids to give a signal three times as high as the noise under our FID-GC conditions were 0.2–4.0 ng as injection amounts.
16. The contents of Orn and 21 protein amino acids, except for Arg, were calculated from the directly derivatized samples without arginase treatment. On the other hand, Arg content was calculated by subtracting the amount of Orn obtained without arginase treatment from that obtained with arginase treatment (37).
17. NPD-GC is selective and sensitive for nitrogen-containing compounds and the application of this technique to the analysis of amino acids has been reported (8,12,48–52). The NPD-GC system described here was over 10–50 times more sensitive than the FID-GC. Particularly, the increase of sensitivity was remarkable for the nitrogen-rich amino acids such as DAPA, DABA, Orn, Lys, and His (40).
18. The calibration curves for protein and nonprotein amino acids by NPD-GC were conducted using 4-piperidinecarboxylic acid, which showed similar behavior to other amino acids during the derivatization and was well separated from other amino acids on a chromatogram as the IS. Various amounts of protein and non-protein amino acids ranging from 0.02 to 2  $\mu\text{g}$  were derivatized in the mixture, and aliquots representing 0.2–20 ng were injected into the NPD-GC system. As pointed previously (12), a significant curvature of the detector response was observed by NPD-GC, particularly at a low concentration of amino acid. Therefore, the calibration curves for each amino acid were constructed from both logarithmic plots of the peak height ratios and the amino acid amounts. A linear relationship was obtained with correlation coefficients being above 0.990, in the range 0.04–2  $\mu\text{g}$  for Asn and Gln, and 0.02–1  $\mu\text{g}$  for other amino acids (Table 1).
19. The minimum detectable amounts to give a signal-to-noise ratio of 3 under our NPD-GC conditions were 6–150 pg as injection amount (Table 1) (41).
20. In order to prevent the oxidation of Tyr and sulfur amino acids during acid hydrolysis, phenol was added to 6 M HCl at the concentration of 1%. In this hydrolysis, Trp is generally destroyed, Asn and Gln are quantitatively converted to Asp and Glu.
21. The FID-GC method developed was successfully applied to acid hydrolysate samples of proteins (10–100  $\mu\text{g}$ ) (37). Acid hydrolysis of proteins and peptides using a Pico-Tag workstation is not contaminated with reagent because of proceeding in the HCl vapor phase (43).
22. Typical chromatograms obtained from some proteins by FID-GC are shown in Fig. 6. The recoveries of total amino acids were 90–104%, and the reproducibility for each amino acid was satisfactory. The amino acid compositions of four well-characterized proteins, bovine serum albumin, hen ovalbumin, calf thymus

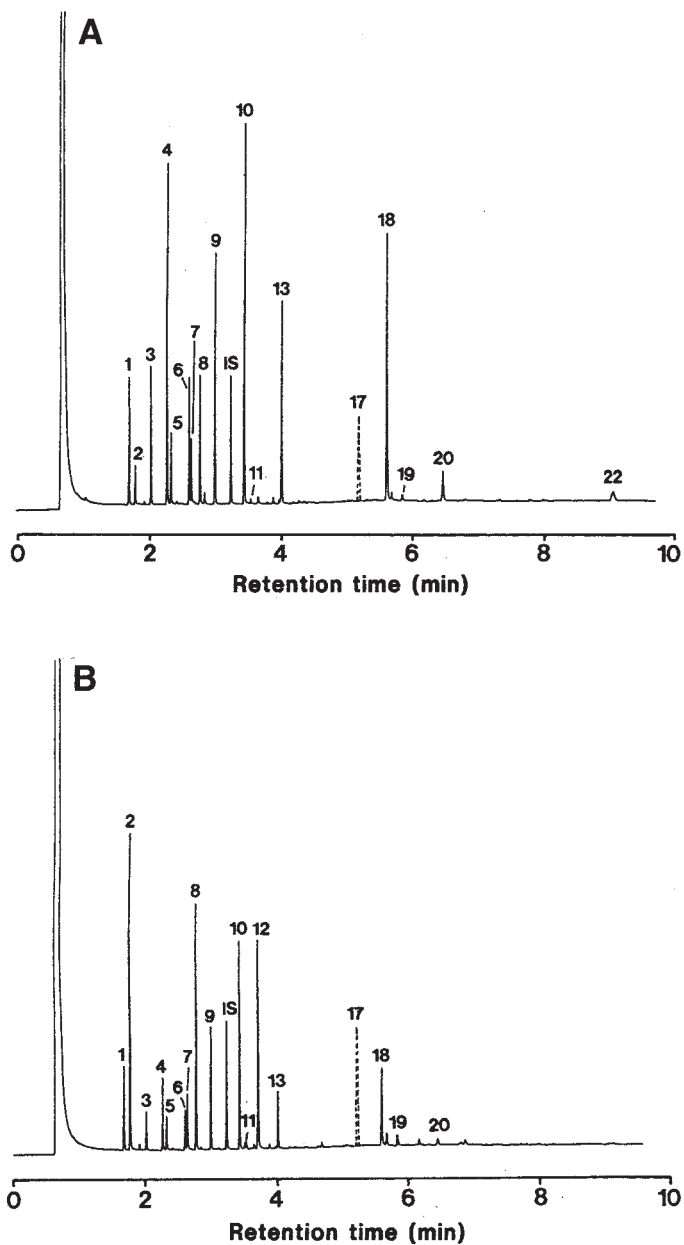


Fig. 6. Typical chromatograms obtained from acid hydrolysates of (A) bovine serum albumin (100  $\mu$ g) and (B) collagen (100  $\mu$ g) by FID-GC. GC conditions and peak number: *see* Fig. 4. Broken line shows the peak of arginine, which is analyzed as ornithine after arginase treatment (from *ref. 37*, with permission).

**Table 1**  
**Linear Regression Data and Detection Limits for Protein**  
**and Nonprotein Amino Acids**

| Peak No. | Amino acid*                    | Abbreviation    | Regression line <sup>a</sup> |                    | Correlation coefficient <i>r</i> | Detection limit (pg) |
|----------|--------------------------------|-----------------|------------------------------|--------------------|----------------------------------|----------------------|
|          |                                |                 | Slope <i>a</i>               | Intercept <i>b</i> |                                  |                      |
| 1        | $\alpha$ -Aminoisobutyric acid | $\alpha$ -AIBA  | 0.9981                       | 0.9016             | 0.9967                           | 9                    |
| 2        | Alanine                        | Ala             | 1.0901                       | 0.9456             | 0.9944                           | 10                   |
| 3        | Glycine                        | Gly             | 1.1637                       | 1.0627             | 0.9952                           | 9                    |
| 4        | $\alpha$ -Aminobutyric acid    | $\alpha$ -ABA   | 1.0942                       | 0.8731             | 0.9986                           | 10                   |
| 5        | Valine                         | Val             | 1.0465                       | 0.7292             | 0.9988                           | 11                   |
| 6        | $\beta$ -Alanine               | $\beta$ -Ala    | 1.2122                       | 0.8740             | 0.9987                           | 11                   |
| 7        | $\beta$ -Aminobutyric acid     | $\beta$ -ABA    | 1.0833                       | 0.8355             | 0.9990                           | 11                   |
| 8        | $\beta$ -Aminoisobutyric acid  | $\beta$ -AIBA   | 1.1227                       | 0.7655             | 0.9993                           | 11                   |
| 9        | Norvaline                      | NVal            | 1.0645                       | 0.7950             | 0.9991                           | 11                   |
| 10       | Leucine                        | Leu             | 1.0282                       | 0.6375             | 0.9997                           | 12                   |
| 11       | <i>allo</i> -Isoleucine        | AlIe            | 1.0163                       | 0.6492             | 0.9989                           | 12                   |
| 12       | Isoleucine                     | Ile             | 1.0185                       | 0.6627             | 0.9997                           | 12                   |
| 13       | Norleucine                     | NLeu            | 1.0670                       | 0.6796             | 0.9999                           | 12                   |
| 14       | $\gamma$ -Aminobutyric acid    | GABA            | 1.1886                       | 0.8782             | 0.9991                           | 11                   |
| 15       | Threonine                      | Thr             | 1.2551                       | 0.7440             | 0.9995                           | 28                   |
| 16       | Serine                         | Ser             | 1.4961                       | 0.7727             | 0.9953                           | 50                   |
| 17       | Proline                        | Pro             | 1.0113                       | 0.8217             | 0.9996                           | 9                    |
| 18       | Pipecolic acid                 | PCA             | 1.0078                       | 0.7320             | 0.9998                           | 11                   |
| 19       | Homoserine                     | HSer            | 1.2982                       | 0.6959             | 0.9992                           | 32                   |
| 20       | $\delta$ -Aminolevulinic acid  | $\delta$ -ALA   | 1.13 13                      | 0.5799             | 0.9992                           | 25                   |
| 21       | Aspartic acid                  | Asp             | 0.9660                       | 0.6543             | 0.9997                           | 14                   |
| 22       | Thioprolin                     | TPro            | 1.2683                       | 0.5937             | 0.9984                           | 30                   |
| 23       | $\epsilon$ -Aminocaproic acid  | $\epsilon$ -ACA | 1.0060                       | 0.5151             | 0.9994                           | 16                   |
| 24       | Glutamic acid                  | Glu             | 1.0266                       | 0.6256             | 0.9998                           | 16                   |

(continued)

histone, and calf skin collagen, determined by this method were good agreement with the literature values. By using this method, protein amino acid contents in serum could be rapidly and simply determined (38).

23. The NPD-GC method developed was successfully applied to the analysis of free amino acids in human urine (40,41) and serum (41) samples without prior clean-up procedures such as deproteinization, ion-exchange column chromatography, solid phase extraction, and subsequent eluate evaporation.
24. Typical chromatograms obtained from 25  $\mu$ L of urine and 50  $\mu$ L of serum by NPD-GC were shown in Fig. 7. In urine sample, hippuric acid was observed between TPro and Glu, but it was not overlapped with  $\epsilon$ -ACA. Some unknown peaks were observed on the chromatogram, but 37 amino acids detected in the urine could be analyzed without any influence from coexisting substances.

**Table 1 (continued)**  
**Linear Regression Data and Detection Limits for Protein and Nonprotein Amino Acids**

| Peak No. | Amino acid*                 | Abbreviation   | Regression line <sup>a</sup> |                    | Correlation coefficient <i>r</i> | Detection limit (pg) |
|----------|-----------------------------|----------------|------------------------------|--------------------|----------------------------------|----------------------|
|          |                             |                | Slope <i>a</i>               | Intercept <i>b</i> |                                  |                      |
| 25       | Methionine                  | Met            | 1.3815                       | 0.2873             | 0.9971                           | 135                  |
| 26       | Hydroxyproline              | Hyp            | 1.1833                       | 0.8384             | 0.9987                           | 16                   |
| 27       | $\alpha$ -Amino adipic acid | $\alpha$ -AAA  | 1.0597                       | 0.5232             | 0.9992                           | 24                   |
| 28       | Phenylalanine               | Phe            | 0.9609                       | 0.5718             | 0.9996                           | 15                   |
| 29       | $\alpha$ -Aminopimelic acid | $\alpha$ -APA  | 1.0683                       | 0.4584             | 0.9983                           | 28                   |
| 30       | Asparagiine                 | Asn            | 1.3192                       | 0.3684             | 0.9961                           | 56                   |
| 31       | <i>p</i> -Aminobenzoic acid | <i>p</i> -ABzA | 1.1047                       | 0.6917             | 0.9980                           | 20                   |
| 32       | 2,3-Diaminopropionic acid   | DAPA           | 1.0236                       | 1.0170             | 0.9984                           | 9                    |
| 33       | Cysteine                    | Cys            | 1.1029                       | 0.5802             | 0.9974                           | 24                   |
| 34       | Glutamine                   | Gln            | 1.3119                       | 0.1786             | 0.9968                           | 150                  |
| 35       | 2,4-Diaminobutyric acid     | DABA           | 1.0891                       | 1.0254             | 0.9991                           | 10                   |
| 36       | Homocysteine                | HCys           | 0.9898                       | 0.1160             | 0.9931                           | 60                   |
| 37       | Methioninesulphone          | Met-S          | 1.0781                       | 0.7250             | 0.9933                           | 25                   |
| 38       | Ornithine                   | Orn            | 1.0267                       | 1.1324             | 0.9982                           | 6                    |
| 39       | Lysine                      | Lsy            | 1.0214                       | 1.1195             | 0.9961                           | 6                    |
| 40       | Histidine                   | His            | 1.0178                       | 1.0684             | 0.9963                           | 7                    |
| 41       | Tyrosine                    | Tyr            | 1.4371                       | 0.6400             | 0.9964                           | 64                   |
| 42       | $\delta$ -Hydroxylysine     | $\delta$ -HLys | 1.1698                       | 0.6803             | 0.9934                           | 34                   |
| 43       | Tryptophan                  | Trp            | 1.2711                       | 0.8663             | 0.9951                           | 24                   |
| 44       | Cystathionine               | CTH            | 1.4594                       | 0.7296             | 0.9936                           | 30                   |
| 45       | Cystine                     | Cyt            | 1.2572                       | 0.7120             | 0.9964                           | 40                   |
| 46       | Homocystine                 | HCyt           | 1.2690                       | 0.4084             | 0.9901                           | 70                   |

<sup>a</sup> $\log y = a \log x + b$ : *y*, peak height ratio against the I.S.; *x*, amount of each amino acid ( $\mu\text{g}$ ); *a*, slope; *b*, intercept. Range: 0.04–2.0  $\mu\text{g}$  for asparagine and glutamine; 0.02–1.00  $\mu\text{g}$  for other amino acids (from ref. 41, with permission).

25. To confirm validity of NPD-GC method, known amounts of amino acids were spiked to human urine and serum, and their recoveries were calculated. As shown in **Table 2**, the overall recoveries of these amino acids were 83.5–112.0% and the relative standard deviations were 0.3–14.9% ( $n = 3$ ). The quantitation limits of amino acids in urine and serum samples were approx 0.1–0.4 mg/mL. The intra-assay C.V.s and inter-assay C.V.s for these samples throughout the overall procedure consisting of derivatization and GC analysis were 0.3–8.9% ( $n = 3$ ) and 1.9–15.8% ( $n = 3$ ), respectively.

**Table 2**  
**Recoveries of Amino Acids Added to Urine and Serum Samples**

| Amino acid | Urine         |                                   |              |              | Serum         |                      |              |              |
|------------|---------------|-----------------------------------|--------------|--------------|---------------|----------------------|--------------|--------------|
|            | Added (µg/mL) | Amount found (µg/mL) <sup>a</sup> |              | Recovery (%) | Added (µg/mL) | Amount found (µg/mL) |              | Recovery (%) |
|            |               | Nonaddition                       | Addition     |              |               | Nonaddition          | Addition     |              |
| α-AIBA     | 2             | 8.61 ± 0.16                       | 10.59 ± 0.80 | 99.0         | 1             | 0.91 ± 0.08          | 1.89 ± 0.08  | 98.0         |
| Ala        | 20            | 18.48 ± 1.41                      | 38.12 ± 2.17 | 98.2         | 10            | 25.68 ± 0.45         | 36.07 ± 2.09 | 103.9        |
| Gly        | 20            | 36.49 ± 0.43                      | 57.59 ± 1.62 | 105.5        | 10            | 18.05 ± 0.35         | 28.23 ± 0.91 | 101.8        |
| α-ABA      | 2             | 0.88 ± 0.07                       | 2.96 ± 0.07  | 104.0        | 1             | 3.15 ± 0.13          | 4.10 ± 0.04  | 95.0         |
| Val        | 2             | 3.69 ± 0.06                       | 5.59 ± 0.22  | 95.0         | 10            | 29.33 ± 0.74         | 39.75 ± 1.48 | 104.2        |
| β-Ala      | 2             | 2.45 ± 0.09                       | 4.41 ± 0.20  | 98.0         | 1             | ND                   | 1.04 ± 0.12  | 104.0        |
| β-ABA      | 2             | ND <sup>b</sup>                   | 2.13 ± 0.02  | 106.5        | 1             | ND                   | 1.10 ± 0.09  | 110.0        |
| β-MBA      | 20            | 108.2 ± 2.3                       | 127.8 ± 3.0  | 98.0         | 1             | ND                   | 0.99 ± 0.03  | 99.0         |
| NVal       | 2             | ND                                | 2.15 ± 0.05  | 107.5        | 1             | ND                   | 1.01 ± 0.06  | 101.0        |
| Lcu        | 2             | 3.80 ± 0.07                       | 5.80 ± 0.12  | 100.0        | 10            | 17.52 ± 0.52         | 26.60 ± 1.68 | 90.8         |
| Alle       | 2             | ND                                | 2.03 ± 0.10  | 101.5        | 1             | ND                   | 1.12 ± 0.14  | 112.0        |
| Ile        | 2             | 1.33 ± 0.02                       | 3.41 ± 0.08  | 104.0        | 10            | 10.11 ± 0.20         | 19.87 ± 1.01 | 97.6         |
| NLeu       | 2             | ND                                | 2.16 ± 0.12  | 108.0        | 1             | ND                   | 0.98 ± 0.05  | 98.0         |
| GABA       | 2             | ND                                | 220 ± 007    | 110.0        | 1             | ND                   | 1.05 ± 0.05  | 105.0        |
| Thr        | 20            | 11.13 ± 0.19                      | 29.80 ± 0.72 | 93.4         | 10            | 19.85 ± 0.09         | 29.34 ± 0.74 | 94.9         |
| Ser        | 20            | 19.77 ± 0.99                      | 38.16 ± 1.96 | 92.0         | 10            | 14.03 ± 0.19         | 23.43 ± 0.76 | 94.0         |
| Pro        | 2             | 0.79 ± 0.06                       | 2.80 ± 0.07  | 100.5        | 10            | 27.65 ± 1.16         | 37.65 ± 0.71 | 100.0        |
| PCA        | 2             | ND                                | 1.95 ± 0.06  | 97.5         | 1             | ND                   | 1.04 ± 0.04  | 104.0        |
| HSer       | 2             | 0.88 ± 0.08                       | 2.91 ± 0.08  | 101.5        | 1             | ND                   | 0.97 ± 0.11  | 97.0         |
| δ-ALA      | 2             | 1.20 ± 0.07                       | 3.35 ± 0.23  | 107.5        | 1             | ND                   | 1.07 ± 0.03  | 107.0        |
| Asp        | 2             | 0.80 ± 0.04                       | 2.75 ± 0.16  | 97.5         | 1             | 1.10 ± 0.10          | 209 ± 008    | 99.0         |
| TPro       | 2             | 1.12 ± 0.07                       | 3.05 ± 0.26  | 96.5         | 1             | ND                   | 0.97 ± 0.12  | 97.0         |
| ε-ACA      | 2             | ND                                | 1.88 ± 0.04  | 94.0         | 1             | ND                   | 1.07 ± 0.03  | 107.0        |



|        |    |              |              |       |    |              |              |       |
|--------|----|--------------|--------------|-------|----|--------------|--------------|-------|
| Glu    | 2  | 2.28 ± 0.08  | 4.35 ± 0.14  | 103.5 | 10 | 13.86 ± 0.52 | 23.51 ± 0.54 | 96.5  |
| Met    | 2  | 1.93 ± 0.22  | 3.60 ± 0.14  | 83.5  | 1  | 1.00 ± 0.03  | 1.86 ± 0.04  | 86.0  |
| Hyp    | 2  | ND           | 1.83 ± 0.16  | 91.5  | 1  | 1.95 ± 0.07  | 2.95 ± 0.21  | 100.0 |
| α-AAA  | 2  | 6.31 ± 0.02  | 8.25 ± 0.30  | 97.0  | 1  | ND           | 1.04 ± 0.02  | 104.0 |
| Phe    | 2  | 5.65 ± 0.06  | 7.75 ± 0.12  | 105.0 | 10 | 10.94 ± 0.39 | 21.06 ± 0.65 | 101.2 |
| α-APA  | 2  | 3.60 ± 0.12  | 5.51 ± 0.30  | 95.5  | 1  | ND           | 0.99 ± 0.02  | 99.0  |
| Asn    | 4  | 16.40 ± 0.48 | 20.40 ± 0.22 | 100.0 | 2  | 7.52 ± 0.62  | 9.66 ± 0.38  | 107.0 |
| p-ABzA | 2  | 6.16 ± 0.08  | 8.28 ± 0.16  | 106.0 | 1  | ND           | 1.09 ± 0.04  | 109.0 |
| DAPA   | 2  | ND           | 2.08 ± 0.18  | 104.0 | 1  | ND           | 0.99 ± 0.08  | 99.0  |
| Cys    | 2  | 8.41 ± 0.13  | 10.39 ± 0.46 | 99.0  | 1  | 0.94 ± 0.04  | 1.87 ± 0.24  | 93.0  |
| Gln    | 40 | 59.54 ± 1.48 | 97.02 ± 6.58 | 93.7  | 20 | 100.2 ± 2.8  | 119.1 ± 3.2  | 94.5  |
| DABA   | 2  | 6.57 ± 0.05  | 8.43 ± 0.53  | 93.0  | 1  | 0.10 ± 0.01  | 1.07 ± 0.04  | 97.0  |
| HCys   | 20 | 16.68 ± 0.08 | 34.87 ± 2.36 | 91.0  | 1  | ND           | 0.97 ± 0.08  | 97.0  |
| Met-S  | 2  | 1.68 ± 0.14  | 3.55 ± 0.26  | 93.5  | 1  | 0.51 ± 0.06  | 1.48 ± 0.22  | 97.0  |
| Orn    | 2  | 6.60 ± 0.08  | 8.49 ± 0.13  | 94.5  | 10 | 8.97 ± 0.37  | 18.55 ± 1.23 | 95.8  |
| Lsy    | 20 | 25.89 ± 0.54 | 45.17 ± 1.61 | 96.4  | 10 | 34.15 ± 1.37 | 44.64 ± 2.85 | 104.9 |
| His    | 20 | 158.0 ± 7.0  | 178.2 ± 9.5  | 101.0 | 1  | 3.55 ± 0.29  | 4.49 ± 0.52  | 94.0  |
| Tyr    | 20 | 10.33 ± 0.67 | 31.76 ± 0.76 | 107.2 | 1  | 8.49 ± 0.23  | 9.48 ± 0.63  | 99.0  |
| δ-HLys | 20 | 10.71 ± 0.05 | 28.73 ± 0.94 | 90.1  | 1  | 0.54 ± 0.05  | 1.46 ± 0.05  | 92.0  |
| Trp    | 20 | 16.39 ± 0.22 | 36.01 ± 2.39 | 98.1  | 1  | 6.47 ± 0.30  | 7.38 ± 0.19  | 91.0  |
| CTH    | 2  | 5.01 ± 0.06  | 7.04 ± 0.31  | 101.5 | 1  | 3.32 ± 0.10  | 4.23 ± 0.36  | 91.0  |
| Cyt    | 20 | 10.33 ± 0.40 | 30.05 ± 1.24 | 98.6  | 1  | 0.51 ± 0.04  | 1.55 ± 0.10  | 104.0 |
| HCyt   | 2  | 6.44 ± 0.04  | 8.27 ± 0.05  | 91.5  | 1  | ND           | 1.08 ± 0.02  | 108.0 |

<sup>a</sup>Mean ± SD (*n* = 3).

<sup>b</sup>Not detectable (from **ref. 41**, with permission).

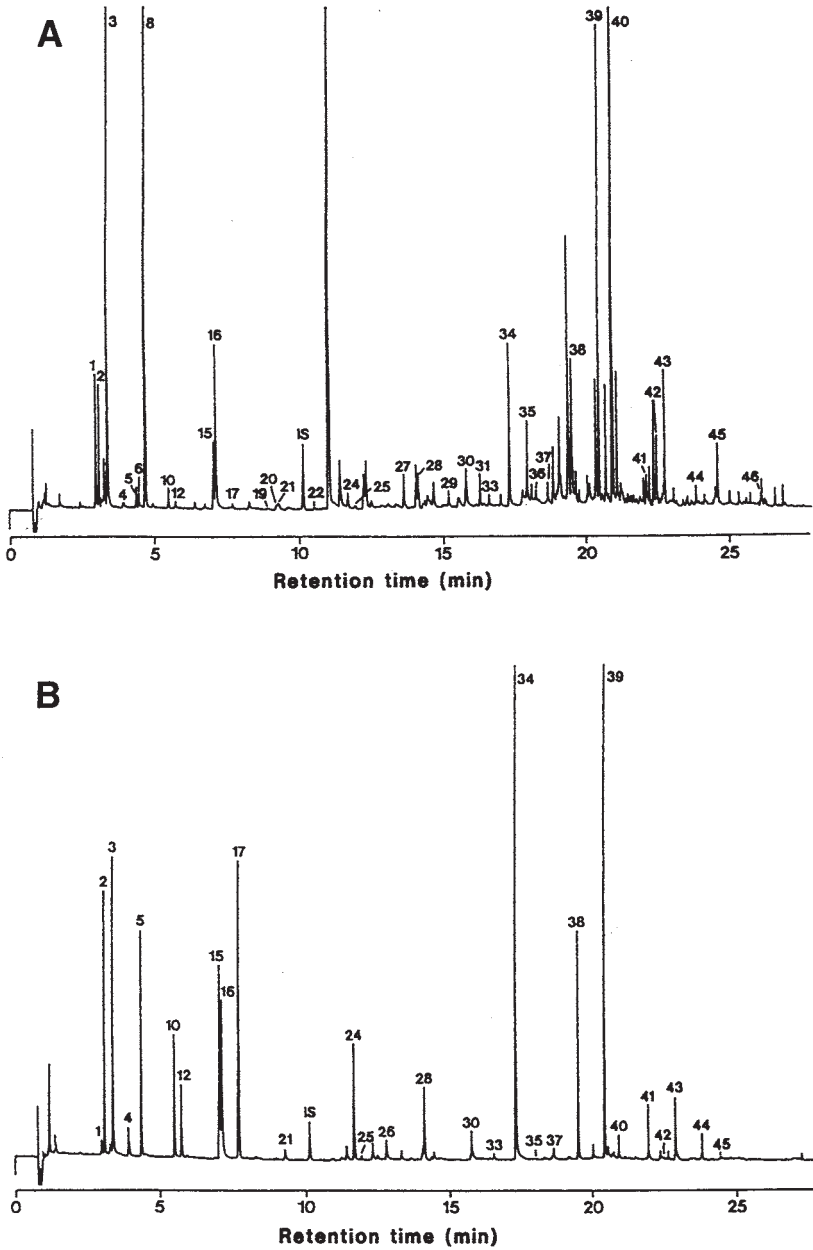


Fig. 7. Typical chromatograms obtained from (A) urine (25  $\mu$ L) and (B) serum (50  $\mu$ L) samples by NPD-GC. GC conditions and peak number: see Fig. 5.

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## Measurement of Blood Plasma Amino Acids in Ultrafiltrates by High-Performance Liquid Chromatography with Automatic Precolumn O-Phthaldialdehyde Derivatization

Hua Liu

### 1. Introduction

Amino acid analysis is an important technique that has many applications in biochemical, pharmaceutical, and biomedical fields. Profiling of plasma amino acids is also of great interest in clinical practice. Many diseases are known to be associated with disorders in amino acid metabolism. The analysis of amino acids offers the possibility of genetic prevention in both premarital and prenatal stages for those diseases resulting from inborn errors of metabolism. The plasma amino acid pattern has been also used to follow the course of prolonged dietary treatment (1,2).

For nearly 40 years, amino acid separations have been carried out mainly by an amino acid analyzer by means of ion-exchange chromatography and detected after postcolumn derivatization (1,3). The use of this analyzer has been widely advocated in the past, especially in routine application because of its high reliability (4). However, these analyzers are somewhat laborious, costly, time-consuming, and usually performed on dedicated instruments (1,5,6). In recent years, methods employing precolumn derivatization of amino acids, combined with reversed-phase (RP) high-performance liquid chromatography (HPLC) have gained increasing importance and have partially replaced the classical amino acid analyzer (6). Publications have demonstrated the usefulness of this technique for the determination of amino acids in physiological fluids, and the results achieved by HPLC methods compared favorably with those

obtained with the amino acid analyzer (6–14). In comparison with the postcolumn derivatization and ion-exchange chromatographic amino acid analyzers, precolumn derivatization and HPLC methods has the advantages of reducing analysis times, enhancing sensitivity and flexibility, and lowering cost of instrumentation and maintenance (7–16).

Among the methods of precolumn derivatization, *O*-phthaldialdehyde (OPA) has become the most popular derivatization reagent. In the presence of a thiol compound and at alkaline pH, OPA reacts with primary amino acids and forms highly fluorescent isoindole derivatives. This derivatization procedure is relatively easy and the reaction occurs rapidly at room temperature and in aqueous solution. No laborious purification procedures are required. The OPA derivatives are less polar than the original amino acids and can be well separated from each other by RP-HPLC (17,18). Moreover, the reagent itself does not fluoresce and consequently produces no interfering peaks.

One disadvantage of OPA derivatization has been the lack of stability of the OPA adducts when 2-mercaptoethanol (2-ME) is used as a sulfhydryl reagent (17,19). The reaction products are not stable and have a short half-life, possibly because of a spontaneous intramolecular rearrangement, with sulfur being displaced by oxygen from the ethanolic portion (19). Consequently, time differences between the reaction and injection during a manual procedure may cause significant errors in quantitation. An automated on-line OPA derivatization procedure is described in this chapter. By using an autoinjector, the exact time of each step from the beginning of reaction to injection can be controlled according to an injector program. The precision of the reaction time and sample volume in an automatic procedure eliminates the human errors that may occur during a manual derivatization. In addition, 2-ME was replaced by 3-mercaptopropionic acid (3-MPA) in our experiment with considerable improvement in stability because the stability of isoindoles formed. The other advantages of 3-MPA are that it is nonvolatile and less toxic because of its carboxylic moiety (20).

Another disadvantage of OPA derivatization is that it reacts only with primary amines, so secondary amino acids (imino acids) are not detected (19,21). 9-fluorenylmethyl-chloroformate (FMOC-Cl) is a highly reactive reagent that has been used as an amino-protective group in peptide synthesis. In 1983, Einarsson applied FMOC-Cl to a precolumn derivatization of amino acids (22). It was as sensitive as OPA, but reacted with secondary, as well as primary amine. The derivatives produced were stable and highly fluorescent. This method has been used to measure hydroxyproline, sarcosine and proline in serum, cerebrospinal fluid, and urine (22,23). FMOC-Cl itself is fluorescent and may obscure some amino acid peaks. Therefore, excess FMOC-Cl has to be removed by pentane extraction or reacted with hydrophobic amine 1-amin-



oadamantane (ADAM) to form an amine-FMOC complex (24). We obviated this problem by using a double-derivative technique (FMOC-Cl is incorporated to this automatic derivatization procedure as a second reagent for the derivatization of secondary amino acids) and using both photodiode array and programmable fluorescence detectors. This procedure has the advantage of different absorption and fluorescent spectra of OPA and FMOC-Cl derivatives. Therefore, both primary and secondary amino acids can be detected simultaneously.

Although there are many techniques available for the analysis of amino acids, and the sample can be analyzed quickly, accurately and sensitively, several precautions need to be taken in order to obtain reliable data on the concentration of plasma amino acids. These include the sample collection, centrifugation, storage conditions, and the deproteinization method. Deproteinization is one of the major problems in the analysis of amino acids in physiological fluid (25). The whole plasma contains soluble peptides and proteins that should be removed from the sample. Otherwise, these substances will clog the chromatographic column, increase instrumental backpressure, and interfere with separation.

The method used to prepare the plasma and to remove the plasma protein has a marked effect on the final results (1,3). The most widely used method of deproteinization is precipitation with 5-sulfosalicylic acid (SSA) followed by centrifugation to remove the precipitated protein (1). We have not had success in using SSA as the deproteinization agent for the analysis of plasma amino acids by an HPLC method with automatic precolumn OPA/3-MPA and FMOC-Cl derivatization (11). When this method was used for the analysis of plasma samples deproteinized by SSA, several major problems were encountered. First, the yield of the derivatization was low in the SSA supernatant. This is probably because the strong acidic nature of SSA inhibits the formation of OPA-amino acid derivatives, which require an alkaline pH. Second, the large peak of deproteinizing agent SSA superimposed the first three amino acid peaks of the chromatogram (*O*-phospho-l-serine (OPS), Asp and Glu). The SSA peaks were higher than 4000, 2000, and 300 mAu at the UV sample wavelengths 230, 260, and 338 nm, respectively. The third problem was the adverse affects of SSA supernatant on the separation and quantitation of other amino acids. Other investigators (11,18) have observed similar problems. When ethanol or methanol was used for the deproteinization, the sample was diluted and some of plasma amino acids with lower levels (OPS, Asp, Glu, AABA, and Trp) became undetectable by the UV detector at 338 nm. Other problems were the high level of organic solvent in the injected solution resulting in broad peaks in the early part of the chromatogram, and the increased volatility of the sample made it difficult to store for long periods of time (11).

Ultrafiltration has the advantage of achieving a protein-free sample without adding chemical agents, thus keeping the sample close to a physiological state. It is an attractive alternative to equilibrium dialysis because of the ease and speed with which it can be accomplished. Another potential benefit of ultrafiltration is that the platelets and leukocytes are removed from the plasma, and thus the contamination of amino acids from these blood components can be eliminated. The ultrafiltration methods are not widely applied in the ion-exchange chromatography of amino acids because, for unknown reasons, they decrease the retention time during chromatographic separation and consequently leads to distorted separations of critical pairs of amino acids (*I,3*).

Ultrafiltration has been chosen for the preparation of protein-free sample in the analysis of amino acid by HPLC methods with precolumn derivatization (*18,26,27*). The results indicate that ultrafiltration of plasma may replace chemical deproteinization in the HPLC analysis of free amino acids. We investigated several factors that may have an influence on the final results of the ultrafiltration (*11*). These conditions were then standardized in our procedure. Satisfactory results were achieved for the analysis of plasma amino acids by the automatic precolumn OPA/3-MPA and FMOC-Cl derivatization and RP-HPLC method.

It appears that the combination of: (1) automatic on-line derivatization (2) the improvement in the stability of the reaction by using 3-MPA (3) the use of OPA and FMOC-Cl as dual derivatizing reagents (4) the optimized gradient elution program (5) and the simple mobile phase composition render this method suitable for the quantitative analysis of plasma amino acids in a clinical laboratory. This procedure can yield more rapid and sensitive results than that of classic amino acid analyzer, with comparable accuracy and precision. We have used this method for the determination of plasma amino acids in more than 3000 plasma samples during a period of 10 yr with satisfactory results (*see Notes 1 and 2*).

## 2. Materials

### 2.1. Equipment

1. A Hewlett-Packard HP 1090 *M* series HPLC system (*see Note 3*). The system consists of a DR 5 solvent delivery system with 3 solvent channels, a variable volume autoinjector and an autosampler, an HP 1040A photo-diode array UV detector, and HP 1046 programmable fluorescence detector (*see Note 4*).
2. An HP 79994A analytical workstation for data processing.
3. Two HP Hypersil-ODS 5- $\mu$ m columns (100  $\times$  2.1 mm).
4. One guard column (20  $\times$  2.1 mm) (*see Note 5*).

## 2.2. Reagents

Water, methanol, and acetonitrile were HPLC grade (Curtin Matheson Scientific, Inc., Houston, TX). Other chemicals used were analytical grade including sodium acetate, glacial acetic acid, boric acid, sodium hydroxide, *O*-phthalaldehyde, 3-mercaptopropionic acid, and 9-fluorenylmethyl-chloroformate (all from Sigma Chemical Company, St. Louis, MO).

## 2.3. Solutions

1. Standard solution: An amino acid standard solution containing 29 amino acids was prepared by adding crystalline *O*-phospho-L-serine (OPS), reduced glutathione (GSH), asparagine, glutamine, citrulline, taurine,  $\alpha$ -amino-N-butyric acid (AABA), tryptophan, ornithine, hydroxy-proline (Hyp), and sarcosine to a 50% methanol solution. Then this solution (containing 11 amino acids) was mixed with a commercial AA-S-18 Amino Acid Standard Solution (from Sigma). Nor-Valine (N-Val) can be used as an internal standard (United States Biochem Corp, Cleveland, OH). The concentration of working standard was 250  $\mu\text{mol/L}$  for each amino acid. The mixture solution was kept in  $-76^\circ\text{C}$  (see **Note 6**).
2. OPA derivatization reagent: The OPA derivatization reagent was prepared by dissolving 3 mg of OPA in 50  $\mu\text{L}$  of methanol, adding 450 mL of sodium borate buffer (0.5 mol/L, pH 10.2) and 5  $\mu\text{L}$  of 3-MPA. Borate buffer was prepared from 0.5 M boric acid solution adjusted to pH 10.2 with 5 M sodium hydroxide solution. This OPA solution was placed in an amber crimp top vial with a silicone rubber PTFE-coated cap and kept in the dark at  $-20^\circ\text{C}$ . Fresh solution was prepared each week.
3. FMOC derivatization reagent: FMOC-Cl solution was prepared by dissolving 1.29 mg of FMOC-Cl in 1 mL of acetonitrile and stored at  $-20^\circ\text{C}$  (see **Note 7**).
4. Mobile phase solution: The sodium acetate buffer (0.1 M) in mobile phase was prepared by dissolving sodium acetate in HPLC-grade water and titrating to pH 6.8 with glacial acetic acid. The buffer was then diluted to 0.015 M and 0.01 M for mobile phase solutions A and C, respectively (see **Note 8**). The mobile phases were filtered by passing through a 0.45- $\mu\text{m}$  Durapore membrane filter (Millipore Inc., Milford, MA) and continuously degassed by helium.
5. Blood sample: Blood samples were collected from subjects into a heparinized vacutainer tube (see **Note 9**). The plasma samples were ultrafiltrated by using the Centrifree System (Amicon, Beverly, MA).

## 3. Methods

1. Collect 1.5 mL of venous blood samples from subjects by venipuncture between 7 AM to 9 AM (see **Note 10**) after overnight fasting and put into a vacutainer tube containing heparin (see **Note 11**). Centrifuge the blood samples at 2000g for 15 min at  $10^\circ\text{C}$  (see **Note 12**) and remove the plasma.
2. Ultrafiltrate the heparinized plasma samples by using the commercially available Centrifree System (see **Notes 13** and **14**). During the ultrafiltration, the sample is

**Table 1**  
**Injector Program<sup>a</sup>**

| Line# | Function | Amount                     | Vial No. | Reagent   |
|-------|----------|----------------------------|----------|-----------|
| 1     | Draw :   | 0.0 $\mu\text{L}$ from :   | Vial#:4  | (Water)   |
| 2     | Draw :   | 2.5 $\mu\text{L}$ from :   | Vial#:5  | (OPA)     |
| 3     | Draw :   | 0.0 $\mu\text{L}$ from :   | Vial#:4  |           |
| 4     | Draw :   | 2.5 $\mu\text{L}$ from :   | Vial#:X  | (Sample)  |
| 5     | Mix :    | 5.0 $\mu\text{L}$ cycles 2 |          |           |
| 6     | Draw :   | 0.0 $\mu\text{L}$ from :   | Vial#:4  |           |
| 7     | Draw :   | 1.0 $\mu\text{L}$ from :   | Vial#:8  | (FMOC-Cl) |
| 8     | Mix :    | 6.0 $\mu\text{L}$ cycles 2 |          |           |
| 9     | Wait :   | 2.5 min                    |          |           |
| 10    | Inject   |                            |          |           |

<sup>a</sup>Reprinted from **ref. 8**, p. 3326 by courtesy of Marcel Dekker, Inc.

**Table 2**  
**Time Table for Gradient Elution**

| Time (Minute) | Solvent A% | B%   | C%  |
|---------------|------------|------|-----|
| 0.05          | 100        | 0    | 0   |
| 15.00         | 60         | 40   | 0   |
| 18.50         | 57.5       | 42.5 | 0   |
| 22.00         | 45         | 55   | 0   |
| 25.00         | 0          | 0    | 100 |
| 30.00         | 0          | 0    | 100 |

Solvent A: 0.015 *M* NaAc buffer (pH 6.8).

Solvent B: Methanol.

Solvent C: 0.010 *M* NaAc buffer (pH 6.8).

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deproteinized by filtration of plasma through the ultrafiltration membrane. The protein is retained by the membrane, whereas the ultrafiltrate (containing free amino acids) pass through and collect in the filtrater cup. Put a volume of 0.4 mL of plasma in the sample reservoir, and then place the device in a centrifuge with a 45° fixed-angle rotor (*see Note 15*). About 80  $\mu\text{L}$  of ultrafiltrates should be collected after centrifugation at 1000*g* for 15 min. Store the ultrafiltrates at -80°C until analyzed (*see Note 16*).

- To perform the automatic precolumn derivatization procedure using an injector program (**Table 1**), place the OPA-reagent at vial number 5 and the FMOC-Cl reagent at vial number 8. First, 2.5  $\mu\text{L}$  of the OPA reagent is drawn into the sample loop. Then, 2.5  $\mu\text{L}$  of sample is drawn into the loop and mixed with the OPA reagent. Finally, 1  $\mu\text{L}$  of FMOC-Cl reagent is drawn and mixed in the sample

- loop. After waiting 2.5 min for the reaction, a total 6  $\mu\text{L}$  of sample and reagents mixture is injected into the column for gradient elution (*see Note 17*). After drawing from sample or reagents, the needle of the injector should always be dipped into 1 mL of water (vial number 4) for cleaning.
4. The mobile phases are 0.015 M sodium acetic buffer (pH 6.8, for solvent A), methanol (solvent B) and 0.015 M of sodium acetic buffer (pH 6.8, for solvent C) (*see Notes 18 and 19*). The separation of amino acids in both the standard solution containing 29 amino acids or plasma samples is carried out by a gradient elution according to a chromatographic time-table (**Table 2**). The flow-rate is 0.3 mL/min and the stop time is 30 min after the injection. Place two HP Hypersil-ODS 5- $\mu\text{m}$  columns (100  $\times$  2.1-mm id) in series in a thermostatically controlled column compartment preceded by a guard column (20  $\times$  2.1-mm id). The column temperature should be maintained at 40°C for the separation of amino acids (*see Note 20*).
  5. For the detection of amino acid derivatives, set the photo-diode array detector at three sample wavelengths: 338, 266, and 230 nm with bandwidths of 10, 4, and 4 nm, respectively (*see Note 21*). The reference wavelength is 550 nm with a bandwidth of 100 nm. The initial parameters for the fluorescence detector are excitation wavelength (Ex) 230 nm and emission wavelength (Em) 450 nm. Twenty minutes after the injection, change the Ex and Em to 260 nm and 315 nm, respectively, for the determination of secondary amino acids proline and hydroxyproline (*see Note 22*).
  6. Amino acid peaks are identified with reference to retention times of standard amino acids injected. Coinjection of standard amino acids and the plasma samples may be needed to identify the amino acid peaks in some samples. Chromatograms of an amino acids standard mixture and a representative plasma sample (*see Note 11*) are demonstrated in **Figs. 1 and 2**. These chromatograms showed a satisfactory separation of 29 primary and secondary amino acids.
  7. The precision of analysis is observed from the reproducibility of the peak areas of eight consecutive injections of 29 amino acid mixture solution. The results are listed in **Table 3**. The coefficients of variation for peak areas ranged from 0.78% to 2.92%, with a mean of  $1.73\% \pm 0.67\%$  SD (*see Note 23*). The high precision of this method would allow analysis without an internal standard for quantitation. The amino acid standard should be first analyzed twice for calibration and thereafter every eighth analysis in an automated series.

#### 4. Notes

1. The levels of 25 plasma amino acids from 75 boys and 85 girls are listed in **Table 4**. In general, amino acid levels of boys were higher than that of girls. The mean values of aspartic acid, methionine, isoleucine, and hydroxy-proline in boys were significantly higher than that in girls ( $p < 0.01$ ). The girls had significant higher level of histidine than boys ( $p < 0.001$ ). Compared with boys below 6 yr old ( $n = 23$ ), boys over 6 yr old ( $n = 52$ ) showed significant higher value of glycine, threo-

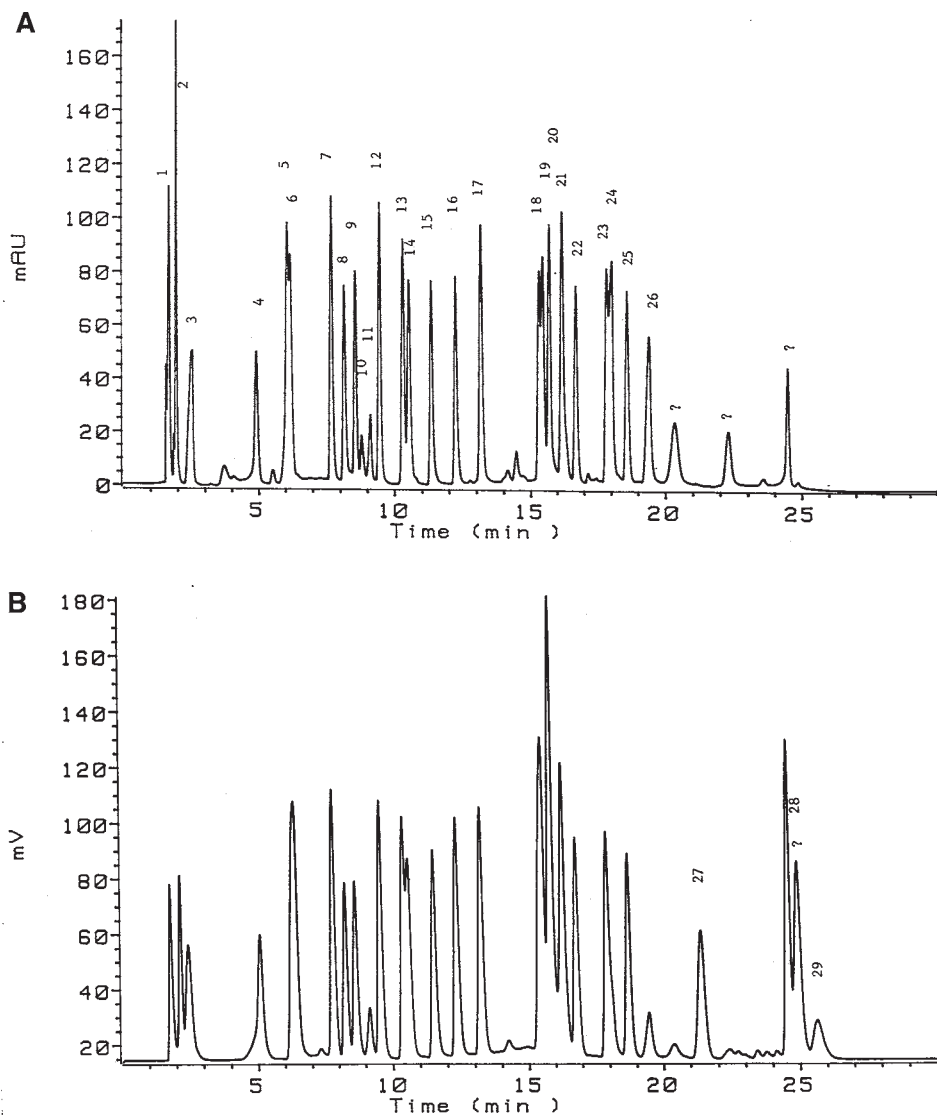


Fig. 1. Typical chromatograms showing the separation of 29 OPA/3-MPA and FMOC-Cl derivatized standard amino acids (500  $\mu\text{mol/L}$ ). (A) Signal from UV detection at 338 nm. (B) Signal from Fluorescence detection. For chromatographic conditions, see **Subheadings 2** and **3**. Peaks: 1 = *O*-phospho-L-serine, 2 = Aspartic Acid, 3 = Glutamic Acid, 4 = Glutathione (reduced), 5 = Asparagine, 6 = Serine, 7 = Glutamine, 8 = Glycine, 9 = Threonine, 10 = Histidine, 11 = Cystine, 12 = Citrulline, 13 = Taurine, 14 = Alanine, 15 = Arginine, 16 = Tyrosine, 17 = Alpha-amino-*N*-butyric Acid, 18 = Methionine, 19 = Valine, 20 = Nor-Valine, 21 = Tryptophan, 22 = Phenylalanine, 23 = Isoleucine, 24 = Ornithine, 25 = Leucine, 26 = Lysine, 27 = Hydroxy-proline, 28 = Sarcosine, 29 = Proline.

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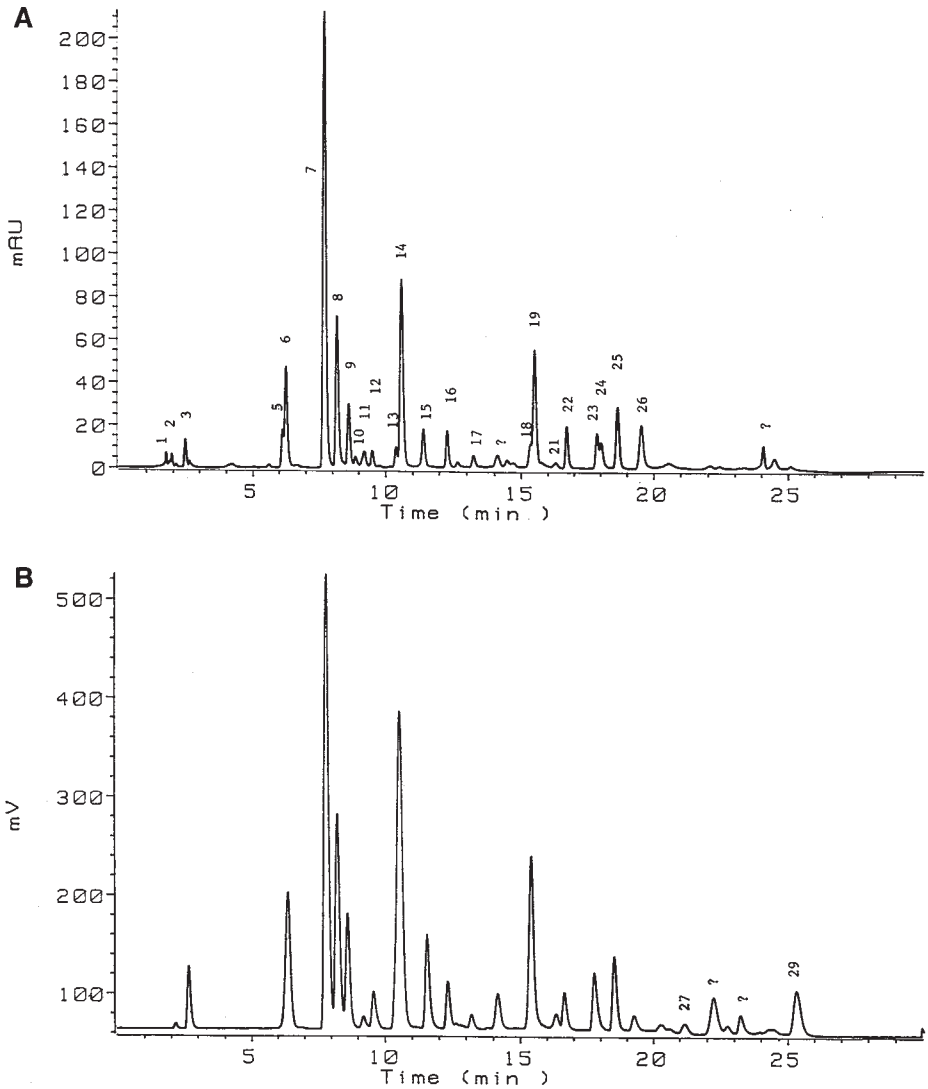


Fig. 2. Chromatograms of a representative plasma sample under the identical experimental conditions as in Fig. 1. (A) Signal from UV detection at 338 nm. (B) Signal from fluorescence detection. Peak numbers are identified in Fig. 1A and B.

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nine, histidine, taurine, alanine, arginine, tyrosine, valine, phenylalanine, isoleucine, ornithine and leucine. For girls over 6 yr old ( $n = 56$ ), only threonine, histidine, and alanine were significantly higher than that for girls below 6 yr old ( $n = 29$ ) (8).

**Table 3**  
**Reproducibility of Peak Areas for Standard Amino Acids**

| Amino Acid <sup>a</sup> | Mean ( <i>n</i> = 8) | SD    | CV%  |
|-------------------------|----------------------|-------|------|
| OPS                     | 527.6                | 5.47  | 1.03 |
| Asp                     | 528.1                | 4.13  | 0.78 |
| Glu                     | 545.4                | 15.49 | 2.84 |
| GSH                     | 532.4                | 13.69 | 2.57 |
| Asn                     | 581.9                | 16.58 | 2.84 |
| Ser                     | 694.3                | 8.96  | 1.29 |
| Gln                     | 676.1                | 8.99  | 1.32 |
| Gly                     | 498.8                | 9.97  | 1.99 |
| Thr                     | 536.4                | 10.78 | 2.00 |
| His                     | 106.0                | 1.59  | 1.50 |
| Cys                     | 170.8                | 3.94  | 2.30 |
| Cit                     | 637.7                | 4.88  | 0.76 |
| Tau                     | 582.4                | 5.76  | 0.98 |
| Ala                     | 589.0                | 5.22  | 0.88 |
| Arg                     | 554.4                | 7.37  | 1.32 |
| Tyr                     | 553.7                | 8.14  | 1.47 |
| AABA                    | 707.5                | 13.78 | 1.94 |
| Met                     | 548.4                | 10.52 | 1.91 |
| Val                     | 660.3                | 7.69  | 1.16 |
| N-Val                   | 829.4                | 12.16 | 1.46 |
| Trp                     | 638.4                | 16.49 | 2.58 |
| Phe                     | 535.1                | 11.55 | 2.15 |
| Ile                     | 646.1                | 18.51 | 2.86 |
| Orn                     | 540.6                | 15.80 | 2.92 |
| Leu                     | 545.1                | 10.01 | 1.83 |
| Lys                     | 587.8                | 9.54  | 1.62 |
| Hyp                     | 1469                 | 28.91 | 1.91 |
| Sar                     | 2427                 | 20.85 | 0.85 |
| Pro                     | 645.9                | 8.86  | 1.37 |

<sup>a</sup>OPS = O-phospho-L-serine, Asp = Aspartic Acid, Glu = Glutamic Acid, GSH = Glutathione (reduced), Asn = Asparagine, Ser = Serine, Gln = Glutamine, Gly = Glycine, Thr = Threonine, His = Histidine, Cys = Cystine, Cit = Citrulline, Tau = Taurine, Ala = Alanine, Arg = Arginine, Tyr = Tyrosine, AABA = Alpha-amino-N-butyric Acid, Met = Methionine, Val = Valine, N-Val = Nor-Valine, Trp = Tryptophan, Phe = Phenylalanine, Ile = Isoleucine, Orn = Ornithine, Leu = Leucine, Lys = Lysine, Hyp = Hydroxy-proline, Sar = Sarcosine, Pro = Proline.

- The plasma amino acid levels of 160 children obtained by this method showed a good agreement with the reference values reported previously by using ion-exchange chromatography (28). The findings about the difference between amino acid levels of boys and girls and between the children with different age were



**Table 4**  
**Free Plasma Amino Acid Levels (mmol/L) of 160 Children**

| Amino Acid <sup>a</sup> | Girls (n = 85)     |   |       | Boys (n = 75)      |   |       |
|-------------------------|--------------------|---|-------|--------------------|---|-------|
|                         | Mean               | ± | SD    | Mean               | ± | SD    |
| Asp                     | 6.19               |   | 1.86  | 7.62 <sup>b</sup>  |   | 2.13  |
| Glu                     | 39.28              |   | 13.95 | 48.23              |   | 17.84 |
| Asn                     | 53.99              |   | 10.16 | 56.53              |   | 10.92 |
| Ser                     | 126.30             |   | 23.57 | 133.50             |   | 28.79 |
| Gln                     | 532.80             |   | 82.00 | 540.70             |   | 93.75 |
| Gly                     | 242.50             |   | 47.29 | 262.60             |   | 59.35 |
| Thr                     | 142.40             |   | 27.86 | 146.20             |   | 28.87 |
| His                     | 96.09 <sup>c</sup> |   | 16.51 | 85.07              |   | 16.95 |
| Cys                     | 88.96              |   | 14.91 | 87.21              |   | 15.29 |
| Cit                     | 24.29              |   | 4.10  | 26.00              |   | 4.15  |
| Tau                     | 41.57              |   | 7.73  | 44.35              |   | 11.53 |
| Ala                     | 423.30             |   | 82.15 | 425.10             |   | 96.85 |
| Arg                     | 95.05              |   | 18.18 | 100.60             |   | 19.52 |
| Tyr                     | 74.88              |   | 15.45 | 73.99              |   | 13.51 |
| AABA                    | 18.23              |   | 3.52  | 18.66              |   | 3.93  |
| Met                     | 23.88              |   | 2.60  | 25.17 <sup>b</sup> |   | 3.65  |
| Val                     | 221.80             |   | 39.27 | 235.20             |   | 49.11 |
| Trp                     | 5.67               |   | 2.73  | 6.38               |   | 3.33  |
| Phe                     | 64.94              |   | 11.15 | 65.32              |   | 5.96  |
| Ile                     | 66.36              |   | 15.35 | 74.28 <sup>b</sup> |   | 19.63 |
| Orn                     | 42.11              |   | 12.76 | 45.03              |   | 14.53 |
| Leu                     | 137.90             |   | 25.92 | 144.30             |   | 32.03 |
| Lys                     | 130.00             |   | 25.99 | 126.10             |   | 26.91 |
| Hyp                     | 18.96              |   | 4.20  | 21.34 <sup>b</sup> |   | 5.65  |
| Pro                     | 153.00             |   | 42.24 | 149.90             |   | 45.66 |

<sup>a</sup>Amino acid abbreviations as in **Table 3**.

<sup>b</sup>Significantly higher than girls ( $p < 0.01$ ).

<sup>c</sup>Significantly higher than boys ( $p < 0.001$ ).

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also similar to the results observed by Armstrong and Stave (**28,29**). Patients suffered from hyperlysinemia, nonketotic hyperglycinemia, disorders of branched-chain amino acid metabolism, and argininosuccinase deficiency have been detected by this procedure.

3. This instrument is a low-pressure gradient system. A high-pressure gradient system can be used for the analysis. We recommend that the buffers in a mobile phase be premixed with organic solvents (5% to 10% buffer in organic solvent and 5% to 10% organic solvent in buffer solution), and thorough degassed be performed when a high-pressure gradient system is being used.

4. The contribution to dispersion by the instrument itself must be minimized. Two detectors should be physically close each other. The internal diameter of the capillaries should be as small as possible.
5. When contamination was detected (baseline drift during the gradient elution) or separation deteriorated, and the change of guard column could not correct the problem, the first column (next to the guard column) was discarded, the second one moved to the first position, and a new column was added at second position. Under a normal situation, the guard or analytical column can be used for analysis up to 100 or 500 plasma samples, respectively.
6. Other individual amino acid (such as argininosuccinic acid) can be added to this amino acid standard mixture for peak identification and quantitation.
7. The purity of the reagent was important for extending detection limit. A previously unused and aged reagent (> 24-h old) was necessary for a higher sensitivity level detection (21). The sensitivity depended mainly on the ability to eliminate or subtract background levels and reduce the interfering substances present in solvents and reagents.
8. It is recommended that the concentration of the buffer solutions in the mobile phase be less than 0.1 M to avoid the potential precipitation during the gradient elution.
9. A siliconized tube is recommended to prevent blood platelets becoming activated with the resultant release of Tau and phosphoethanolamine (PEA). Hemolysis should also be prevented because it may lead to false increases in the concentrations of Asp, Glu, Tau, and PEA (4).
10. The time of blood withdrawal and the relation to dietary intake may be another reason that may cause varying results between laboratories. If it is possible, the blood sample should be collected between 8 and 9 AM before breakfast.
11. Many analytical factors could result in variation in the plasma amino acid profile. These include: (1) sample collection (2) centrifugation and ultrafiltration (3) storage conditions (4) contamination by platelets and leukocytes or hemolysis (5) shift in baseline during chromatographic analysis, and (6) interference or overlap of one or more amino acids. The physiological factors such as circadian rhythm, protein intake, pregnancy, nutritional status of the subject, physical activity before blood collecting, menstrual cycle, sex, and age differences and medication that interferes with the analysis may also have influence on plasma amino acids. All of these factors should be controlled to assess with certainty the clinical data under various pathological conditions.
12. Although most of the amino acids remained stable for up to 30 min at room temperature, delayed deprotenization may cause an increase of Asp, Glu, Gly, Ala, and Orn, whereas there is a decrease of Asn, Gln, His, and 3-MH (2). When deproteination of physiological samples cannot be done immediately after centrifugation, it is recommended to store the specimens at temperature lower than  $-18^{\circ}\text{C}$  to prevent further hydrolysis of protein. In particular, the amino acids Asp and Glu increased markedly when the samples were not frozen immediately and not stored at  $-68^{\circ}\text{C}$  or lower.

13. In dealing with physiological fluid samples, the protein should be completely removed from the sample prior to analysis. Direct injection onto RP columns are known to affect the retention behavior of the solutes and cause increase back pressure by irreversible adsorption of protein to the stationary phase.
14. Precaution should be taken when this method is used for some samples with higher protein concentration, such as blood cell lysates or tissue extracts. It has been reported that for the lipemia samples, the time required to filter sufficient sample was very variable and occasionally no filtrate could be obtained, presumably because the membrane pores got blocked. The alternative method is chemical precipitation by using various acids, bases, or organic solvents. Precaution should be taken to avoid hydrolysis, degradation, and oxidation of amino acids in selecting any such procedure for the deproteinization. The main factors to be considered are the amount of protein to be removed, the comparability or interference with the chromatographic analysis, the thermal and chemical stability, the solubility of the analytes, and finally, the cost, time, and labor. The use of 24–30% SSA to deprotein 500- $\mu$ L plasma samples (w/v) has been reported yield consistent results with high recoveries of most amino acids (2,4). Ethanol or acetonitrile (ACN) can also be used as precipitants (ACN or ethanol to plasma ratio 2–4:1, v/v) and has the advantage of resulting in full recovery of total Trp, whereas using SSA precipitation cannot completely recover this amino acid (4).
15. Several factors such as rotor, membrane, and the time of the ultrafiltration may have an influence on the final results of the ultrafiltration (II). These factors were investigated in our experiment. The recovery rates of amino acid in ultrafiltrates obtained by fixed-angle rotor were higher than that by swinging-bucket rotor. This may be caused by the different polarization control between fixed-angle and swinging-bucket rotors. The use of a fixed-angle rotor provides polarization control. The angle counteracts the buildup of retained protein at the membrane surface, because this dense layer slides outward and accumulates at the edge of membrane. In a swinging-bucket rotor, the polarization layer is compacted over the entire membrane surface, restricting the passage of solute and solvents through the membrane. Our experiment suggests that the polarization occurring during the ultrafiltration may be one of the important factors that can influence the recovery rate of amino acids. This might be the reason why some other authors could not achieve satisfactory results by ultrafiltration, because it was difficult to control the polarization when ultrafiltration was performed under nitrogen pressure, or a syringe was used as the driving force for the ultrafiltration.
16. The recovery from ultrafiltration and its reproducibility of standard amino acids were tested by 10 aliquots of amino acid standards: 5 aliquots were analyzed by HPLC before ultrafiltration, another 5 aliquots were analyzed after ultrafiltration. The results showed that the recovery of all of the standard amino acids was excellent, ranging from 95–102% (II). The accuracy of measurement was tested by adding a known quantity of amino acid standards to a plasma sample, then the sample was ultrafiltered and derivatized for analysis. The analytic recovery rate

**Table 5**  
**Recovery of Amino Acid Standard Added to Plasma**

| Amino Acid <sup>a</sup> | Plasma alone (μmol/L) | Plasma+250 mmol/L (expected) | Actual | Recovery Rate % |
|-------------------------|-----------------------|------------------------------|--------|-----------------|
| Ops                     | 32.65                 | 282.65                       | 271.87 | 97              |
| Asp                     | 18.98                 | 268.98                       | 273.34 | 101             |
| Glu                     | 28.76                 | 278.76                       | 290.15 | 104             |
| GSH                     | —                     | 250.00                       | 80.73  | 32              |
| Asn                     | 35.89                 | 285.89                       | 284.52 | 100             |
| Ser                     | 128.27                | 378.27                       | 401.06 | 106             |
| Gln                     | 538.74                | 788.74                       | 757.19 | 96              |
| Gly                     | 219.74                | 469.74                       | 465.04 | 100             |
| Thr                     | 132.14                | 382.14                       | 384.32 | 100             |
| His                     | 31.71                 | 281.71                       | 290.16 | 103             |
| Cys                     | 64.12                 | 314.12                       | 307.83 | 98              |
| Cit                     | 18.36                 | 268.36                       | 287.14 | 107             |
| Tau                     | 38.85                 | 288.85                       | 291.73 | 101             |
| Ala                     | 376.32                | 626.32                       | 613.79 | 98              |
| Arg                     | 103.50                | 353.50                       | 352.22 | 100             |
| Tyr                     | 62.38                 | 312.38                       | 303.01 | 97              |
| AABA                    | 29.09                 | 279.09                       | 287.46 | 103             |
| Met                     | 19.60                 | 269.60                       | 258.82 | 96              |
| Val                     | 170.04                | 420.04                       | 407.73 | 97              |
| N-Val                   | —                     | 250.00                       | 252.36 | 101             |
| Trp                     | 14.26                 | 264.26                       | 214.05 | 81              |
| Phe                     | 49.82                 | 299.82                       | 297.42 | 99              |
| Ile                     | 60.21                 | 310.21                       | 313.08 | 100             |
| Orn                     | 50.18                 | 300.18                       | 285.17 | 95              |
| Leu                     | 100.92                | 350.92                       | 349.15 | 100             |
| Lys                     | 137.24                | 387.24                       | 385.69 | 99              |
| Hyp                     | 24.49                 | 274.49                       | 277.23 | 101             |
| Pro                     | 180.86                | 430.85                       | 439.46 | 102             |

<sup>a</sup>Amino acid abbreviations as in **Table 3**.

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of each amino acid was calculated after HPLC quantitation (**Table 5**). It should be mentioned that when standard amino acids were added to the plasma, Trp and GSH (a peptide) showed poor recovery (81% and 32%, respectively), whereas all of the other amino acids remained at a similar recovery. The reason was unknown. Changing the membrane in the ultrafiltration system to a membrane with higher molecular weight cutoff might be helpful for the improvement of the recovery rate of Trp and GSH.

The multiplier in calibration form was set at 1.19 for the quantitation of plasma tryptophan. No effort has been made for the determination of GSH in plasma.

17. Although the stability of OPA/3-MPA derivative is better than OPA/2-ME derivative, the OPA/3-MPA derivative of various amino acids does not have the same stability (8). When the waiting time was increased beyond 2.5 min, the stability of various amino acids was different. The UV absorbency of GSH and OPS increased, whereas asparagine, glutamine, glycine, histidine, taurine, ornithine, and lysine decreased. Others were unchanged. Therefore, the precise control of reaction time is still important in the derivatization using OPA/3-MPA as reagents. The results from our experiment indicated that OPA/3-MPA derivatives were relative stable during the waiting time from 0.5 to 5 min (8).
18. When analyzing physiological samples with interested amino acids covering a broad retention range, it is necessary to modify the elution conditions during the analysis to optimize the separation. The combinations of the organic solvents and buffer solution; the pH and ion concentration of the buffer in the mobile phase; the patterns of gradient elution, and the different column and column temperature should be investigated to optimize the separation.
19. Comparison of buffer pH from 6.4 to 7.2 was made in our experiment with all other conditions held constant (8). The pH at 6.8 appeared to give optimal separation. The pH of the buffer in the mobile phase should be close to neutral. A lower pH of the buffer would result in weaker UV absorbency and fluorescent intensity.
20. Maintaining a stable column temperature is critical to the reproducibility of retention time, in addition to improving the separation of amino acids. When the column temperature was kept at 40°C, the coefficients of variation for retention time ranged from 0.02% to 0.97%, with a mean of 0.26%+0.21 SD. Column temperature was also tested for its effect on separation. When column temperature was maintained at 30°C, the separation of amino acids at first half of the chromatogram was better than that at temperature 35°C or 40°C, but the separation of second half was not satisfactory. Increasing the temperature to 40°C gave better separation for amino acids eluted later and the overall results were the best, together with a lower-column pressure. We therefore set the column temperature at 40°C for the analysis (8).
21. The OPA/3-MPA and FMOC-Cl derivatives were detected in our experiment by a photo-diode array detector at three different sample wavelengths. The intensity of the signals, the baseline noises, and interference were different at various wavelength settings. At sample wavelengths of 338 nm and 266 nm, we compared two reference wavelengths at 390 nm and 550 nm. The use of 550 nm as reference wavelength produced more stable baseline than that at 390 nm. Therefore, 550 nm was used as reference wavelength for the photo-diode array detector. Under the chromatographic conditions we had chosen, there was no interfering peak on either UV or fluorescence signals during a blank solvent gradient elution. When HPLC water was used as a blank sample, there was no peak interfering with amino acid peak at UV sample wavelength 338 nm, and fluorescence signals. But at UV 266 nm, and especially at 230 nm, several peaks were

observed that were large enough and close enough to interfere with about 10 amino acid peaks.

Although detection at UV 230 nm was more sensitive than at 338 nm, the interference could seriously compromise the quantitation. For this reason, UV 338 nm is recommended as the main sample wavelength for the quantitation of primary amino acids whereas wavelength 266 nm is recommended for the FMOC-Cl derivatives of secondary amino acids. Extremely high sensitivity is not required for the analysis of plasma amino acids, because the limitation is not sample size, rather the amount of plasma required for deproteinization procedure (such as ultrafiltration).

22. For the fluorescence detector we used to detect OPA and FMOC-Cl derivatives, both 230 nm and 340 nm were evaluated as Ex for OPA primary amino acid derivatives. The results showed that the detection of OPA primary amino acid derivatives at Ex 230 nm gave a response over seven times stronger than that at 340 nm, with an acceptable baseline. For the detection of FMOC-Cl secondary amino acid derivatives, three excitation wavelengths of 254, 260, and 266 nm were compared. The results showed 260 nm had the strongest signal. Therefore, we set Ex 230 nm and Em 450 nm during the first 20 min for the measurement of OPA derivatives of primary amino acids, then switched to Ex 266 nm and Em 315 nm for the determination of FMOC-Cl derivatives of secondary amino acids. The hydroxyproline, sarcosine, and proline were all eluted after 20 min in our chromatographic system.
23. Analysis of physiological samples with different amino acid concentrations is required during the practical applications. The linear relationship of amino acid concentrations in the ultrafiltrates was investigated. The linearity of response was estimated by injecting derivatized amino acids with different concentrations and constructing regression equations for UV and fluorescence response-concentration curves. The linear relationship between the concentration and peak areas of each standard amino acid was determined by analyzing the standard amino acid mixture at concentrations ranging from 31.25 to 500  $\mu\text{mol/L}$  ( $n = 5$ , by serial dilution). These concentrations cover the normal range of most plasma amino acids. For plasma, the original plasma sample and plasma samples diluted by HPLC water to 75%, 50%, and 25% ( $n = 4$ ) of the plasma were analyzed. The linear regression analysis showed satisfactory coefficients of correlation ( $>0.99$ ) between the concentration and peak areas of each amino acid from both UV and fluorescent signals and in both standard amino acids and plasma samples (11).

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# Determination of Amino Acids in Foods by Reversed-Phase High-Performance Liquid Chromatography with New Precolumn Derivatives, Butylthiocarbamyl, and Benzylthiocarbamyl Derivatives Compared to the Phenylthiocarbamyl Derivative and Ion Exchange Chromatography

Kang-Lyung Woo

## 1. Introduction

Amino acid analysis with reverse-phase high-performance liquid chromatography (RP-HPLC) and ultraviolet (UV) detection following precolumn derivatization is popular owing to the greater versatility at the instrument, sensitivity and speed of analysis compared to specialized ion-exchange amino acid analyzers. Phenylthiocarbamyl (PTC) amino acid derivative is a precolumn derivatization method that has been widely used for analysis of amino acid by RP-HPLC (1–8). This method is an excellent method for the derivatization of secondary amino acids, proline, and hydroxyproline (2–4,7). However, its disadvantages are that it requires a high-vacuum system and it takes a long time to remove the byproducts produced in the process of derivatization and the excess reagent in order to avoid interfering peaks.

A need for simple, sensitive, stable, and more volatile precolumn derivatization reagent for analysis of amino acid with RP-HPLC and UV-detection still remains, although there have been developments of many reagents for precolumn derivatization, because these derivatives have some faults. The widely used precolumn derivatives, except PTC-amino acids, in RP-HPLC are *o*-phthalaldehydes (OPA) (9,10), dansyl (11,12), dabsyl (13,14), and 4-nitrophenylthiocarbamyl (NPTC) (7,15) derivatives. With the OPA derivatives, secondary amino acids, proline, and hydroxyproline, were not detected

because OPA does not react with secondary amines in the absence of oxidizing agents. Moreover, as OPA derivatives are unstable, complete automation of the precolumn reaction with accurate control of the reaction time is essential for acceptable reproducibility (*1*).

Dansyl derivatives are formed in the dark and are unstable toward prolonged reaction times, solvents, and exposure to light and interfering peaks arise because of the byproducts during the derivatization process (*12*). In the dansyl derivatives, it was reported that the sulphonamide bond in the derivative was very stable. The limitation of the dansyl derivatization method is that the presence of an excess amount of urea, salt, phosphate, or ammonium hydrogen carbonate will change the pH of the reaction buffer and interfere with derivatization (*13,14*).

Derivatization with 4-nitrophenylisothiocyanate (NPITC) forms the stable nitrophenylthiocarbamyl (NPTC) derivatives that are suitable for analysis by RP-HPLC and UV-detection at 254 nm or 340 nm (*15*). The disadvantage of NPITC is that the excess reagent cannot be easily removed under a high-vacuum system. Extraction with toluene can remove the excess reagent (*7*).

A more volatile reagent for precolumn derivatization compared to other reagents developed to recent year, butylisothiocyanate (BITC), an aliphatic compound, was successfully adopted for derivatization to butylthiocarbamyl (BTC) derivatives of 22 protein standard amino acids. The BTC-amino acids were successfully analyzed via C<sub>18</sub> RP-HPLC and UV-detector at 250 nm (*16*). The BITC reagent was also successfully adapted for the analysis of amino acids in foods (*17*).

The advantages of BITC were high volatility and the ability of separate derivatization on the cysteine and cystine, which had not been found with PTC-derivatives. The high volatility of this reagent substantially reduced the analysis time because the excess reagent and byproducts produced during the reaction could be easily removed. The BTC-derivatives of the secondary amino acids, proline, and hydroxyproline were also detected with high sensitivity. But asparagine and serine peaks overlapped completely and the stability of BTC-derivatives at room temperature was estimated to be only approx 8 h.

Benzylisothiocyanate (BZITC), the analog of phenylisothiocyanate (except NPITC), was successfully derivatized to benzylthiocarbamyl (BZTC) derivatives on all of the 22 protein amino acids and the derivatives were completely separated on a reversed-phase Nova-Pak C<sub>18</sub> column (*18*). The BZITC reagent was less volatile compared to BITC but the volatility was similar to PITC. The advantages of BZTC-derivatives compared to PTC-derivatives was superior resolution on reversed-phase column and superior reproducibility to PTC-derivatives using the same experimental condition.

In this chapter, more detail experimental methods, results, and discussions on the BTC and BZTC-derivatives developed at the most recent years will be described.

## 2. Materials

### 2.1. Equipment

1. A water aspirator for vacuum (*see Note 1*).
2. Spectra-Physics 8800 ternary solvent delivery system with solvent stabilization and degassing system.
3. Spectra 200 programmable wavelength UV detector.
4. Nova-Pak C<sub>18</sub> (300 × 3.9 id, 4 μm dimethyloctadecylsilyl-bonded amorphous silica, Waters).
5. Eppendorf CH-30 column heater.

### 2.2. Reagents

1. BITC, BZITC, and PITC are obtained from Aldrich (Milwaukee, WI) and were stored at 0–5°C (*see Note 2*).
2. Bovine serum albumin (BSA), standard amino acids, and norleucine are obtained from Sigma (St. Louis, MO). Standard amino acids were stored at room temperature and BSA was stored at 2–8°C.
3. HPLC-grade acetonitrile, methanol, and tetrahydrofuran from Merck (Darmstadt, Germany) were stored at room temperature.
4. All other reagents were of analytical grade.
5. Food samples, whole egg, and soybean purchased from a commercial market.

### 2.3. Solutions

1. Standard amino acid solution: A mixture solution of standard amino acids, except glutamine, cysteine, and cystine was prepared at a concentration of 2.5 μmol/mL of 0.01 M HCl. Standard solutions of glutamine and cysteine were prepared with water (*see Note 3*). Cystine was prepared at a concentration of 0.5 μmol/mL of 0.01 M HCl because of the solubility.
2. Preparation of coupling buffer solution: Coupling buffer (acetonitrile-methanol-triethylamine [10:5:2]) containing L-norleucine (2.5 μmol/mL) as an internal standard (*see Note 4*) was stored at 0–5°C and prepared again after 1 mo.

## 3. Method

### 3.1. Hydrolysis of BSA and Food Protein Samples (Figs. 1 and 2)

1. Place BSA (4 mg) and grinded soybean or homogenized whole egg food sample (0.2 g) in a 5-mL and 25-mL test tube, respectively, with an open-hole screw cap with a septum (**Fig. 3**).
2. Add 0.5 mL and 15 mL of 6 M HCl containing 0.1% phenol into the 5-mL and 25-mL test tubes containing the BSA and food samples, respectively.

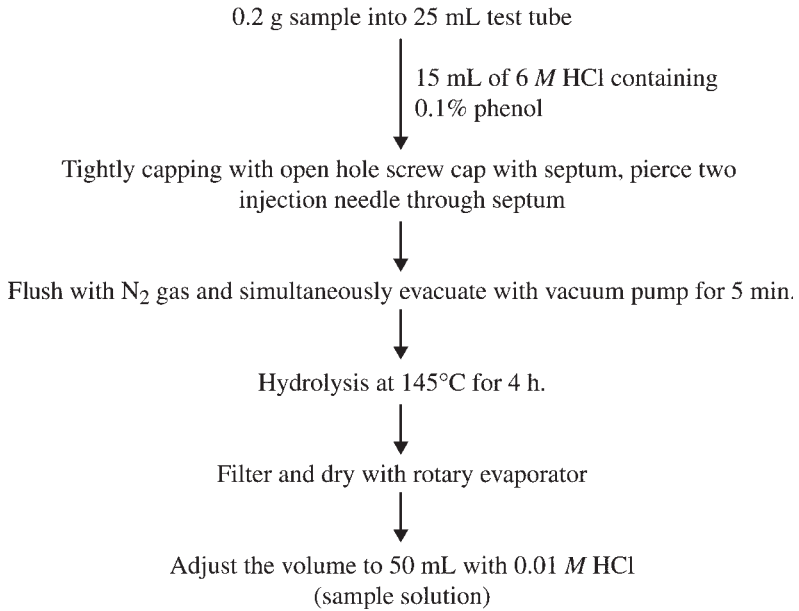


Fig. 1. Flow chart for hydrolysis of food samples.

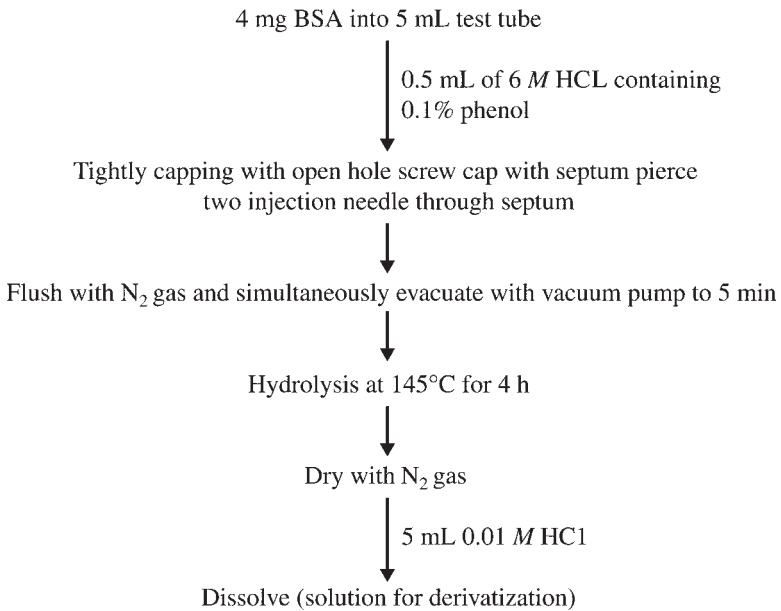


Fig. 2. Flow chart for hydrolysis of BSA.

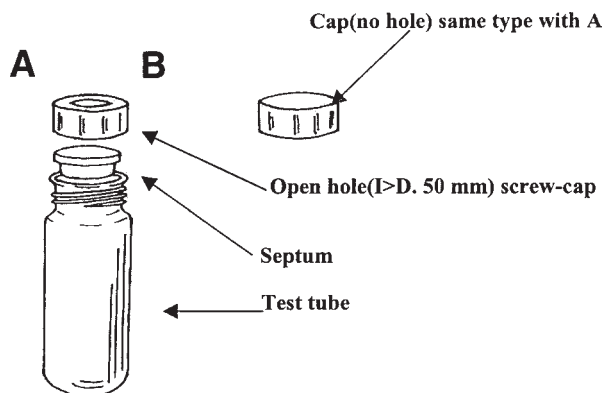


Fig. 3. Test tube for hydrolysis of food protein samples.

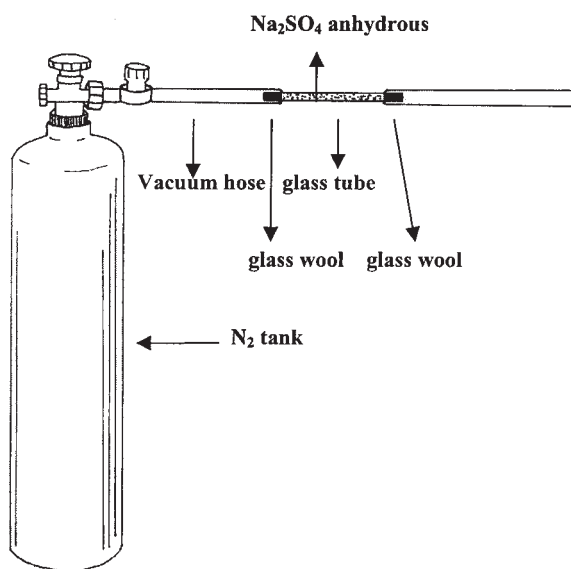


Fig. 4. The device for supply of the dried  $N_2$  gas.

3. After tightly capping, pierce the septum with two stainless steel injection needles.
4. Connect one needle, immersed in the sample solution, to a dried nitrogen supply (*see Note 5, Fig. 4*). Connect the other needle, not immersed, to the vacuum pump (*Fig. 5*).
5. Evacuate the test tubes with a vacuum pump for 5 min and simultaneously flush with nitrogen gas.

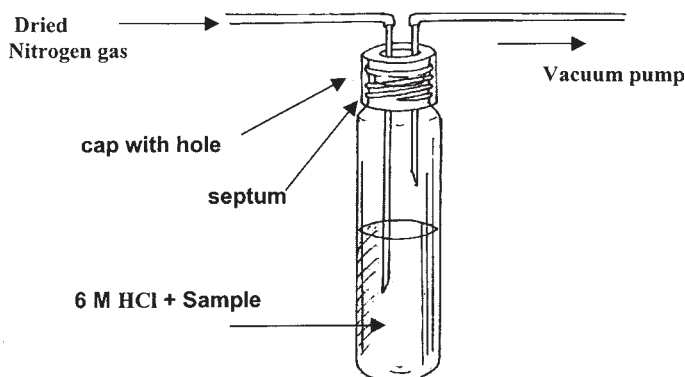


Fig. 5. The diagram of device for nitrogen saturation into the sample.

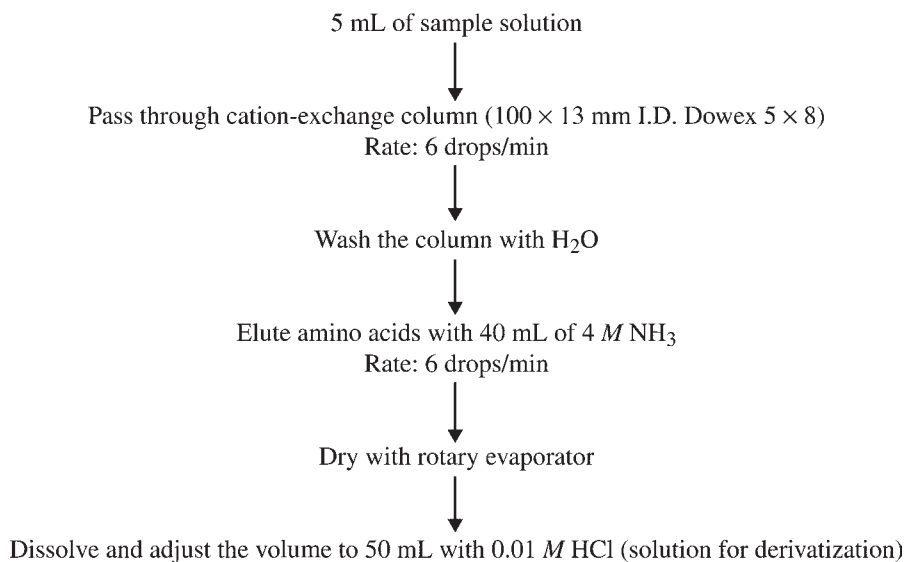


Fig. 6. Flow chart for clean up with cation-exchange column.

6. Remove the needle connected to the vacuum pump before removing the needle connected to the nitrogen.
7. Carefully remove the cap with the holes and change for a cap without holes (*see Note 6*), because small holes may be formed by the high pressure.
8. Carry out hydrolysis at 145°C for 4 h.
9. Dry the hydrolysate of BSA with nitrogen at 50°C. Redissolve with 5 mL of 0.01 M HCl (hydrolysate solution). This is now ready for derivatization.

10. Filter the hydrolysates of soybean meal and whole egg, then dry with a rotary evaporator. Redissolve and adjusted the volume to 50 mL with 0.01 M HCl (hydrolysate solution).
11. To clean up the hydrolysate solutions of soybean meal and whole egg use cation-exchange chromatography as follows (**Fig. 6.**).
12. Pass 5 mL of hydrolysate solution through a 100 × 13-mm id cation-exchange column (Dowex 5 × 8) at the rate of 6 drops/min, to retain the amino acids on the cation-exchange resin (*see Note 7*).
13. Wash the column several times with 20 mL of H<sub>2</sub>O (*see Note 8*).
14. Elute the retained amino acids on the cation-exchange column with 40 mL of 4 M ammonia solution with the rate of 6 drops/min.
15. Dry the eluted solutions in a rotary evaporator at 50°C. Redissolve in 0.01 M HCl (hydrolysate solution) and adjust the volumes to 50 mL. These samples are now ready for derivatization (*see Note 9*).

### 3.2. Derivatization (**Fig. 7**)

1. Place 20 µL of the mixture solution of standard amino acids, 50 µL of the standard solution of cystine, and sample hydrolysates (BSA; 100 µL, soybean; 500 µL, whole egg; 500 µL) into separate 2-mL conical vials with an open-hole screw cap and a septum (**Fig. 8**).
2. Dry the solutions completely with nitrogen gas at 50°C.
3. Add an appropriate amount of acetonitrile to each vial and dried again (*see Note 10*).
4. Redissolve the residues in 50 µL of coupling buffer.
5. Add of 3 mL of BITC, BZITC, and PITC to each of the dissolved solutions for BTC, BZTC, and PTC derivatives, respectively.
6. After tightly capping the vials with open-hole screw-caps, the derivatizations are carried out at 40°C for 30 min for BTC and PTC derivatives and at 50°C for 30 min for BZTC derivatives.
7. After derivatizations, use two stainless steel injection needles to pierce through the septum into the vials. Connect one needle with the nitrogen supply and the other with the vacuum pump (**Fig. 9**).
8. Infuse nitrogen into the vials and simultaneously evacuate with the vacuum pump to complete dryness at room temperature for approx 10 min for BTC derivative and for approx 40 min for BZTC and PTC derivatives.
9. Inject 100 µL of acetonitrile into the vials with a microinjection syringe and redry the contents for 5 min for BTC-derivative and for 30 min for BZTC and PTC derivatives (*see Note 11*).
10. Dissolve the residue of BTC-derivatives in 1 mL of 0.02 M ammonium acetate.
11. Dissolve the residue of BZTC and PTC-derivatives in 1 mL of 0.02 M NaH<sub>2</sub>PO<sub>4</sub> containing 5% methanol and 1.5% tetrahydrofuran (pH 6.8, adjusted with phosphoric acid).
12. Filter the dissolved solutions through a 0.25-µm membrane filter.
13. Inject 10-µL aliquots of the filtrates onto the respective HPLC system.

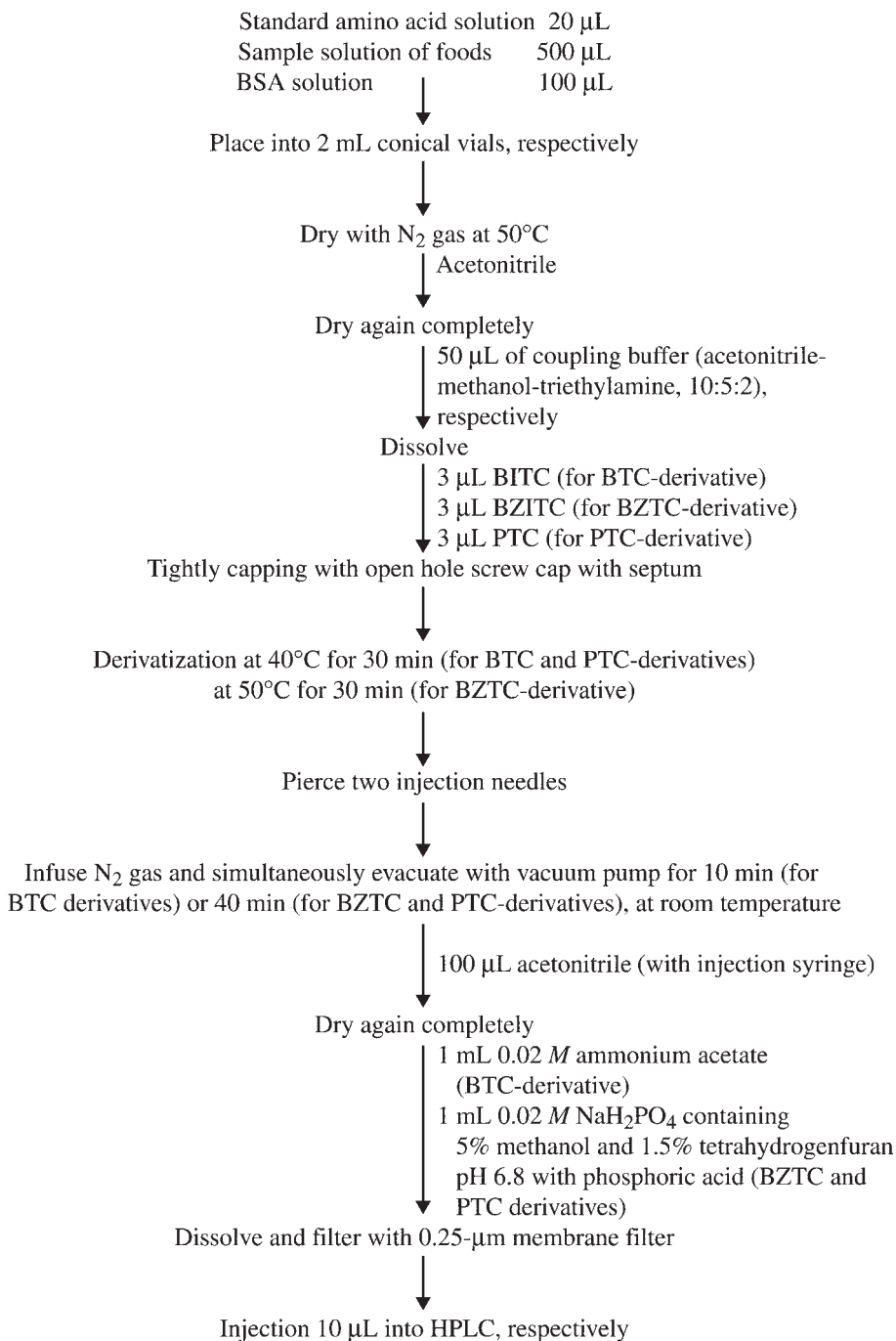


Fig. 7. Flow chart for derivatization.



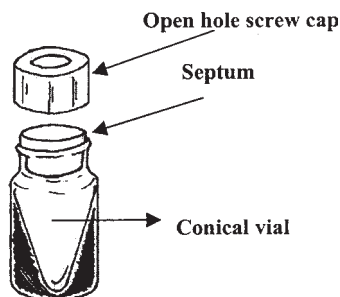


Fig. 8. 2-mL conical vial for derivatization.

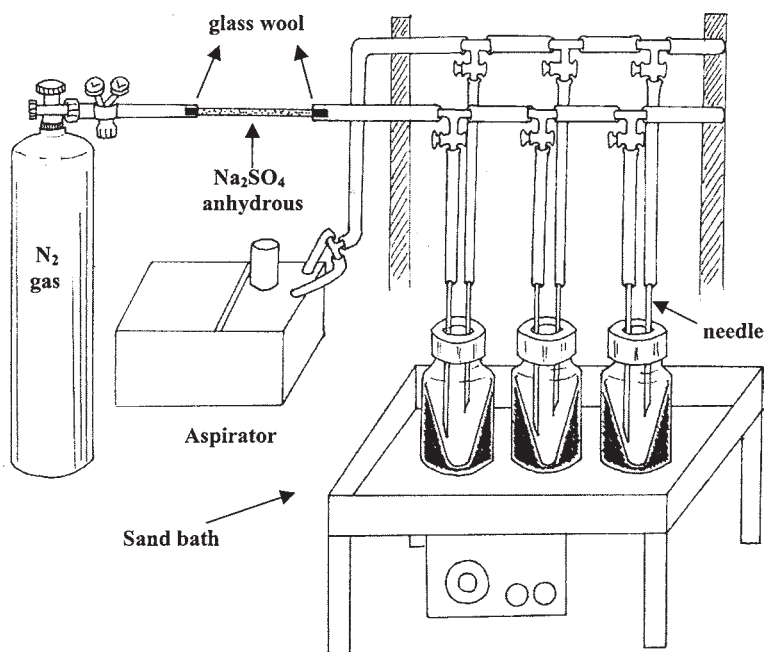


Fig. 9. The system drying several vials simultaneously.

### 3.3. Chromatography

HPLC conditions for the analysis were elucidated with **Table 1**.

### 3.4. Sensitivities of BTC, BZTC, and PTC-Derivatives

The sensitivities of BTC, BZTC, and PTC-derivatives are detected at 0.05 AUFS, which is the limit that gives a stable baseline with the smallest quantity.

It is possible to determine a linear relationship with the quantitative analysis (see **Notes 21** and **22**).

**Table 1**  
**HPLC Conditions for the Analysis of BTC, BZTC,**  
**and PTC-Amino Acids Derivatives**

|                          |                                                                                                                                                                                                                                                                                |      |    |    |            |
|--------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------|----|----|------------|
| HPLC system              | Spectra-Physics 8800 ternary solvent delivery system                                                                                                                                                                                                                           |      |    |    |            |
| Detector                 | Solvent stabilization and degassing system with a blanket of helium                                                                                                                                                                                                            |      |    |    |            |
| Wavelength for detection | Spectra 200 programmable wavelength UV detector                                                                                                                                                                                                                                |      |    |    |            |
| Column                   | 240 nm for BTC-derivatives ( <i>see Note 12</i> )                                                                                                                                                                                                                              |      |    |    |            |
|                          | 246 nm for BZTC-derivatives ( <i>see Note 13</i> )                                                                                                                                                                                                                             |      |    |    |            |
|                          | 254 nm for PTC-derivatives ( <i>see Note 13</i> )                                                                                                                                                                                                                              |      |    |    |            |
| Column                   | Nova-Pak C <sub>18</sub> (300 × 3.9 id, 4 μm dimethyloctadecylsilyl-bonded amorphous silica, Waters). For all derivatives column-temperature; 40°C with Eppendorf CH-30 column heater                                                                                          |      |    |    |            |
| Solvent system           | For BTC-derivatives ( <i>see Notes 14–17</i> ):                                                                                                                                                                                                                                |      |    |    |            |
|                          | A solution; 0.05 M ammonium acetate (pH 6.7 adjusted with phosphoric acid)                                                                                                                                                                                                     |      |    |    |            |
|                          | B solution; 0.02 M sodium phosphate dibasic solution containing 5% methanol and 1.5% tetrahydrofuran-acetonitrile (50:50)                                                                                                                                                      |      |    |    |            |
|                          | C solution; acetonitrile-water (70:30)                                                                                                                                                                                                                                         |      |    |    |            |
|                          | Solvent gradient                                                                                                                                                                                                                                                               |      |    |    |            |
|                          |                                                                                                                                                                                                                                                                                | A    | B  | C  | Flow rate  |
|                          | 0.0 min                                                                                                                                                                                                                                                                        | 100% | 0% | 0% | 1 mL/min   |
|                          | 5.0 min                                                                                                                                                                                                                                                                        | 85   | 15 | 0  | "          |
|                          | 14.0 min                                                                                                                                                                                                                                                                       | 70   | 20 | 10 | "          |
|                          | 20.0 min                                                                                                                                                                                                                                                                       | 60   | 20 | 20 | "          |
|                          | 25.0 min                                                                                                                                                                                                                                                                       | 30   | 20 | 50 | "          |
|                          | 30.0 min                                                                                                                                                                                                                                                                       | 10   | 20 | 70 | "          |
|                          | For BZTC-derivatives ( <i>see Notes 14–17</i> ) and PTC-derivatives ( <i>see Note 14–17</i> )                                                                                                                                                                                  |      |    |    |            |
|                          | A solvent; 0.02 M NaH <sub>2</sub> PO <sub>4</sub> containing 5% methanol and 1.5% tetrahydrofuran (pH 6.8 adjusted with phosphoric acid)                                                                                                                                      |      |    |    |            |
|                          | B solvent; A solvent-acetonitrile (50:50)                                                                                                                                                                                                                                      |      |    |    |            |
|                          | C solvent; acetonitrile-water (70:30)                                                                                                                                                                                                                                          |      |    |    |            |
|                          | Solvent gradient                                                                                                                                                                                                                                                               |      |    |    |            |
|                          |                                                                                                                                                                                                                                                                                | A    | B  | C  | Flow rate  |
|                          | 0.0 min                                                                                                                                                                                                                                                                        | 100% | 0% | 0% | 1.2 mL/min |
|                          | 15.0 min                                                                                                                                                                                                                                                                       | 76   | 20 | 4  | "          |
|                          | 20.0 min                                                                                                                                                                                                                                                                       | 70   | 20 | 10 | "          |
|                          | 30.0 min                                                                                                                                                                                                                                                                       | 50   | 30 | 20 | "          |
|                          | 40.0 min                                                                                                                                                                                                                                                                       | 30   | 35 | 35 | "          |
|                          | After this gradient program, a washing step for 20 min with solvent C substantially protected the column damage. Above two types of solvent gradients were adapted to the samples amino acid derivatives as well as standard amino acid derivatives ( <i>see Note 18–20</i> ). |      |    |    |            |

### 3.5. Stabilities of the BTC-Derivatives and BZTC-Derivatives

The variations in the peak-area responses of BTC and BZTC-derivatives with storage time at room temperature can be determined for the stabilities of the BTC and BZTC-derivatives (*see* **Notes 23** and **26**).

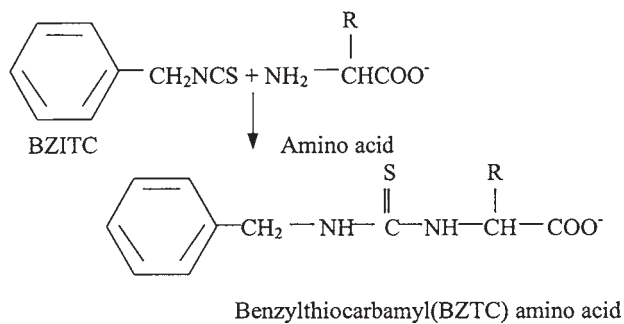
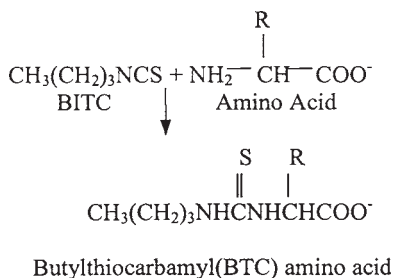
### 3.6. Statistical Analysis

1. To determine the reproducibilities, all experiments should be repeated more than three times.
2. Relative standard deviations (RSDs) on the relative molar response (RMR) can be calculated for comparison of precisions on the derivatives. Linearity of calibration graphs on the appropriate ranges are also detected by the determination of statistical significance of correlation coefficient ( $\gamma$ ) of calibration graphs (*see* **Notes 27** and **33**).
3. The reproducibility and accuracy on the BSA and food samples can be compared to ion-exchange chromatography and the data of the other literatures (*see* **Notes 34** and **38**).

## 4. Notes

1. If you use a vacuum pump that is not using water, you must install a device to absorb the evaporating excess reagents and byproducts produced during derivatization because the vacuum pump is rendered useless by these evaporating materials.
2. These reagents seem to be stable for several years unopened, but when the caps were opened several times, the reagents must be used in about 6 mo even if they are stored at  $-20^{\circ}\text{C}$ . All of these reagents are very harmful and toxic.
3. Standard solutions of glutamine and cysteine were prepared with water because of the conversion to pyroglutamic acid and cystine, respectively, on prolonged storage of these amino acids in HCl solution (**19**).
4. For BZTC-derivatives of sample internal standard (L-norleucine) should not be used.
5. For supply of the dried nitrogen, connect the tube for absorption of any moisture as shown in **Fig. 2**. This  $\text{Na}_2\text{SO}_4$  anhydrous tube must be periodically changed every 1 or 2 mo.
6. When the cap with a hole is changed to the cap without a hole, you have to be carefully that the septum is not removed.
7. Be careful that the sample, washing, and eluting solutions do not drop under the level of cation-exchange resin in the column.
8. The rate of drops during the washing is not important.
9. For the analysis with ion-exchange chromatography and ninhydrin derivatization method, dry the clean-up sample solutions in a rotary evaporator. Redissolve in 0.2 M sodium citrate buffer (pH 2.2) and inject into the automation amino acid analyzer (LKB 4150 Alpha, Ultrapac-11 cation-exchange column).
10. By the synergy effect of acetonitrile evaporating, the residues of water are completely dried.

11. The chemical reactions of BTC and BZTC-derivatives are as follows:



12. UV-spectra of the BTC-amino acid mixture and BITC, the coupling reagent, are shown in **Fig. 10**. The  $\lambda_{\text{max}}$  of BTC-amino acids was about 234 nm, but the most efficient wavelength was 250 nm, which avoided the absorption spectra of the impurities and the electrolyte, ammonium acetate in the solvent (**16**).
13. The UV spectra of the BZTC-amino acids mixture and the PTC-amino acid mixture dissolved in 0.02 M NaH<sub>2</sub>PO<sub>4</sub> are shown in **Fig. 11**. The wavelengths giving strong absorbance were 220 nm and 238 nm in the BZTC derivative and 215 and 270 nm in the PTC derivative, but the most efficient wavelengths were 246 nm in the BZTC derivative and 254 nm in the PTC derivative, which avoided interference by the absorption spectra of the impurities and electrolyte and showed a stable baseline (**18**).
14. The standard amino acid chromatograms of BTC and BZTC-derivatives compared to PTC-derivatives separated on the Nova-Pak C<sub>18</sub> column are shown in **Fig. 12**. All of the 22 standard amino acids were derivatized with BTC, BZTC, and PTC-derivatives and resolved on C<sub>18</sub> reversed column.
15. In the BTC-derivatives, asparagine and serine completely overlapped but BTC-cysteine and cystine were individually eluted even though the cystine peak was resolved with a tailing peak. In the PTC-derivatives, cystine and cysteine peaks appeared at the same position, from which it could be assumed that cysteine might be completely converted into cystine during derivatization because of the fact that cysteine can be oxidized (**20**). Other articles have also reported that PTC-cysteine and cystine were eluted at the same position (**2,4,7**).

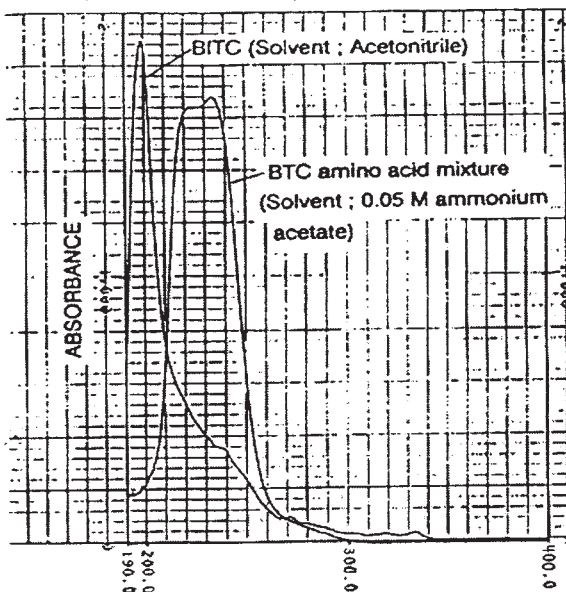


Fig. 10. UV spectra (x-axis in nm) of BTC-amino acid mixture and the coupling reagent BTC. Absorbance of solvents at 250 nm = 0.

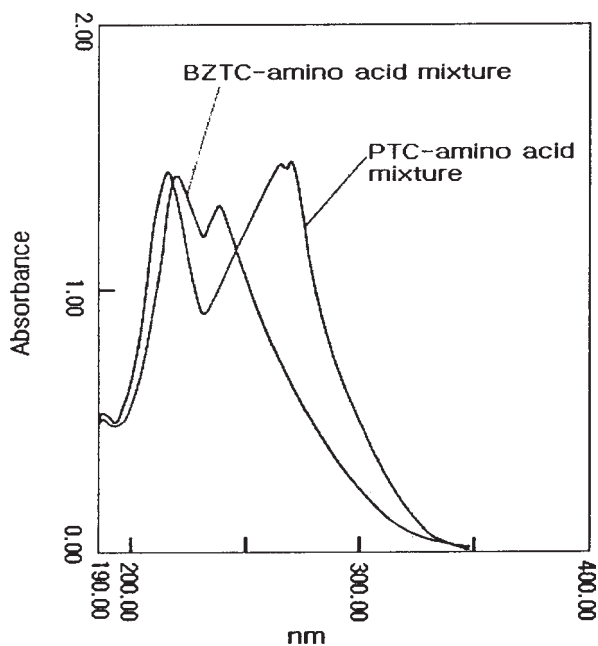


Fig. 11. UV spectra of the BZTC-amino acid mixture and the PTC-amino acid mixture. Solvent, 0.05 M  $\text{NaH}_2\text{PO}_4$ . Absorbance of solvent at 254 nm = 0.

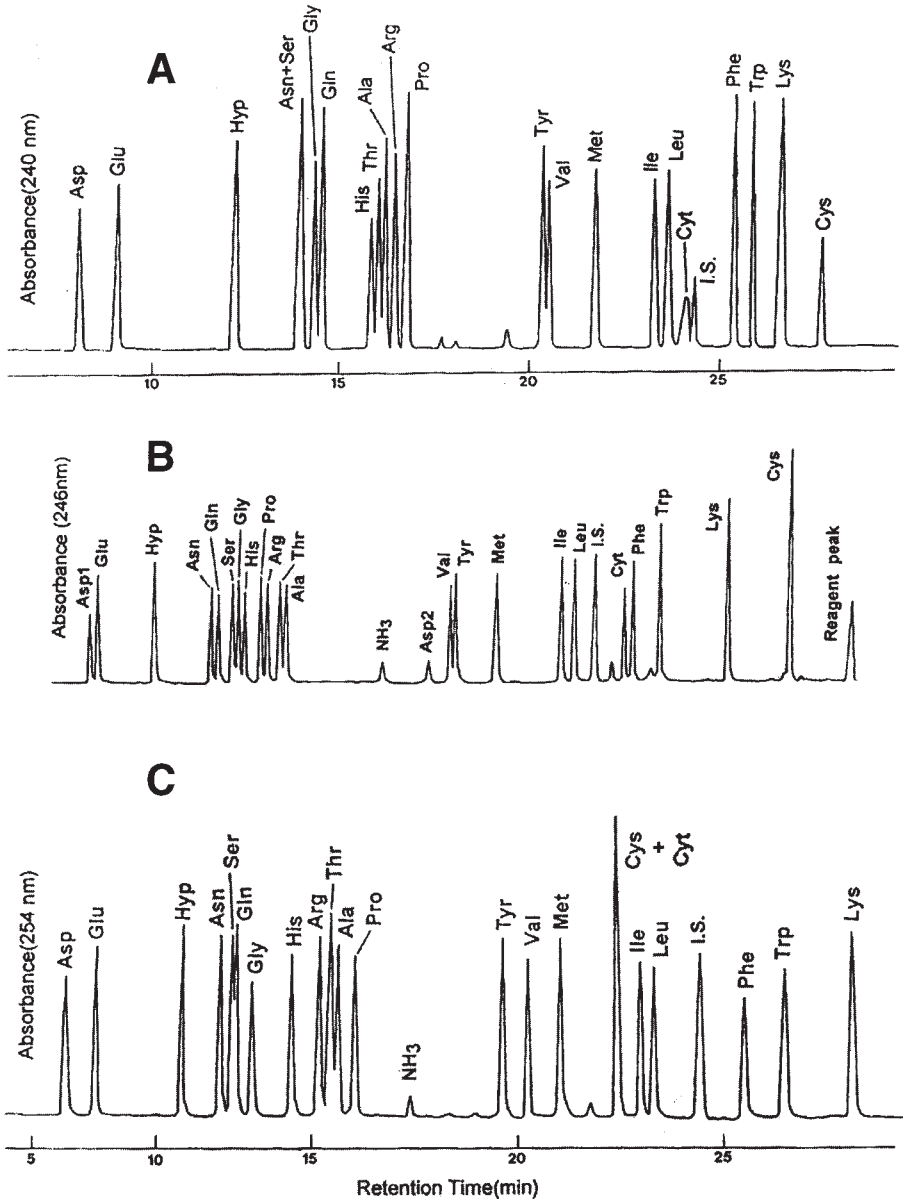


Fig. 12. Chromatogram of standard protein amino acid derivatives resolved on a Nova-Pak (30 cm  $\times$  3.9 mm)  $C_{18}$  column. I.S. = norleucine, injected amount 0.625 nmol. (A) BTC-amino acid; (B) BZTC-amino acid; (C) PTC-amino acid. Cyt = cystine; Cys = cysteine.

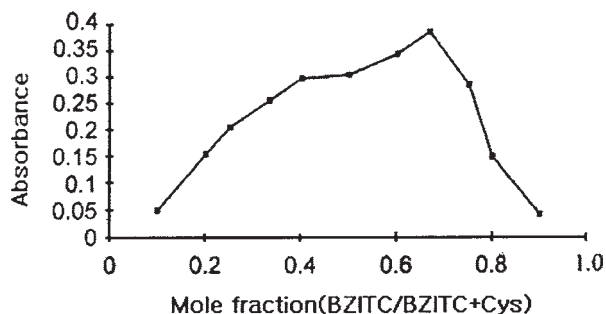


Fig. 13. Absorbance plot of BZTC-cysteine derivative at 238 nm showing maximum absorption of the mole fraction of BZITC in the derivatizing solution for the reaction between benzylisothiocyanate and cysteine.

16. Unlike BTC and PTC-derivatives, in the BZTC-derivatives all of 22 standard amino acids were nearly completely separated. Also, in the BZTC-derivatives as in the BTC-derivatives, the cysteine and cystine peaks were markedly separated.
17. Unlike the PTC-derivatives, in the BZTC and BTC-derivatives, the -SH moiety of cysteine would be converted to the thiocyanate derivatives by BZITC and BITC. To find out whether 1 mol of cysteine reacts with 2 mol of BZITC, absorbances of the BZTC-derivatives on the mole fraction of BZITC in the solution for derivatization between BZITC and cysteine were determined (**Fig. 13**). The determined wavelength was 238 nm, which gave maximum absorption for the BZTC-derivatives. Maximum absorbance appeared at 0.667 of the mole fraction, which meant that the mole ratio of the reaction between cysteine and BZITC was 1:2. So we could draw the conclusion that the -SH group, as well as the amino group of cysteine was derivatized. Intelligence due to the fact that -SH moiety of cysteine was readily converted to the thiocyanate derivative with 2-nitro-5-thiocyanobenzoic acid (**21**) could support this conclusion. BZTC and BTC-cysteine might be eluted last because of the long less-polar side chain by the derivatization of -SH group.
18. Chromatograms of BTC, BZTC, and PTC-amino acid derivatives of soybean meal hydrolysate are shown in **Fig. 14**. In the BTC and PTC-derivatives we could detect the same kinds of amino acids of 16, but in the BZTC-derivatives cystine and cysteine in addition to above 16 amino acids were detected. We think that the sensitivity of BZTC-derivatives are more superior to the BTC and PTC-derivatives.
19. Chromatograms of BTC and PTC-amino acid derivatives on the whole egg hydrolysate are shown in **Fig. 15**. Because unfortunately, we did not detect on the BZTC-amino acid derivatives of the whole egg hydrolysate, it is really regrettable that we could not compare to BTC and PTC-derivatives. The same kinds of 17 amino acids in the BTC and PTC-derivatives were detected. As is shown in the chromatograms of food samples (**Figs. 14 and 15**), assuming that the some kinds

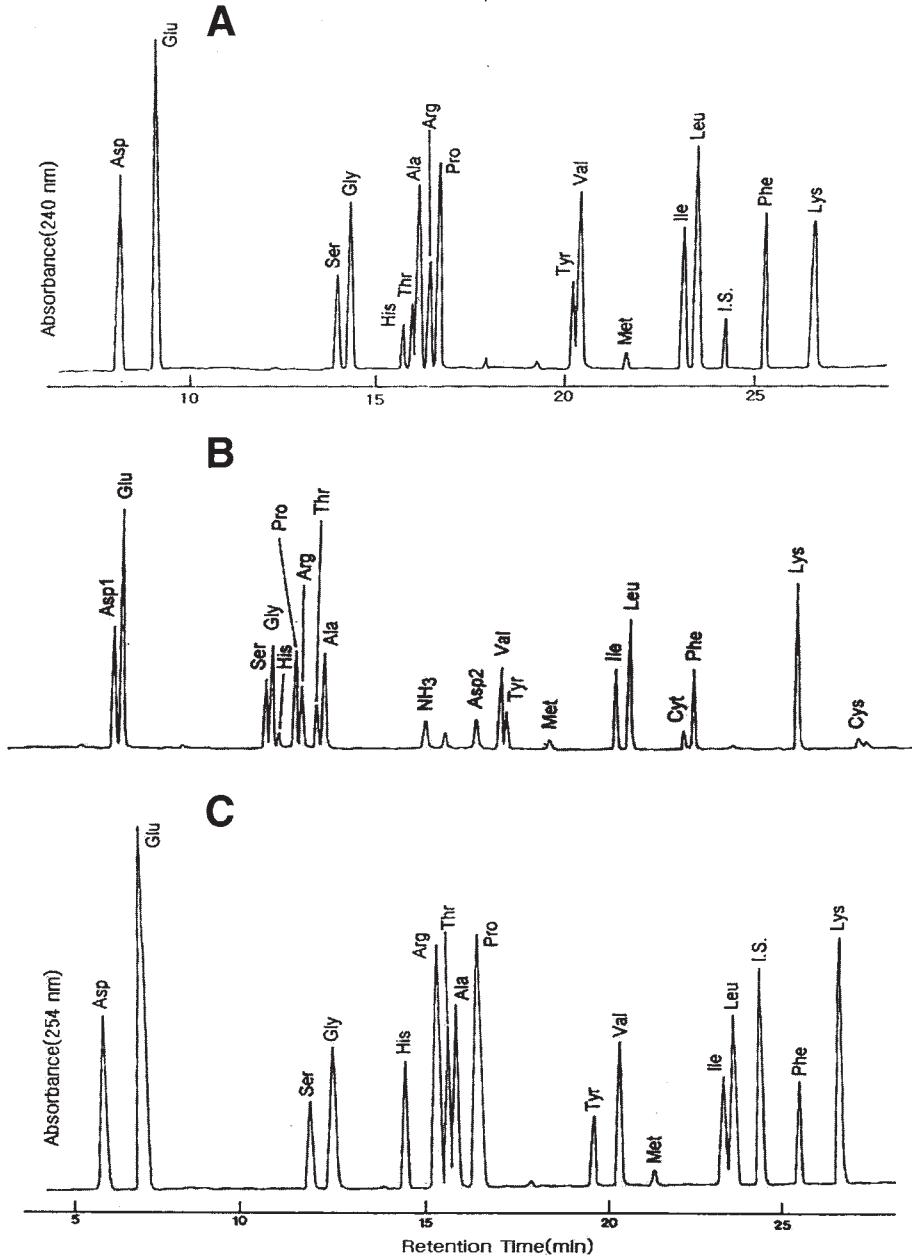


Fig. 14. Chromatogram of amino acids in soybean hydrolysate. (A) BTC derivatives; (B) BZTC derivatives; (C) PTC derivatives.



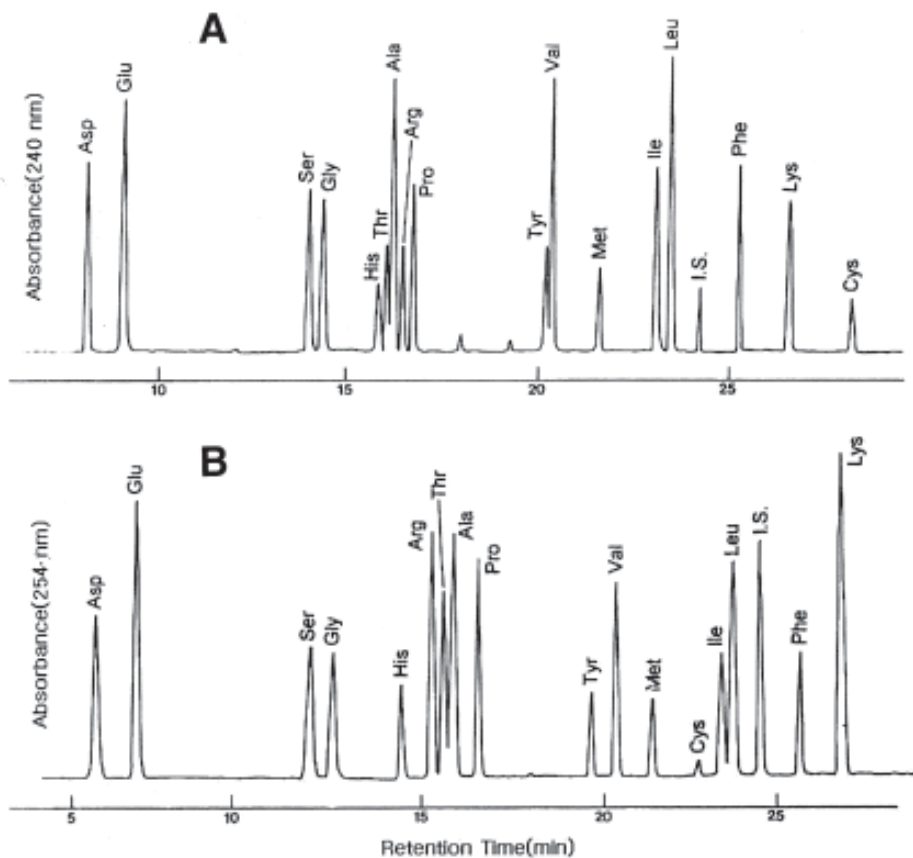


Fig. 15. Chromatogram of amino acids in whole egg hydrolysate. (A) BTC derivatives; (B) PTC derivatives. Cys = cysteine.

of contaminants were contained there were few ghost peaks. This phenomenon indicates that BITC, BZITC, and PITC are very good reagents having superior selective reactivity with amino acids.

20. Chromatograms at BTC, BZTC, and PTC-derivatives of BSA hydrolysate are shown in **Fig. 16**. In the soybean meal, on the other hand, the BTC and PTC-derivatives detected 17 amino acids, in the BZTC-derivatives 18 amino acids were detected with the additional detection of cystine.
21. Sensitivities of BTC, BZTC, and PTC-derivatives were about 3.9 pmol at 0.05 AUFS (**Fig. 17**). At levels lower than 3.9 pmol, several amino acids in the all of the three derivatives were not detected and not showed the linearity in the calibration graphs for the quantitative analysis.
22. There is a report that the sensitivity of the PTC-derivatives was 1 pmol in the detector at 0.005 AUFS (signal-to-noise ratio 5:1) (2). But in our experiment, at

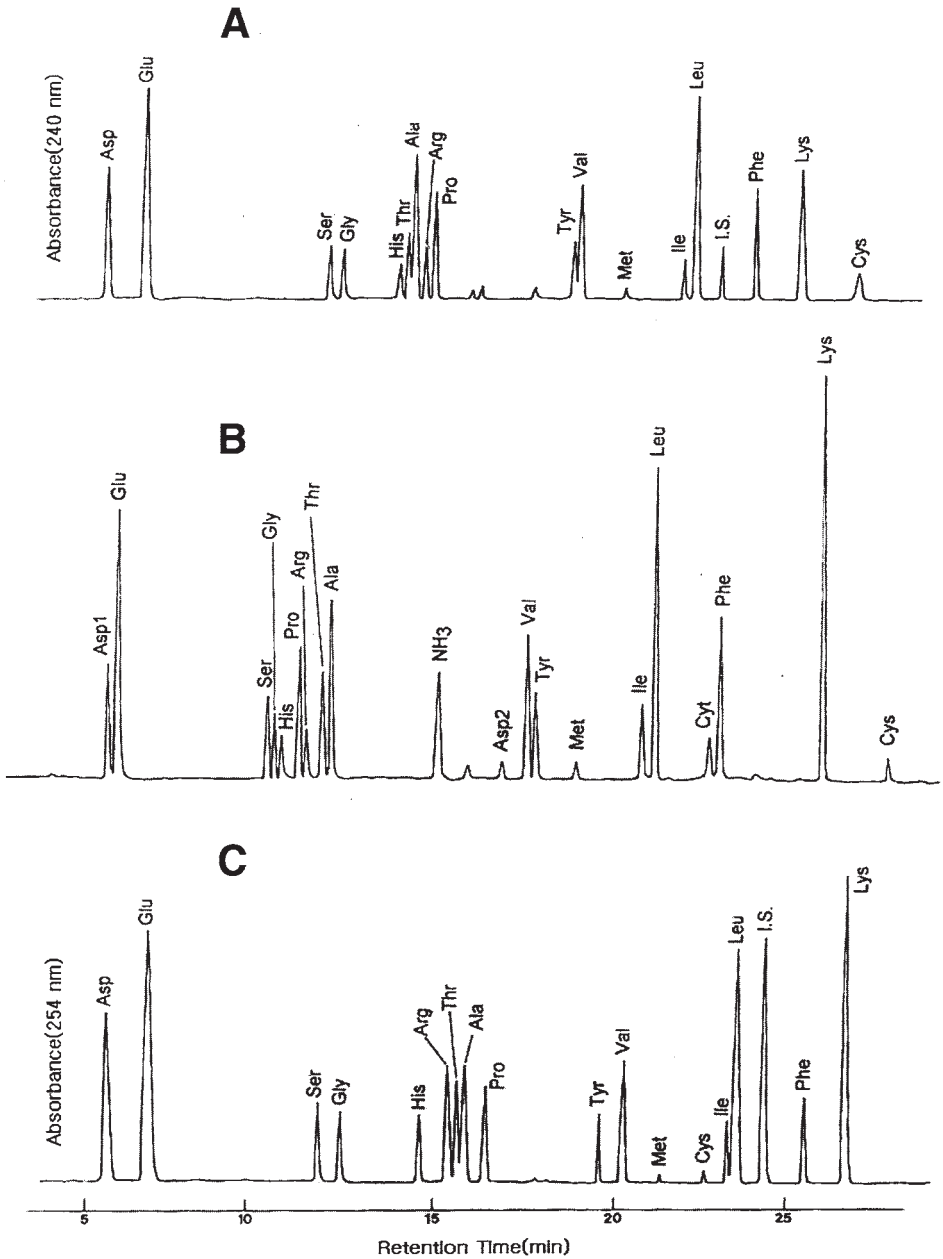


Fig. 16. Chromatogram of amino acids in bovine serum albumin hydrolysate. (A) BTC derivatives; (B) BZTC derivatives; (C) PTC derivatives. Cys = cysteine.

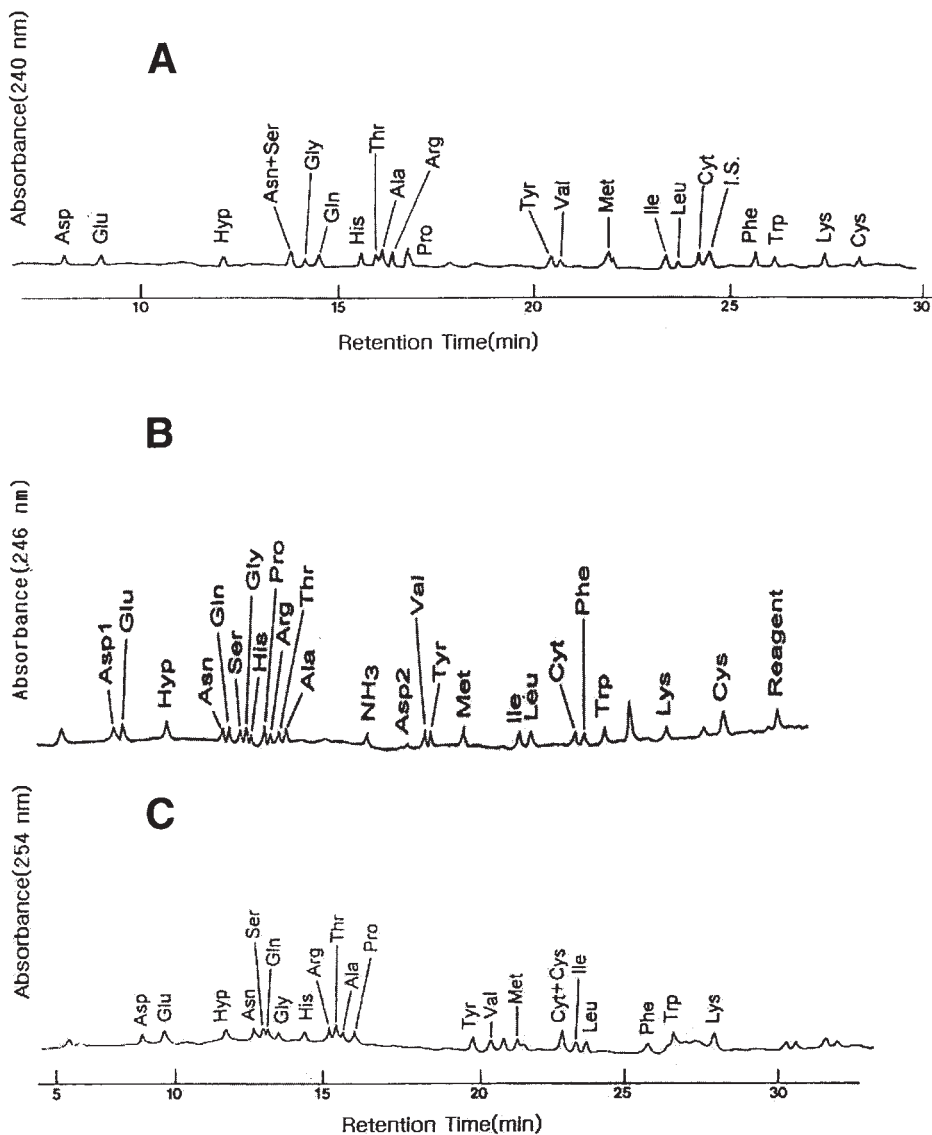


Fig. 17. Chromatogram of standard protein amino acid derivatives showing the sensitivity. Injected amount 3.9 pmol. Range 0.05 AUFS. (A) BTC-amino acid; (B) BZTC-amino acid; (C) PTC-amino acid. Cyt = cystine; Cys = cysteine.

this level, because the contamination of amino acids by the reagents, instrument, environment, and solvents was serious and the baseline was very unstable, the adaptation of the practical analysis was impossible.

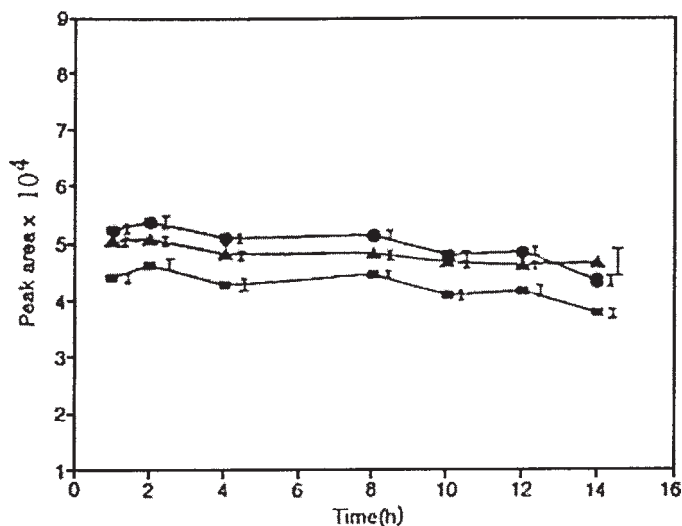


Fig. 18. Stability of BTC-amino acids at room temperature. ■ = Glu; ● = Ile; ▲ = Leu.

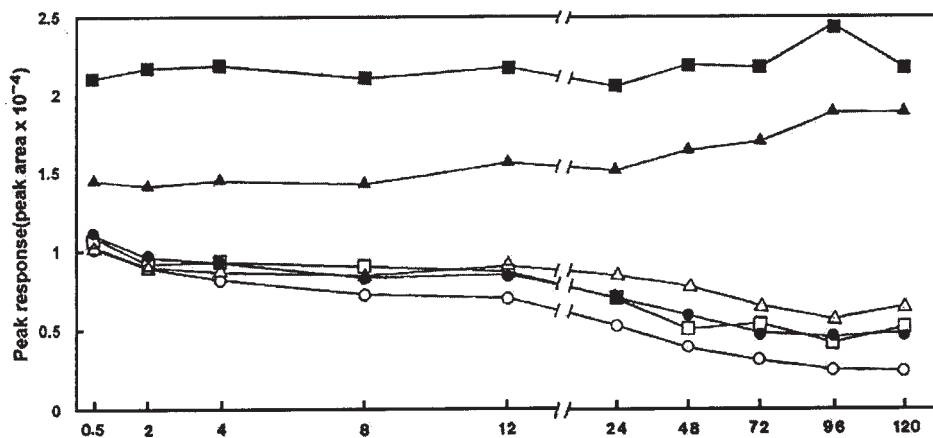


Fig. 19. Stability of BZTC derivatives of hydroxyproline (■), proline (▲), serine (●), cystine (□), alanine (△), and threonine (○). Each point is the mean value of three measurements.

23. The stability of BTC-derivatives and BZTC-derivatives is shown in **Figs. 18** and **19**, respectively.
24. In the BTC-derivatives, up to 2 h of storage, the peak area responses increased for most of the derivatives, which indicated the derivatization reaction was continuing at room temperature. After 8 h of the storage, the derivative that showed

**Table 2**  
**RMR of BTC, BZTC, and PTC Derivatives of Standard Amino Acids**

| Amino acid | BTC derivatives   |      | BZTC derivatives |      | PTC derivatives   |      |
|------------|-------------------|------|------------------|------|-------------------|------|
|            | RMR <sup>a</sup>  | RSD  | RMR <sup>b</sup> | RSD  | RMR <sup>b</sup>  | RSD  |
| Asp        | 0.50              | 1.40 | 1.89             | 3.82 | 0.89              | 2.48 |
| Glu        | 0.48              | 1.25 | 1.21             | 0.83 | 0.84              | 1.83 |
| Hyp        | 0.07              | 1.64 | 1.61             | 4.50 | 0.93              | 1.62 |
| Asn        | 0.66 <sup>c</sup> | 0.45 | 1.12             | 4.48 | 0.91              | 2.37 |
| Gln        | 0.71              | 1.27 | 1.07             | 2.80 | 1.03              | 5.89 |
| Ser        | —                 | —    | 1.05             | 1.46 | 0.76              | 2.02 |
| Gly        | 0.52              | 0.96 | 0.96             | 2.63 | 0.65              | 1.68 |
| His        | 0.62              | 1.13 | 0.84             | 2.49 | 0.83              | 0.87 |
| Pro        | 0.71              | 1.55 | 1.17             | 3.42 | 0.85              | 7.37 |
| Arg        | 0.44              | 0.68 | 1.00             | 1.15 | 0.91              | 3.02 |
| Thr        | 0.29              | 1.72 | 0.92             | 1.09 | 1.01              | 16.6 |
| Ala        | 0.75              | 2.53 | 0.88             | 1.73 | 0.90              | 6.64 |
| Val        | 0.88              | 3.86 | 0.95             | 1.08 | 0.83              | 4.68 |
| Tyr        | 0.83              | 1.33 | 1.39             | 2.20 | 0.92              | 2.64 |
| Ile        | 0.84              | 0.03 | 1.05             | 0.95 | 0.85              | 3.76 |
| Leu        | 0.86              | 2.21 | 0.84             | 1.38 | 0.84              | 6.86 |
| Cyt        | 0.49              | 4.29 | 0.79             | 3.64 | —                 | 10.5 |
| Phe        | 1.01              | 2.57 | 1.52             | 1.32 | 0.76              | 2.46 |
| Trp        | 0.93              | 3.87 | 2.04             | 0.98 | 0.92              | 1.64 |
| Lys        | 1.10              | 3.27 | 1.27             | 0.91 | 1.24              | 10.8 |
| Cys        | 0.63              | 5.24 | 3.63             | 7.22 | 1.62 <sup>d</sup> | —    |

RMR, relative molar response

RSD, relative standard deviation (n = 3)

<sup>a</sup>Values are relative to methionine.

<sup>b</sup>Values are relative to norleucine.

<sup>c</sup>Asn+Ser

<sup>d</sup>Cyt+Cys. Cyt = Cystine, Cys = cysteine.

the greatest decrease was cysteine (17.3%), the losses of the other derivatives after 8 h were in the range 0–6.9%. The derivatives that decreased to less than 5% after 14 h were glutamic acid, asparagine+serine, glutamine, threonine, tyrosine, proline, lysine, and tryptophan (16).

25. In the BZTC-derivatives, the peak area responses were decreased to less than 5% up to 120 h storage, but threonine, alanine, cystine, and serine were decreased to 18.9, 13.7, 15.1, and 15.6% after 4 h, respectively. Regarding the stability, we know that the optimum pH of the solvent dissolving the derivatives was very important. For the PTC-derivatives, the optimum pH of the solvent was 7.5. The loss of PTC-derivatives at this pH was in the range 0–10% after 10 h storage at room temperature (7).

**Table 3**  
**Amino Acid Compositions of Soybean Determined by the BTC, BZTC, and PTC-Derivatives Compared to the Ion-Exchange Chromatography (g/100 g dry matter)**

| Amino acid | BTC derivatives <sup>a</sup> | BZTC derivatives <sup>b</sup> | PTC derivatives <sup>a</sup> | PTC derivatives <sup>b</sup> | Ion-exchange chromatography <sup>a</sup> |
|------------|------------------------------|-------------------------------|------------------------------|------------------------------|------------------------------------------|
| Asp        | 4.17 ± 0.24                  | 4.45 ± 0.25                   | 4.64 ± 0.28                  | 4.72 ± 0.63                  | 4.25 ± 0.32                              |
| Glu        | 6.96 ± 0.42                  | 7.41 ± 0.14                   | 7.25 ± 0.34                  | 7.76 ± 0.84                  | 7.19 ± 0.12                              |
| Ser        | 1.61 ± 0.12                  | 2.00 ± 0.16                   | 1.59 ± 0.02                  | 2.03 ± 0.43                  | 1.65 ± 0.01                              |
| Gly        | 1.59 ± 0.07                  | 1.56 ± 0.03                   | 1.77 ± 0.10                  | 2.09 ± 0.13                  | 1.62 ± 0.05                              |
| His        | 1.50 ± 0.06                  | 1.06 ± 0.08                   | 1.52 ± 0.04                  | 1.15 ± 0.04                  | 1.32 ± 0.05                              |
| Arg        | 3.38 ± 0.17                  | 3.61 ± 0.13                   | 2.28 ± 0.08                  | 2.65 ± 0.36                  | 3.50 ± 0.09                              |
| Thr        | 1.19 ± 0.04                  | 1.78 ± 0.11                   | 1.79 ± 0.17                  | 1.79 ± 0.05                  | 1.24 ± 0.10                              |
| Ala        | 1.72 ± 0.10                  | 2.00 ± 0.12                   | 1.89 ± 0.35                  | 1.89 ± 0.35                  | 1.80 ± 0.12                              |
| Pro        | 2.15 ± 0.12                  | 1.96 ± 0.20                   | 2.06 ± 0.08                  | 2.38 ± 0.18                  | 2.20 ± 0.06                              |
| Tyr        | 1.20 ± 0.04                  | 1.20 ± 0.05                   | 1.29 ± 0.08                  | 1.26 ± 0.11                  | 1.15 ± 0.05                              |
| Val        | 2.88 ± 0.24                  | 1.97 ± 0.10                   | 2.47 ± 0.10                  | 2.12 ± 0.10                  | 3.01 ± 0.14                              |
| Met        | 0.42 ± 0.08                  | 0.73 ± 0.01                   | 0.54 ± 0.06                  | 0.50 ± 0.25                  | 0.40 ± 0.03                              |
| Ile        | 2.46 ± 0.14                  | 1.70 ± 0.05                   | 2.21 ± 0.06                  | 1.83 ± 0.16                  | 2.25 ± 0.18                              |
| Leu        | 3.38 ± 0.18                  | 2.95 ± 0.18                   | 3.03 ± 0.31                  | 3.10 ± 0.24                  | 3.57 ± 0.21                              |
| Cyt        | —                            | 0.49 ± 0.12                   | —                            | —                            | —                                        |
| Phe        | 2.27 ± 0.16                  | 2.06 ± 0.15                   | 2.10 ± 0.10                  | 2.63 ± 0.18                  | 2.19 ± 0.14                              |
| Lys        | 3.03 ± 0.13                  | 2.72 ± 0.10                   | 3.15 ± 0.16                  | 3.73 ± 0.25                  | 2.86 ± 0.19                              |
| Cys        | —                            | 0.17 ± 0.04                   | —                            | 0.11 ± 0.01 <sup>c</sup>     | —                                        |

All values are mean ± SD.

<sup>a</sup>*n* = 4. Data from **ref. 17**.

<sup>b</sup>*n* = 5. Data from **ref. 18**.

<sup>c</sup>Cys+Cyt. Cyt = cystine, Cys = cysteine.

26. We assumed that the stability of the BZTC-derivatives was superior to those of the PTC-derivatives and BTC-derivatives, except for threonine, alanine, cystine, and serine.
27. The relative molar responses of BTC, BZTC, and PTC-derivatives were shown in **Table 2**.
28. In the BTC-derivatives, RMR values determined with relative to methionine. RSDs on the RMR were less than 5% except cysteine (5.24%) in the BTC-derivatives. In the BZTC and PTC-derivatives, RMR values were determined with relative to norleucine, the internal standard.
29. In the BZTC-derivatives, the R.S.D values of all derivatives except cysteine (7.22%) were less than 5%. However, for the PTC-derivatives of glutamine, proline, alanine, and leucine the RSDs exceeded 5% and for threonine, cystine+cysteine, and lysine the RSDs exceeded 10%. In the PTC derivatization,

**Table 4**  
**Amino Acid Compositions of Whole Egg Determined by the BTC and PTC-Derivatives Compared to the Ion-Exchange Chromatography (g/100 g dry matter)**

| Amino acid | BTC derivatives | PTC derivatives | Ion-exchange chromatography |
|------------|-----------------|-----------------|-----------------------------|
| Asp        | 4.20 ± 0.09     | 4.74 ± 0.07     | 4.23 ± 0.05                 |
| Glu        | 5.48 ± 0.05     | 5.85 ± 0.59     | 5.76 ± 0.07                 |
| Ser        | 3.50 ± 0.09     | 2.61 ± 0.42     | 3.54 ± 0.10                 |
| Gly        | 1.56 ± 0.09     | 1.54 ± 0.31     | 1.53 ± 0.02                 |
| His        | 1.90 ± 0.14     | 1.91 ± 0.15     | 1.79 ± 0.06                 |
| Thr        | 1.75 ± 0.13     | 2.15 ± 0.11     | 1.82 ± 0.15                 |
| Ala        | 2.71 ± 0.16     | 3.29 ± 0.19     | 2.70 ± 0.11                 |
| Arg        | 3.55 ± 0.10     | 3.43 ± 0.48     | 3.61 ± 0.16                 |
| Pro        | 1.95 ± 0.15     | 1.87 ± 0.24     | 1.98 ± 0.22                 |
| Tyr        | 2.36 ± 0.62     | 1.96 ± 0.20     | 2.23 ± 0.30                 |
| Val        | 3.64 ± 0.20     | 3.83 ± 0.36     | 3.59 ± 0.32                 |
| Met        | 1.68 ± 0.11     | 1.58 ± 0.18     | 1.63 ± 0.06                 |
| Ile        | 3.42 ± 0.16     | 2.92 ± 0.28     | 3.26 ± 0.21                 |
| Leu        | 4.63 ± 0.21     | 4.32 ± 0.42     | 4.35 ± 0.09                 |
| Phe        | 2.94 ± 0.14     | 2.65 ± 0.22     | 2.88 ± 0.15                 |
| Lys        | 4.16 ± 0.20     | 4.30 ± 0.33     | 4.18 ± 0.30                 |
| Cys        | 1.35 ± 0.16     | 0.40 ± 0.10     | 1.41 ± 0.23                 |

All values are mean ± SD,  $n = 4$ . Cys = cysteine.

the PTC reagent must be stored at  $-20^{\circ}\text{C}$  under the inert gas to prevent breakdown (7) and a high-vacuum system (50–100 mtorr; 1 torr = 133.322 Pa) is needed to remove the excess reagent and byproducts that could interfere with the main peaks and to improve the reproducibility (3). In this chapter, we did not use the high-vacuum system, so this seems to be one of the reasons why the RSD values in the PTC-derivatives exceeded 5%.

30. We concluded that BTC and BZTC derivatization was less fastidious compared to PTC derivatization.
31. Calibration graphs for BTC-derivatives showed good linearity in the range 0.5–2.5 nmol (not shown with Fig. or Table). The correlation coefficients ( $\gamma$ ) of the calibration graphs for all of the derivatives were highly significant ( $p < 0.001$ ), and the lowest value was  $\gamma = 0.926$  for the cystine derivative, but it could be used for quantitative analysis.
32. Calibration graphs for all of the BZTC derivatives also showed good linearity in the measured range (0.125–5 nmol). The correlation coefficients ( $\gamma$ ) of the calibration graphs were also highly significant ( $p < 0.001$ ) and exceeded 0.99, except for cysteine (0.952) and glutamine (0.976).

**Table 5**  
**Amino Acid Compositions of BSA Determined by BTC and PTC-Derivatives Method Compared to the Ion-Exchange Chromatography and the Analytical Results Reported in the Literature**

| Amino acid | BTC derivatives <sup>a</sup> | BZTC derivatives <sup>c</sup> | PTC derivatives <sup>a</sup> | PTC derivatives <sup>a,c</sup> | Ion-exchange chromatography <sup>a</sup> | Automatic analyzer <sup>d</sup> | Sequence <sup>e</sup> | Sequence <sup>f</sup> |
|------------|------------------------------|-------------------------------|------------------------------|--------------------------------|------------------------------------------|---------------------------------|-----------------------|-----------------------|
| Asp        | 54.85 ± 3.01 <sup>b</sup>    | 57.5 ± 4.76 <sup>b,g</sup>    | 53.78 ± 4.75 <sup>b</sup>    | 52.2 ± 0.42 <sup>b,h</sup>     | 54.49 ± 3.51 <sup>b</sup>                | 55.0                            | 53                    | 41                    |
| Glu        | 73.52 ± 5.04                 | 79.8 ± 5.54                   | 72.61 ± 0.94                 | 78.5 ± 2.92                    | 75.99 ± 4.14                             | 82.0                            | 78                    | 59                    |
| Ser        | 24.60 ± 0.45                 | 24.4 ± 1.60                   | 26.22 ± 4.38                 | 22.3 ± 1.92                    | 25.84 ± 0.15                             | 26.9                            | 28                    | 28                    |
| Gly        | 16.68 ± 0.28                 | 13.8 ± 1.42                   | 14.96 ± 0.50                 | 16.3 ± 2.03                    | 16.31 ± 0.37                             | 15.3                            | 15                    | 16                    |
| His        | 22.39 ± 0.49                 | 16.4 ± 3.22                   | 20.85 ± 2.40                 | 18.4 ± 0.70                    | 17.60 ± 0.87                             | 17.0                            | 17                    | 18                    |
| Arg        | 21.94 ± 0.22                 | 23.0 ± 2.55                   | 31.06 ± 1.51                 | 23.9 ± 1.80                    | 22.98 ± 0.17                             | 23.6                            | 23                    | 25                    |
| Thr        | 30.35 ± 0.82                 | 32.4 ± 1.52                   | 31.16 ± 0.76                 | 32.6 ± 0.80                    | 32.05 ± 0.94                             | 33.0                            | 34                    | 34                    |
| Ala        | 46.07 ± 0.94                 | 47.6 ± 2.24                   | 38.17 ± 0.27                 | 40.0 ± 0.28                    | 46.06 ± 1.33                             | 45.2                            | 46                    | 46                    |
| Pro        | 26.85 ± 1.13                 | 28.3 ± 2.65                   | 29.27 ± 0.78                 | 32.7 ± 0.35                    | 28.60 ± 2.11                             | 28.5                            | 28                    | 29                    |
| Tyr        | 20.47 ± 0.55                 | 22.9 ± 1.56                   | 20.72 ± 0.60                 | 20.3 ± 2.12                    | 20.22 ± 1.22                             | 19.9                            | 19                    | 20                    |
| Val        | 38.61 ± 0.85                 | 38.2 ± 1.50                   | 36.87 ± 0.84                 | 40.2 ± 2.42                    | 37.28 ± 0.96                             | 35.4                            | 36                    | 36                    |
| Met        | 4.61 ± 0.28                  | 4.2 ± 0.25                    | 6.97 ± 0.48                  | 4.6 ± 0.56                     | 4.43 ± 0.11                              | 3.8                             | 4                     | 4                     |
| Ile        | 15.25 ± 0.29                 | 15.9 ± 1.09                   | 18.67 ± 0.55                 | 16.9 ± 0.25                    | 15.65 ± 0.32                             | 14.1                            | 14                    | 14                    |
| Leu        | 61.77 ± 0.26                 | 66.6 ± 2.02                   | 60.88 ± 0.65                 | 63.3 ± 3.68                    | 61.13 ± 0.70                             | 59.2                            | 61                    | 60                    |
| Phe        | 29.20 ± 1.07                 | 28.7 ± 2.44                   | 29.26 ± 0.65                 | 30.3 ± 1.70                    | 27.27 ± 0.80                             | 2.1                             | 26                    | 26                    |
| Lys        | 58.75 ± 1.97                 | 58.1 ± 2.73                   | 54.58 ± 4.65                 | 64.4 ± 2.42                    | 59.42 ± 3.65                             | 36.0                            | 59                    | 58                    |
| Trp        | —                            | —                             | —                            | —                              | —                                        | —                               | 2                     | 2                     |
| Cys        | 26.82 ± 3.60                 | 12.4 ± 1.60                   | 9.32 ± 0.59                  | 25.8 ± 5.30 <sup>g</sup>       | 28.66 ± 5.05                             | —                               | 35                    | 35                    |
| Cyt        | —                            | 12.5 ± 1.63                   | —                            | —                              | —                                        | —                               | —                     | —                     |
| Gln        | —                            | —                             | —                            | —                              | —                                        | —                               | —                     | 20                    |
| Asn        | —                            | —                             | —                            | —                              | —                                        | —                               | —                     | 12                    |
| Total      | 572.73                       | 582.7                         | 555.35                       | 582.7                          | 573.98                                   | 582.2                           | 578                   | 583                   |

Cys = cysteine.

<sup>a</sup>*n* = 3, Data from **ref. 17**.

<sup>b</sup>The values were recalculated on the basis of a total of 583 residues considering the fact that two Trp residues were completely destroyed and cysteine residues were substantially destroyed during acid hydrolysis.

<sup>c</sup>Data from **ref. 18**.

<sup>d</sup>Data from **ref. 24**.

<sup>e</sup>Data from **ref. 25**.

<sup>f</sup>Data from **ref. 26**.

<sup>g</sup>*n* = 12.

<sup>h</sup>*n* = 4.



33. In the PTC-derivatives, cystine+cysteine showed the lowest  $\gamma$  value (0.967), but it could be used for quantitative analysis. However, there is a report that the linearity for PTC-cystine was so poor that it could not be used for quantitative analysis (22).
34. Amino acid compositions of soybean meal determined by the BTC, BZTC, and PTC-derivatives compared to the ion-exchange chromatography were shown in **Table 3**.
35. In the BTC-derivatives compared to ion-exchange chromatography, most of amino acids showed a deviation of less than 5%, except for histidine (13%) and isoleucine (9%). But BZTC and PTC-derivatives, most of amino acids showed a deviation of more than 5%. Reproducibility was superior in the BTC and BZTC-derivatives compared to PTC-derivatives.
36. **Table 4** shows the amino acid compositions of whole egg determined by the BTC and PTC-derivatives compared to ion-exchange chromatography. We did not determine BZTC-derivatives, so we could not show the data of the BZTC-derivatives. As in the soybean meal, the data of BTC-derivatives were well matched with those of ion-exchange chromatography. Reproducibility was also superior in the BTC-derivatives as in the soybean meal.
37. **Table 5** shows amino acid compositions of BSA recalculated on the basis of a total of 583 residues considering the fact that 2 Trp residues were completely destroyed, and cysteine residues were substantially destroyed during the hydrolysis.
38. It has been shown by sequence analysis that BSA has 17 disulfide bonds (cystine) and one free cysteine (23). In this chapter, the number of BZTC-cystine and BZTC-cysteine residues was 12.5 and 12.4, respectively. It is known that a substantial amount of cysteine and cystine is destroyed during the hydrolysis of protein with 6 M HCl when the hydrolysis was carried out without conversion of these amino acids. Cysteine and cystine determination seems to give unavoidable error, unless they are modified to other compounds, i.e., pyridylethyl-cysteine or cysteic acid, prior to quantification (18,21).

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## Amino Acid Measurement in Body Fluids Using PITC Derivatives

Roy A. Sherwood

### 1. Introduction

Amino acid chromatography is used to detect both primary disorders of amino acid metabolism (e.g., maple syrup urine disease) and disorders of renal tubular reabsorption (e.g., cystinuria) (*see* **Notes 1–4**). In most patients with disorders in the former group, the abnormal amino acids are clearly increased in both plasma and urine, although the abnormality is usually more pronounced in urine. In disorders of renal tubular transport, only the urine amino acids will be abnormal, thus these conditions will be missed if plasma alone is studied. Quantitative amino acids are measured in blood samples from children who are having their amino acids levels manipulated by dietary restriction of natural protein. Derivatization of amino acids is required before analysis by high-performance liquid chromatography (HPLC). A number of alternative pre- or postcolumn derivatization methods have been described (**1**). In the method described here amino acids in standards, urine, or deproteinized serum/plasma are reacted with phenylisothiocyanate (PITC) in the presence of a coupling solvent. The phenylthiocyanate (PTC) derivatives are then applied to a reverse-phase (RP) HPLC column and separated using a gradient elution system. The PTC amino acids are detected using an ultraviolet (UV) detector (254 nm) and an electrochemical detector (ECD) in series (**2**).

### 2. Materials

#### 2.1. Equipment

1. Gradient HPLC system with ODS Hypersil 5  $\mu\text{m}$ , 4 mm  $\times$  25-cm column with precolumn filter.
2. UV detector and amperometric electrochemical detector.

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**Table 1**  
**Composition of Coupling Reagent**

| Chemical             | 2 samples        | 10 samples        | 30 samples         |
|----------------------|------------------|-------------------|--------------------|
| Methanol (Hipersolv) | 70 $\mu\text{L}$ | 350 $\mu\text{L}$ | 1050 $\mu\text{L}$ |
| Triethylamine        | 10 $\mu\text{L}$ | 50 $\mu\text{L}$  | 150 $\mu\text{L}$  |
| Distilled water      | 10 $\mu\text{L}$ | 50 $\mu\text{L}$  | 150 $\mu\text{L}$  |
| neat PITC            | 10 $\mu\text{L}$ | 50 $\mu\text{L}$  | 150 $\mu\text{L}$  |

## 2.2. Reagents and Solutions

1. Acetic acid 6% (v/v).
2. Buffer A: 10 mM sodium acetate adjusted to pH 6.4 with 6% acetic acid. Should be prepared fresh for each run. Two liters will be enough to run 15 samples.
3. Buffer B: 10 mM acetonitrile, sodium acetate 60% (v/v) adjusted to pH 6.4 with 6% acetic acid. Stable at room temperature for 14 d. One liter will be enough to run 25 samples. Acetonitrile is toxic by ingestion, inhalation, and skin contact.
4. Precipitating reagent containing internal standard: 10% sulphosalicylic acid (SSA), 750  $\mu\text{M}$  norleucine. Stable for 6 mo at 4°C.
5. PITC coupling reagent. Prepared fresh, as per **Table 1**. PITC is toxic by ingestion, irritating to skin/eyes, and has been reported to have teratogenic effects. Triethylamine is toxic by ingestion and is irritating to the eyes.
6. Amino acid standards (Sigma Chemicals, Poole, UK). Choice dependent on amino acids of interest. Available standards (17 amino acid standard, acids, and neutrals, basics) can be supplemented by specific amino acids as required.

## 3. Methods

### 3.1. Preparation of Samples

Plasma, random urine (children) or 24 h urine (adults) can be used (*see* **Notes 5 and 6**). Heparinized plasma is preferred, there are differences between plasma and serum amino acids. In acutely ill children with a suspected inherited metabolic defect, blood should be taken on admission. If the acute illness occurred some days before investigation, dietary protein may have been withdrawn and an amino acid abnormality may not be apparent. In a nonacute situation, a postabsorptive or fasting sample is preferred (adults and older children fasted overnight, smaller children 6–8 h). This is because plasma amino acids may be significantly increased 1–2 h after a protein meal. In neonates samples should be taken 2–3 h after feeding.

1. Urine. For urine, determine the creatinine in mmol/L (*see* **Note 7**). For creatinine concentrations below 5 mmol/L, take 50  $\mu\text{L}$  of urine, make up to 100  $\mu\text{L}$  with distilled water, and multiply result by two. For urine creatinine concentrations

above 5 mmol/L, take 20  $\mu\text{L}$  of urine, make up to 100  $\mu\text{L}$  with distilled water, and multiply result by 5. For urine creatinine concentrations below 1 mmol/L, another sample should be requested as the urine is too dilute to give meaningful results.

2. Add 50  $\mu\text{L}$  of SSA precipitating reagent to tube. Mix and leave for 30 min at 4°C.
3. Plasma/serum: Pipet 100  $\mu\text{L}$  of sample into a microfuge tube. Add 50  $\mu\text{L}$  of SSA precipitating reagent to tube. Mix and leave for 30 min at 4°C.
4. Standards: 10  $\mu\text{L}$  added to 90  $\mu\text{L}$  of distilled water. Additional amino acid standards can be added at this point with corresponding reduction in the water, e.g., add 10  $\mu\text{L}$  of a 1 mg/mL standard. Standards as prepared give 250  $\mu\text{mol/L}$  concentrations except cystine (125  $\mu\text{mol/L}$ ). Add 50  $\mu\text{L}$  of precipitating reagent containing internal standard to tube. Mix and leave for 30 min at 4°C.

### 3.2. Derivative Formation

1. Centrifuge tubes for 10 min at 14,000g.
2. Transfer 50  $\mu\text{L}$  of supernatant to an appropriately labeled microfuge tube. Add 200  $\mu\text{L}$  acetonitrile and mix.
3. Add 40  $\mu\text{L}$  PITC/coupling reagent and vortex mix. Leave to react for 20 min.
4. Evaporate to dryness at a temperature less than 45°C. *It is essential that the derivatives are dry, but not cooked/desiccated.* The derivatives (dry) can be left at this stage at 4°C (no more than 72 h).
5. Dissolve in 200  $\mu\text{L}$  of sodium acetate buffer (Buffer 1). Mix.
6. Centrifuge and remove 200  $\mu\text{L}$  of supernatant to HPLC vial with microinsert for chromatography.

### 3.3. Chromatography

1. HPLC operating conditions: Column oven temperature: 30°C, sample cooler, 4°C, injection volume 25  $\mu\text{L}$ . Electrochemical detector settings: potential +1.1 V (oxidation), sensitivity 3  $\mu\text{A}$ . UV detector: wavelength 254 nm. If the UV detector and the electrochemical detector are both used in series the UV detector must be placed first.
2. Gradient: The solvent gradient is set as shown in **Table 2**.
3. Allow the system to equilibrate and run a gradient off-line while preparing samples and standards. The cycle time on the autosampler is 95 min.
4. Inject the first standard in the queue  $\times 2$ . (The first sample retention times are often too variable for calibration purposes.)
5. Typical retention times and relative (to the internal standard norleucine) retention times are shown in **Table 3** (*see Notes 8–12*).
6. Calculation: To obtain calculation factor:

$$\text{Calculation factor} = \frac{\text{conc of std } (\mu\text{mol/L}) \times \text{area/height of Internal Standard}}{\text{area/height of Standard}}$$

The peak area/height of unknowns is multiplied by this factor to give the concentration in  $\mu\text{mol/L}$ .

**Table 2**  
**HPLC Solvent Gradient**

| Time (min) | Buffer 1 (%)<br>Sodium acetate,<br>pH 6.4 | Buffer 2 (%)<br>60% acetonitrile/<br>sodium acetate, pH 6.4 | Flow rate<br>mL/min |
|------------|-------------------------------------------|-------------------------------------------------------------|---------------------|
| 0          | 100                                       | 0                                                           | 1.0                 |
| 20         | 87                                        | 13                                                          | 1.0                 |
| 65         | 45                                        | 55                                                          | 1.0                 |
| 67.5       | 0                                         | 100                                                         | 2.0                 |
| 70         | 0                                         | 100                                                         | 2.0                 |
| 75         | 100                                       | 0                                                           | 2.0                 |
| 80         | 100                                       | 0                                                           | 1.0                 |
| 85         | 100                                       | 0                                                           | 1.0                 |
| 95         | 100                                       | 0                                                           | 1.0                 |

7. Interfering substances: Vigabatrin will form a PTC derivative. The cephalosporin group of antibiotics potentially interfere with the UV detection method and might cochromatograph with glycine.

#### 4. Notes

1. In approx 99% of cases, the amino acid pattern will be within normal limits. Variations occur for physiological reasons, particularly in premature neonates, but in comparison to pathological processes the variation is small. Most inherited disorders of amino acid metabolism produce gross changes in the observed pattern. Children diagnosed with inherited diseases involving amino acids are often treated with artificial feeds excluding certain amino acids, and they can get very low levels of essential amino acids, necessitating frequent monitoring.
2. Pathological cases: Most are fairly obvious and the specific amino acid will increase dramatically, e.g., in phenylketonuria, cystinuria, maple syrup urine disease, cystathioninuria, and hypophosphatasia the increase on HPLC will be off scale by 2–10 fold. Homocystinuria is an exception to this in that homocystine is normally undetectable by HPLC. A positive cystine screen with a normal cystine on thin layer chromatography should alert you to the possibility of homocystinuria. Using the HPLC method, it chromatographs immediately after the internal standard and any visible peak must be considered abnormal, but this may not be easy to pick up. Methionine concentrations are often high, secondary to liver disease, such as tyrosinemia and galactosemia.
3. Symptoms suggestive of specific disorders:
  - a. Cataracts. Common in homocystinuria.
  - b. Deafness. Occurs in arginosuccinic aciduria. Gives three peaks on HPLC (two anhydrides and the acid itself).

**Table 3**  
**Mean Retention Times and Relative Retention Times for the Amino Acids Most Likely to be Found in Body Fluids**

| Amino acid                    | Retention time (min) | Relative retention time |
|-------------------------------|----------------------|-------------------------|
| Phosphoserine                 | 3.82                 | 0.084                   |
| Aspartic acid                 | 5.44                 | 0.12                    |
| Glutamic acid                 | 6.83                 | 0.15                    |
| $\gamma$ -Aminoadipic acid    | 10.35                | 0.229                   |
| Hydroxyproline                | 12.25                | 0.271                   |
| Phosphoethanolamine           | 12.45                | 0.277                   |
| Serine                        | 14.52                | 0.321                   |
| Glycine                       | 15.53                | 0.345                   |
| Asparagine                    | 15.55                | 0.347                   |
| Sarcosine                     | 16.91                | 0.375                   |
| $\beta$ -Alanine              | 17.40                | 0.387                   |
| Taurine                       | 19.00                | 0.421                   |
| $\gamma$ -Aminobutyric acid   | 20.47                | 0.454                   |
| Citrulline                    | 20.65                | 0.458                   |
| Threonine                     | 21.11                | 0.468                   |
| Alanine                       | 21.75                | 0.482                   |
| $\beta$ -Aminoisobutyric acid | 22.27                | 0.494                   |
| Proline                       | 23.49                | 0.521                   |
| Histidine                     | 23.91                | 0.532                   |
| Carnosine                     | 25.71                | 0.569                   |
| Arginine                      | 28.22                | 0.626                   |
| Methyl histidine              | 28.39                | 0.630                   |
| $\alpha$ -Aminobutyric acid   | 28.60                | 0.634                   |
| Anserine                      | 29.06                | 0.644                   |
| Tyrosine                      | 35.02                | 0.778                   |
| Valine                        | 35.76                | 0.795                   |
| Ethanolamine                  | 37.66                | 0.835                   |
| Methionine                    | 38.10                | 0.845                   |
| Cystathionine                 | 38.20                | 0.848                   |
| Cystine                       | 40.88                | 0.907                   |
| Isoleucine                    | 42.79                | 0.950                   |
| Leucine                       | 43.47                | 0.965                   |
| Norleucine                    | 45.00                | —                       |
| Hydroxylysine                 | 47.78                | 1.060                   |
| Phenylalanine                 | 48.20                | 1.069                   |
| Ornithine                     | 49.67                | 1.102                   |
| Tryptophan                    | 50.16                | 1.112                   |
| Lysine                        | 52.79                | 1.170                   |



- c. Abnormal X-ray. Hypophosphatasia (serum alkaline phosphatase activity will be  $< 25$  IU/L).
  - d. Renal Stones. Cystinuria (also possibly xanthine/purine stones which are radiolucent in disorders such as Lesch-Nyhan syndrome).
  - e. Organic acidurias. Most organic acidurias produce a metabolic acidosis, a urine pH  $< 6.0$  and coma. Some include hypoglycemia and abnormal blood lactate/pyruvate ratio. There are often abnormalities in the amino acid pattern, but these are not always helpful.
4. Physiological/spurious increases in amino acids:
    - a. Tyrosine. Transient neonatal tyrosinaemia is common in the first 6–8 wk of life for premature infants. If it persists after this time, then either tyrosinaemia or liver disease should be suspected (the latter is about 10 times more likely). The cutoff for tyrosine levels giving rise to recall in neonatal screening programs is typically  $500 \mu\text{mol/L}$ . However, recent experience has shown that tyrosine concentrations in tyrosinemia type I can be around  $200 \mu\text{mol/L}$  with a neonatal presentation.
    - b. Threonine. Increased in neonates when compared to adults. It is also increased in liver disease.
    - c. Taurine. Increased taurine can occur in catabolic states, but there is no specific disorder involving taurine.
    - d. Glycine/aspartic acid/glutamic acid. Hyperglycinemia can occur in organic acidurias, valproate treatment, and in subjects with a low calorie intake. In urine samples with bacterial contamination, glutamine is deaminated to glutamic acid, aspartic acid may be increased, and hippuric acid is broken down to glycine. A pH  $> 7.0$  is suspicious in such cases.
    - e. Lysine/cystine. Both are often raised in the first 6 mo of life.  
A generalized amino aciduria is often seen in very sick neonates with poor peripheral circulation. Particular increases are seen in alanine, glycine and proline related to disturbances of lactate/pyruvate metabolism.
  5. Samples should *not* be hemolyzed. Samples are stored *frozen* if not assayed immediately. Samples left unseparated will have increases in taurine, phosphoethanolamine, aspartic, and glutamic acid. Free cystine and homocystine in plasma should only be measured in samples that have been separated and frozen within 30 min of being taken. Care should be taken not to aspirate the buffy coat with the plasma, this leads to increases in taurine, aspartic acid, glutamic acid, and phosphoethanolamine.
  6. A 24-h collection is required for monitoring treatment of cystinuric patients (a compromise for adults is to fast overnight and collect a 4-h (still fasting) collection (from 0630–1030) (3).
  7. An aliquot of urine should be taken on receipt for creatinine estimation, the remainder of the urine should be frozen until analyzed. Excretion rates of amino acids may vary independently of the creatinine over a 24-h period. Diurnal variation has been observed.

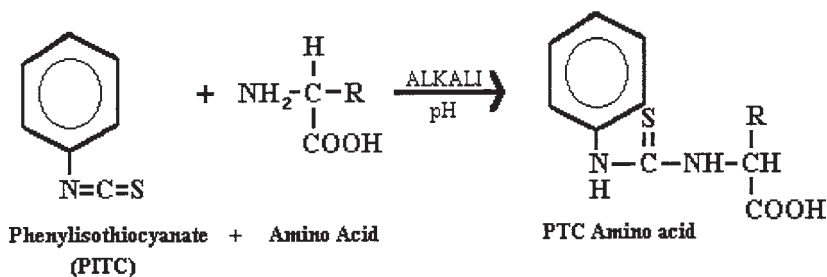


Fig. 1. Formation of the PTC derivatives of amino acids.

8. Identify peaks on samples by cross reference to standards and to relative retention times obtained on previous runs. Peaks can be identified by cochromatography. Mix equal volumes of a derivatized standard and sample, cochromatography and the amino acid is probably the same as in the sample, splitting of the peak — it is not the same.
9. Several of the peaks are pH dependent and move slightly as the buffer ages, in addition, separations tend to vary with different batches of the column. Typically, proline moves to merge with histidine or alanine. Hydroxyproline, ethanolamine, and phosphoethanolamine also move, but this is seldom a problem.
11. Histidine can be identified by its characteristic tailing peak.
12. Threonine and citrulline tend to cochromatograph, citrulline runs slightly fast of threonine.
13. Glycine and asparagine occasionally run together, glycine is faster than asparagine.

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## Determination of Proteins, Phosphatidylethanolamine, and Phosphatidylserine in Lipid-Rich Materials by Analysis of Phenylthiocarbamyl Derivatives

Margareta Stark and Jan Johansson

### 1. Introduction

In classical, ninhydrin-based amino acid analysis (1,2), the ion-exchange matrix used for separation becomes contaminated upon consecutive analyzes of extremely lipid-rich samples; in our experience, already after approx 20–30 samples. Therefore, in analysis of lipid-rich material we focused on amino acid analysis involving reversed phase (RP) chromatography, because lipids are soluble in the organic solvents commonly used for elution and regeneration of such columns (e.g., acetonitrile and 2-propanol). Precolumn derivatization with phenylisothiocyanate (PITC) of protein hydrolysates from physiological samples followed by RP-HPLC is equivalent in analytical quality to the classical ion-exchange chromatography/ninhydrin method (3), and is sensitive down to at least 10 pmol (4). PITC reacts both with primary and secondary amines, the reproducibility is high and the phenylthiocarbamyl (PTC) amino acid derivatives are stable for months when stored dry at  $-20^{\circ}\text{C}$ , or for days in solution at ambient temperature (3,4). This approach can be used conveniently for analysis of lipid-rich material, where at least 300 samples can be analyzed with the same column, i.e., the column lifetime is comparable to those encountered during analysis of lipid-free samples (5).

During isolation of proteins from lipid-rich sources, we have found it necessary to monitor both protein and phospholipid profiles, at least at early stages of the purification scheme. The phospholipids are usually detected by phosphorous analysis, but with the PTC/RP-HPLC method, phospholipids contain-

ing a primary amino group can be determined simultaneously with the proteins. Phosphatidylethanolamine subjected to hydrolysis and subsequent analyzed by the PTC/RP-HPLC method show a peak eluting between PTC-Arg and PTC-Tyr, with a linear range between 1–50 nmol, but lower levels are detectable. With the classical ion-exchange/ninhydrin amino acid analysis, where ethanolamine coelutes with Lys and consequently neither of these compounds can be determined. Phosphatidylserine analyzed in the same way by the PTC/RP-HPLC method gives one peak that coelutes with PTC-Ser, with a linear range between 10 pmol–50 nmol. When monitoring the protein and phospholipid profile after chromatography, it is feasible to determine when serine derived from phosphatidylserine elutes by looking at the overall PTC-amino acid profile and amounts. Phosphatidylcholine contains a quaternary amino group and cannot be detected with the PTC/RP-HPLC method (5). A major advantage with PTC/RP-HPLC analysis, in addition to allowing repeated analysis of lipid-rich samples, is that phosphatidylethanolamine and phosphatidylserine can be determined simultaneously with the proteins, thereby reducing the need for separate phosphorous analysis during protein purification. A sample from a lung phospholipid fraction, which contains about 2% protein and 98% lipids, was hydrolyzed and analyzed by the PTC/RP-HPLC method (Fig. 1).

## 2. Materials

### 2.1. Hydrolysis

1. Glass tubes for hydrolysis and derivatization are 6–7 × 35 mm. Tubes for mixing derivatization solutions are 10 × 75 mm. Submit all glass tubes to pyrolysis (400–500°C for 3–4 h) before use to remove any contaminating material. Recommended glass quality: soda-lime or Duran.
2. Hydrolysis solution: 6 M HCl with 0.5% (w/v) phenol.

### 2.2. Derivatization

1. Amino acid standard in 0.1 M HCl (e.g., Amino Acid Standard H, Pierce). A stock solution of 100 μM of each amino acid in water can be stored at –20°C for several months.
2. Phosphatidylethanolamine (e.g., dipalmitoylphosphatidylethanolamine, Fluka) dissolved in chloroform, e.g., 10 mM stock solution. Store in capped glass tube at –20°C.
2. Triethylamine (sequanal grade, Pierce). Store at 4°C. NB! Avoid inhalation and skin contact. Wear protective clothing. Work in a ventilated hood and use gloves.
3. Phenylisothiocyanate (PITC) (sequanal grade, Pierce). Store opened ampoule in a screw-capped glass tube under nitrogen in the dark at ambient temperature. Discard when yellow. Caution: Avoid inhalation and contact with skin and eyes. Work in a ventilated hood, use gloves, and wear protective clothing.

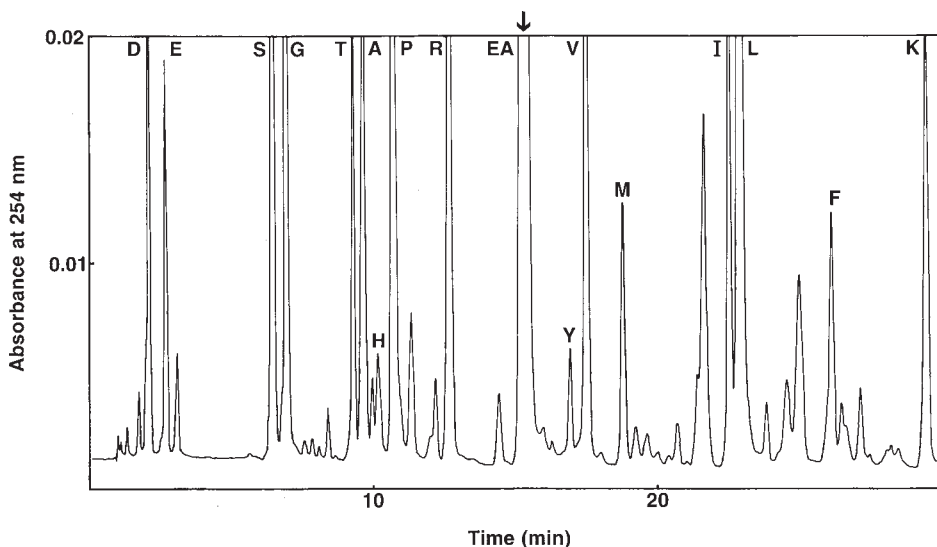


Fig. 1. RP-HPLC of PTC derivatives from a sample obtained from a phospholipid/Lipoprotein fraction from porcine lung. The sample was hydrolyzed and analyzed by the PTC/RP-HPLC method. The PTC derivatives are identified with the one-letter code for the corresponding amino acid or EA for ethanolamine. The arrow identifies the peak that appears when dipalmitoyl-phosphatidylethanolamine was hydrolyzed separately.

4. Pre-derivatization solution: 99.5% ethanol/water/triethylamine (2:2:1, v/v/v), 40  $\mu$ L per sample is required. Mix in a 10  $\times$  75-mm glass tube. Prepare fresh.
5. 50% ethanol: 99.5% ethanol/Milli-Q water (1:1, v/v). Store in a screw-capped glass tube at 4°C.
6. Derivatization solution: 99.5% ethanol/triethylamine/PITC (7:2:1, v/v/v). Mix in a 10  $\times$  75-mm glass tube. Prepare fresh.

### 2.3. Separation

1. Mobile phase A: 30 mM phosphate buffer, pH 6.30–6.80 (*see Note 1*). Dissolve 1.2 g NaOH in approx 800 mL MilliQ water, adjust to the pH required with 1 M  $\text{H}_3\text{PO}_4$ . Adjust to 1000 mL and filter through 0.2  $\mu$ m. Degas before use. Use fresh or store at 4°C overnight. If stored longer (a few days), filter again through 0.2  $\mu$ m before use.
2. Mobile phase B: 60% acetonitrile in water. Mix 600 mL acetonitrile with 400 mL Milli-Q water. Store at ambient temperature. Stable for several weeks. Degas before use.
3. Column:  $\text{C}_{18}$  reversed phase HPLC column, Spherisorb S3 ODS2, 4.6  $\times$  100 mm, packed with 3- $\mu$ m particles (Waters).

**Table 1**  
**Gradient Program for Separation of the PTC-Derivatives**

| Time (min)           | % A     | % B     |
|----------------------|---------|---------|
| 0                    | 100     | 0       |
| 1                    | 95      | 5       |
| 26 ( <b>Note 6</b> ) | 68 (66) | 32 (34) |
| 28                   | 0       | 100     |
| 34                   | 0       | 100     |
| 37                   | 100     | 0       |

Flow rate 1.0 mL/min, the gradient is linear. Next sample injected after 8 min delay (45 min total per sample).

4. High-performance liquid chromatography (HPLC) instrumentation: Any analytical instrument equipped with two separate pumps, absorbance detector (254 nm), integrator, heating block, and a temperature control unit.

### 3. Methods

#### 3.1. Hydrolysis

1. Pipet samples/phospholipid standards (*see Note 2*) present in volatile solution in  $6 \times 35$ -mm glass tubes and dry the samples under a stream of nitrogen or vacuum. For the acid hydrolysis add 40  $\mu\text{L}$  6 M HCl/0.5% phenol to each tube, evacuate and seal the tubes. Incubate at 110°C for 20–24 h, open and dry the samples under vacuum.

#### 3.2. Derivatization

1. Pipet amino acid standard (1000–5000 nmol/aa) in a  $6 \times 35$ -mm glass tube and dry under vacuum (*see Note 3*).
2. Add 40  $\mu\text{L}$  pre-derivatization solution (99.5% ethanol/water/triethylamine; 2:2:1) (**6**) to each of the hydrolyzed samples/phospholipid standards and amino acid standard, seal the tubes with parafilm, and mix for 10 s. Remove the parafilm and redry under vacuum (*see Note 4*).
3. For derivatization add 3  $\mu\text{L}$  50% ethanol to each tube (**7**), make sure that an aqueous film is obtained at the bottom of the tube by mixing or turning the tube. Then add 7  $\mu\text{L}$  of derivatization solution (99.5% ethanol/triethylamine/PITC, 7:2:1) and seal the tubes with parafilm. Mix vigorously (vortex) for 10 s and allow the reaction to occur for 15–30 min at ambient temperature. Remove the parafilm and dry the samples under vacuum overnight to remove excess reagent (*see Note 5*).
4. Seal the tubes with parafilm and store at  $-20^\circ\text{C}$ , if not immediately subjected to chromatography. The samples can be stored for several months in the freezer.

### **3.3. Separation of PTC-derivatives by HPLC**

1. Dissolve the derivatized and dried standard and samples in 100  $\mu$ L mobile phase A. Filter lipid-containing samples through 0.45  $\mu$ m.
2. Equilibrate the column with mobile phase A, flow rate 1.0 mL/min, and column temperature 36°C.
3. Inject 5–90  $\mu$ L standard or sample on the column and carry out gradient elution as shown in **Table 1** at a flow rate of 1.0 mL/min and a column temperature of 36°C. Allow the column to equilibrate with mobile phase A for 8 min between each run, i.e., one sample injected every 45 min.
4. Quantify the PTC-amino acids and PTC-ethanolamine by comparison of the peak areas with those of the standard mixtures.
5. Store the column between runs in 60% acetonitrile (*see Note 7*).

### **4. Notes**

1. Start at pH 6.3–6.4. The elution position of His is pH sensitive. Lower pH — earlier retention time of His, higher pH — later retention time. After several runs His usually elutes close to, or coelutes with Pro, then raise the pH (0.05–0.1 pH U). Do not exceed pH 7.0, the matrix of the column is not stable in basic pH.
2. Phosphatidylethanolamine standard, suggested amounts: hydrolyze 10 nmol and analyze 10% (i.e., 1 nmol) or make a dilution series. Analysis of less than 500 pmol gives unreliable results. A phosphatidylserine standard is not necessary, the serine from the amino acid standard can be used.
3. Do not hydrolyze the amino acid standard. 50 pmol per amino acid analyzed is suitable for AUFS 0.005 at 254 nm.
4. This step is indispensable. PITC do not react with the amino groups if any residual acid from the hydrolysis is present.
5. PITC has low volatility. Remove by high vacuum in a dessicator with solid NaOH for at least 12 h. If the reagent peak is too high, redry for longer time (e.g., 24 h).
6. Lower %B — later retention times from Ser. Higher % B — earlier retention times from Ser.
7. After 30–50 runs, wash the column with methanol/2-propanol, 1/1 (v/v) to minimize lipid contamination. Store the column in at least 60% acetonitrile.

### **Acknowledgments**

We are grateful to Carina Palmberg for skillful assistance and Professor Hans Jörnvall for support.

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## Analysis of *O*-Phosphoamino Acids in Biological Samples by Gas Chromatography with Flame Photometric Detection

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### 1. Introduction

The separation and determination of *O*-phosphoamino acids has been carried out by thin-layer chromatography (1–8), thin-layer electrophoresis (1,6,7,9–14), gel electrophoresis (15), amino acid analyzer (16–18), high-performance liquid chromatography (HPLC) (19–33), capillary zone electrophoresis (CZE) (34,35) and immunoassay (36–38). However, most of these methods require <sup>32</sup>P-labeling for detecting and quantifying *O*-phosphoamino acids, and therefore cannot be used for analysing nonradiolabeled *O*-phosphoamino acids in biological samples. Furthermore, some of them show poor resolution or low sensitivity. HPLC methods based on the precolumn ultraviolet (UV) derivatization with phenyl isothiocyanate (22–24), dabsyl chloride (25), and *N*- $\alpha$ -(2,4-dinitro-5-fluorophenyl)-l-alaninamide (26), the postcolumn fluorescence derivatization with *o*-phthalaldehyde (27–31), the precolumn fluorescence derivatization with 9-fluorenylmethyl chloroformate (32,33), and CZE methods based on the UV derivatization with phenyl isothiocyanate or dabsyl chloride (34,35) were highly sensitive, but some of these methods lack specificity. Furthermore, many of these methods require lengthy separation time, and clean-up of the sample to remove the excess reagent and coexisting substances. On the other hand, the immunoassays by antibodies raised against each *O*-phosphoamino acid have been employed for measuring endogeneously phosphorylated amino acids, but these methods involve some problems in the specificity and crossreactivity of the antibodies. Current methods for the deter-

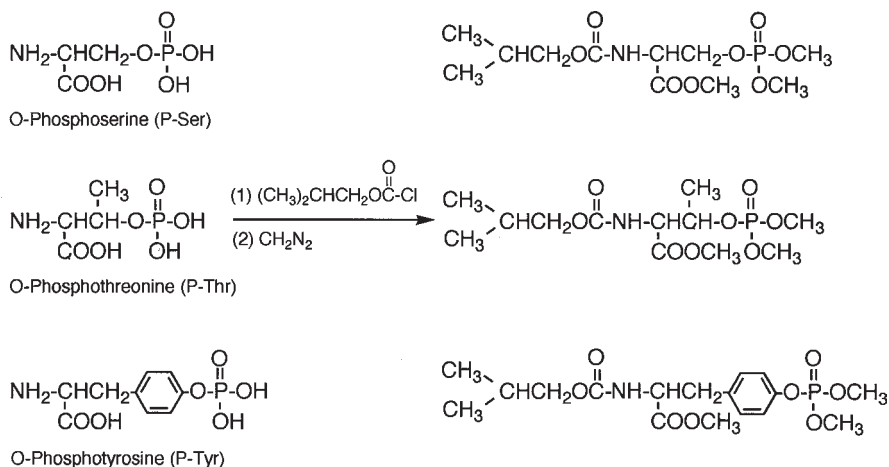


Fig. 1. Derivatization process of *O*-phosphoamino acids.

mination of *O*-phosphoamino acids have also been described in detail in **ref. 39**, and the review in **ref. 40**.

Recently, we have developed a selective and sensitive method for the determination of *O*-phosphoamino acids, such as *O*-phosphoserine (P-Ser), *O*-phosphothreonine (P-Thr), and *O*-phosphotyrosine (P-Tyr) by GC with flame photometric detection (FPD) using a capillary column, in which these compounds were analyzed as their *N*-isobutoxycarbonyl (isoBOC) methyl ester derivatives (**41–49**). By using this method, we demonstrated that the contents of free *O*-phosphoamino acids in tissues (**42,43**) and urine (**48**), and protein-bound *O*-phosphoamino acids in phosphorylated proteins (**44,45**), tissues (**46,49**), and urine (**47–49**) could be rapidly and simply analyzed. In this chapter, selective and sensitive methods for the determination of *O*-phosphoamino acids in biological samples by FPD-GC are described on the basis of the aforementioned results, and optimum conditions and typical problems encountered in the development and application of the methods are discussed.

The derivatization process is shown in **Fig. 1**. *N*-isoBOC methyl esters of *O*-phosphoamino acids can be easily, rapidly, and quantitatively prepared by the reaction with isobutyl chloroformate (isoBCF) by shaking in aqueous alkaline media, followed by esterification with diazomethane. The *O*-phosphoamino acids in peptides or proteins were released from the sample by acid and base hydrolyzes prior to derivatization. The main advantage of this method is that these amino acids can be easily converted in an aqueous medium without any further clean-up procedure, and the derivatives are stable to moisture. Furthermore, the derivatives can be quantitatively and reproducibly resolved as single peaks using a capillary column, and provide an excellent FPD response.

## 2. Materials

### 2.1. Equipment

1. Shimadzu 14A gas chromatograph equipped with a flame ionization detector and a flame photometric detector (P-filter).
2. For the analysis of free *O*-phosphoamino acids, fused silica capillary columns (15 m × 0.53-mm id, 1.0- $\mu$ m film thickness) of crosslinked DB-1701 (14% cyanopropylphenyl-86% methylpolysiloxane, J & W, Folsom, CA) or DB-210 (50% trifluoropropyl-50% methylpolysiloxane, J & W).
3. For the analysis of protein-bound P-Ser/P-Thr and P-Tyr, DB-1707 (15 m × 0.53-mm id, 1.0- $\mu$ m film thickness) and DB-5 (5% phenyl-95% methylpolysiloxane, J & W, Folsom, CA, USA: 15 m × 0.53-mm id, 1.0- $\mu$ m film thickness).
4. Model LK-21 ultradisperser (Yamato Kagaku, Tokyo, Japan).
5. Model RD-41 Centrifugal evaporator (Yamato Kagaku).
6. Pico-Tag workstation (Waters Associates, Milford, MA).
7. Pasteur capillary pipet (Iwaki glass No. IK-PAS-5P).

### 2.2. Reagents

1. Standard *O*-phosphoamino acids, aminophosphonic acids and nonphosphorylated amino acids: *O*-phospho-L-serine (P-Ser), *O*-phospho-D,L-threonine (P-Thr), *O*-phospho-L-tyrosine (P-Tyr), *O*-phosphoethanolamine (PEA), and 2-aminoethylphosphonic acid (AEP) were purchased from Sigma (St. Louis, MO). All of the nonphosphorylated amino acids used were purchased from Ajinomoto (Tokyo, Japan).
2. Internal standards (IS): 2-amino-4-phosphonobutyric acid (APB), 2-amino-7-phosphonoheptanoic acid (APH) and 2-amino-8-phosphonooctanoic acid (APO) as internal standards (IS) were purchased from Sigma.
3. Protein samples: purified commercial proteins, bovine serum albumin (Fraction V, 96–99%), hen egg ovalbumin (Type VII), bovine milk  $\alpha$ -casein, bovine milk  $\beta$ -casein, bovine milk  $\kappa$ -casein, egg yolk phosphovitin, human hemoglobin, bovine heart cytochrome *c*, salmon sperm protamine, calf thymus histone (Type II), bovine brain myelin basic protein, yeast alcohol dehydrogenase, bovine liver catalase, egg white lysozyme, and bovine pancreas chymotrypsin were purchased from Sigma.
4. Derivatizing reagents: isobutyl chloroformate (isoBCF) obtained from Tokyo Kasei Kogyo was used without further purification and stored at 4°C when not in use. *N*-methyl-*N*-nitroso-*p*-toluenesulphonamide and diethyleneglycol monomethyl ether for the generation of diazomethane (50) were obtained from Nacalai Tesuque.
5. Other materials: peroxide-free diethyl ether was purchased from Dojindo Laboratories (Kumamoto, Japan). Distilled water was used after fresh purification with a Model Milli-Q Jr. water purifier (Millipore, Bedford, MA). All other chemicals were analytical grade.

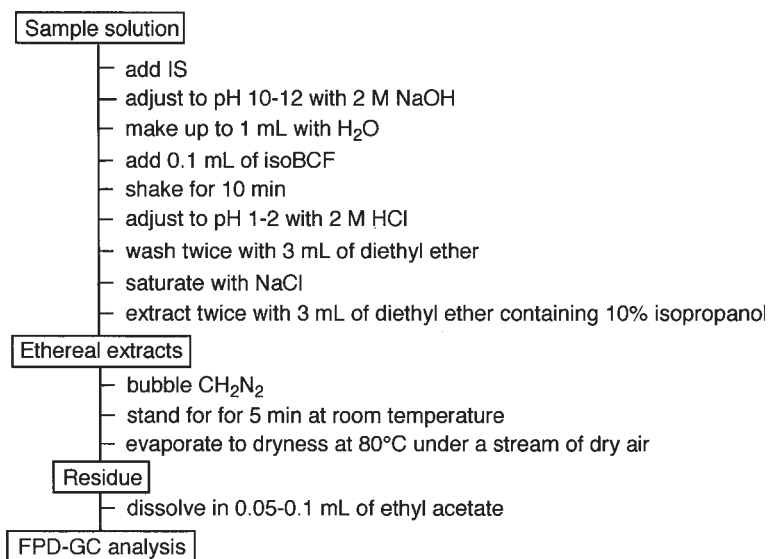


Fig. 2. Schematic flow diagram of the derivatization procedure.

### 2.3. Solutions

1. Standard *O*-phosphoamino acid solutions: each compound is dissolved in distilled water to make up a stock solution at a concentration of 0.1 mg/mL and then stored at 4°C. The working standard solutions are made up freshly, as required by dilution of the stock solution with distilled water. These solutions are stable at 4°C for at least 2 wk.
2. Internal standard solutions: APB, APH, and APO are dissolved in distilled water to make up stock solutions at a concentration of 0.1 mg/mL. These solutions are stable at 4°C for at least 2 wk.
3. Trichloroacetic acid (TCA) solution: TCA is dissolved in distilled water at concentrations of 5% and 20%.

## 3. Method

### 3.1. Derivatization

Derivatization procedure of *O*-phosphoamino acids is shown in **Fig. 2** (see **Note 1**).

1. For the analysis of free *O*-phosphoamino acids, pipet an aliquot of a sample containing 10–2000 ng of each amino acid and 0.1 mL of 2.5 µg/mL APB (IS) (if necessary) into a 10-mL Pyrex glass reaction tube with a PTFE-lined screw cap.
2. For the analysis of *O*-phosphoamino acids in protein hydrolysates, pipet an aliquot of a sample containing 10–500 ng of each amino acid and 0.05–0.1 mL of 1

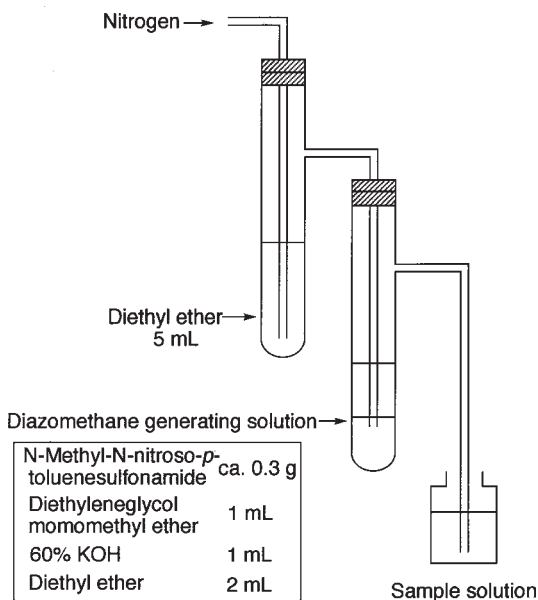


Fig. 3. Apparatus for microscale generation of diazomethane.

$\mu\text{g/mL}$  APB (P-Ser/P-Thr analysis) and APH or APO (for P-Tyr analysis) (IS) (if necessary) into a reaction tube.

- Adjust these mixtures to pH 10.0–12.0 with 2 *M* NaOH and make the total volume up to 1 mL with distilled water.
- Immediately add 0.1 mL of isoBCF to the mixtures and shake at 300 cycles/min (up and down) for 10 min at room temperature (*see Note 2*).
- Readjust the reaction mixture to pH 1.0–2.0 with 2 *M* HCl.
- Extract twice with 3 mL of diethyl ether to remove the excess of reagent. Shake vigorously for 5–10 s by hand, then centrifugation at 2000*g* for 30 s (*see Note 3*). The ethereal extract can then be discarded.
- Saturate the aqueous layer with NaCl and extract twice with 3 mL of diethyl ether containing 10% of isopropanol with vigorous shaking for 5–10 s by hand. After centrifugation at 2000*g* for 30 s, the organic layers should be transferred into another tube by means of a Pasteur capillary pipet (*see Note 4*).
- The pooled ethereal extracts are methylated by bubbling diazomethane, generated according to the microscale procedure (50) (*see Note 5*). As shown in **Fig. 3**, a stream of nitrogen is saturated with diethyl ether in the first side-arm test tube and then passed through a diazomethane generating solution containing *N*-methyl-*N*-nitroso-*p*-toluenesulphonamide, diethylene-glycol monomethyl ether, and KOH. The generated diazomethane is carried into the sample tube via the nitrogen stream until a yellow tinge becomes visible. Leave to stand for more than 5

min at room temperature. The solvents are then removed by evaporation to dryness at 80°C under a stream of dry air. The residue on the walls of the tube are dissolved in 0.05–0.1 mL of ethyl acetate and then 1 µL of this solution is injected into the gas chromatograph (see **Note 6**).

### 3.2. Gas Chromatography

1. DB-1701 (14% cyanopropylphenyl-86% methylpolysiloxane, J & W, GC analysis was carried out with a Shimadzu 14A gas chromatograph equipped with a flame ionization detector and a flame photometric detector (P-filter).
2. For the analysis of free *O*-phosphoamino acids, fused-silica capillary columns (15 m × 0.53-mm id, 1.0-µm film thickness) of crosslinked DB-1701 or DB-210 should be used (see **Note 7**).
3. For a DB-1701 column, the operating conditions are as follows: column temperature program at 5°C/min from 180°C to 280°C; injection and detector temperature, 290°C.
4. For a DB-210 column, the operating conditions are as follows: column temperature program at 3°C/min from 200°C to 260°C; injection and detector temperature, 270°C.
5. For the analyzes of protein-bound P-Ser/P-Thr and P-Tyr, DB-1701 (15 m × 0.53-mm id, 1.0-µm film thickness) and DB-5 (15 m × 0.53-mm id, 1.0-µm film thickness) should be used (see **Notes 8 and 9**).
6. For the *O*-phosphoamino acid analysis of phosphorylated protein samples, a DB-1701 column is used with the following operating conditions: column temperatures, 210°C (for P-Ser/P-Thr) and 270°C (for P-Tyr); injection and detector temperatures, 260°C (P-Ser/P-Thr) and 290°C (for P-Tyr) (see **Note 10**).
7. For the P-Ser/P-Thr analyzes of protein-bound urine and tissue samples, a DB-1701 column is used with the following operating conditions: column temperature, isothermal at 220°C; injection and detector temperature, 260°C (see **Note 11**).
8. For the P-Tyr analyzes of protein-bound urine and tissue samples, a DB-5 column should be used with the following operating conditions: column temperature program at 3°C/min from 230°C to 280°C; injection and detector temperature, 290°C (see **Note 11**).
9. The nitrogen flow rate in each of these GC analyzes is 10 mL/min.
10. Measure the peak heights of each *O*-phosphoamino acid and the IS. Calculate the peak height ratios against the IS to construct calibration curves (see **Notes 12–14**).

### 3.3. Tissue Sample Preparation

#### 3.3.1. Preparation of Tissue Samples for the Free *O*-Phosphoamino Acid Analysis

Preparation of tissue samples for the free *O*-phosphoamino acid analysis (**43**): the following species were used in this experiment: cuttlefish (*Ommastrephes sloani pacificus*), mackerel (*Scomber Japonicus*), chicken, mouse, and pig (see **Note 15**).

1. After dissection of each animal, remove the organs and chill on ice. Chop the organs up into pieces and then store frozen at  $-20^{\circ}\text{C}$  until the assay.
2. Extract the *O*-phosphoamino acids from a tissue sample within 48 h by the following procedure, the assay being repeated three times.
3. Homogenize each tissue (0.2–1 g) with 5 vol of 5% of TCA using a Model LK-21 ultradisperser (Yamato Kagaku, Tokyo, Japan).
4. Centrifuge at 2000g for 10 min, then reextract the precipitate with 5 vol of 5% of TCA (see **Note 16**).
5. The combined supernatants are clarified by passage through a glass filter, if necessary, washed three times with two volumes of diethyl ether to remove TCA, and then warmed at  $60^{\circ}\text{C}$  to remove the ether.
6. After cooling, the solution is adjusted to pH 6.0–7.0 by the addition of a few drops of 1 M NaOH and then run through a Dowex  $1 \times 8$  ( $4 \times 0.7$ -cm id, AcO<sup>-</sup> form, 100–200 mesh) column (see **Note 17**).
7. Wash the column with 15 mL of water and then elute with 0.1 M HCl. The initial 9 mL of the eluate is discarded and the following 5 mL of the eluate (pH 1.0–2.0) is collected as the *O*-phosphoamino acid fraction. To this fraction, add 0.1 mL of 2.5  $\mu\text{g}/\text{mL}$  APB (IS) solution, and the resultant mixture is used for the derivatization.

### 3.3.2. Preparation of Tissue Samples for the Protein-Bound *O*-Phosphoamino Acid Analysis

Preparation of tissue samples for the protein-bound *O*-phosphoamino acid analysis (**46**): seven male ICR mice (6 wk old) were used in the experiments (see **Notes 18** and **19**).

1. Immediately after dissection, remove each organ and store at  $-20^{\circ}\text{C}$  until used. Chop up each pooled tissue. An aliquot (approx 0.5 g) should be homogenized with 5 vol of 5% TCA using an ultradisperser.
2. Centrifuge at 2000g for 10 min and reextract the precipitate with 5 vol of 5% TCA.
3. Wash the resulting pellet twice with 5 mL of diethyl ether to remove the TCA, then warm to dryness at  $60^{\circ}\text{C}$  (see **Note 20**).
4. Weigh the residue, then ground it to a powder, this being called the protein-bound fraction.
5. Hydrolyze an aliquot (5–10 mg) of the powdered protein-bound fraction with acid and base, as described in **Subheading 3.4**.
6. Adjust the pH of the acid and base hydrolyzates to pH 10.0–12.0 with 2 M NaOH or 10 M HCl, as described in **Subheading 3.4**.
7. The samples are then ready for derivatization (see **Notes 21** and **22**).

### 3.3.3. Preparation of Urine Samples for the Free *O*-Phosphoamino Acid Analysis

Preparation of urine samples for the free *O*-phosphoamino acid analysis (**48**).

1. Collect 24-h urine samples from healthy volunteers under toluene and keep frozen, if not analyzed immediately.

2. To 3 mL of urine samples, add 1 mL of 20% TCA and mix.
3. Centrifugation at 2000g for 10 min.
4. Wash the supernatant three times with 3 mL of diethyl ether to remove the TCA, and warm at 60°C to remove the ether.
5. After cooling, the solution should be adjusted to pH 6.0–7.0 by the addition of a few drops of 1 M NaOH and then run through a Dowex 1 × 8 (4.X 0.7-cm id, AcO<sup>-</sup> form, 200–400 mesh) column. Wash the column with 15 mL of water and then elute with 0.1 M HCl. Discard the initial 7.5 mL of the eluate. Collect the following 5 mL of the eluate (pH 1.0–2.0) as the *O*-phosphoamino acid fraction.
6. To this fraction, add 0.05 mL of 1 µg/mL APB and APO (IS) solutions.
7. The resultant mixtures are used for the derivatization.

### 3.3.4. Preparation of Urine Samples for the Total *O*-Phosphoamino Acid Analysis

Preparation of urine samples for the total (free plus protein-bound) *O*-phosphoamino acid analysis (48) (see **Notes 23** and **24**):

1. Collect, under toluene, 24-h urine samples from healthy volunteers and keep frozen, if not analyzed immediately.
2. Perform acid and base hydrolyses of urine samples, as described in **Subheading 3.4**.
3. For the P-Ser/P-Thr analysis, add 0.1 mL of 1 µg/mL APB (IS) to 0.1 mL of the urine sample and evaporated to dryness at 60°C in a centrifugal evaporator. Hydrolyzed the residue with 0.2 mL of 6 M HCl containing 1% phenol in the vapor phase for 2 h at 110°C under vacuum in a Pico-Tag workstation.
4. For the P-Tyr analysis, add 0.05 mL of 1 µg/mL APO (IS) and 1.22 mL of 10.5 M KOH to 3 mL of the urine sample and hydrolyze for 1 h at 130°C in a Pico-Tag workstation.
5. Adjust the resulting acid and base hydrolysates to pH 10.0–12.0 with 2 M NaOH or 10 M HCl, ready for derivatization.

## 3.4. Protein Sample Preparation

### 3.4.1 Acid Hydrolysis

For P-Ser and P-Thr analysis the proteins are hydrolyzed in an acid (see **Notes 25** and **26**).

1. Place an aliquot of the protein sample (0.004–2 mg) in a 5 × 50-mm glass test tube, add 50 µL of 20 µM APB (IS). Dry the mixture in a centrifugal evaporator.
2. Hydrolyze the residue with 0.2 mL of 6 M HCl containing 1% phenol in the vapor phase for 2 h at 110°C under vacuum in a Pico-Tag workstation (51).
3. Extract the resulting hydrolysate twice with 0.5 mL of distilled water, the extracts being transferred to another reaction tube (10-mL Pyrex glass tube with a PTFE-lined screw cap), and then used for derivatization.



### 3.4.2. Base Hydrolysis

For P-Tyr analysis, the proteins are hydrolyzed in a base (*see Note 27*).

1. Place an aliquot of the protein sample (0.2–4 mg) in a 9 × 75-mm polypropylene tube, add 50  $\mu$ L of 4  $\mu$ M APH (IS).
2. Add 0.2 mL of 10 M KOH, and make the total volume up to 0.4 mL with distilled water, if necessary.
3. Hydrolyze the mixture by heating for 1 h at 130°C under vacuum in a Pico-Tag workstation.
4. Add 0.2 mL of 8 M HCl to the resulting hydrolysate and transfer the mixture to another reaction tube. This is now ready for derivatization.

## 4. Notes

1. IsoBCF and diazomethane are used as a derivatizing reagents for the amino group and the carboxyl and phosphoryl groups, respectively (**Fig. 1**).
2. The *N*-isobutoxycarbonylation of *O*-phosphoamino acids with isoBCF proceeds rapidly and quantitatively in aqueous alkaline media (**41**). This reaction is completed with >50  $\mu$ L of isoBCF within 5 min by shaking at room temperature.
3. Although diethyl ether has been used as an extraction solvent for *N*-isoboc aminocarboxylic acids (**52,53**), *N*-isoboc phosphoamino acids cannot be extracted with this solvent. Therefore, the reaction mixture is washed with diethyl ether under acid condition in order to remove not only the excess reagent, but also the amines, phenols, and aminocarboxylic acids, which often coexist with *O*-phosphoamino acids in biological samples and derivatize to corresponding *N*- and *O*-isoboc derivatives under same conditions as described in **Note 2**.
4. In order to extract the remaining *N*-isoboc *O*-phosphoamino acids in aqueous layer, the addition of another solvent to diethyl ether was tested. Among the various solvents tested, isopropanol proved to be the most satisfactory solvent for this purpose, and its optimum concentration was found to be in the range 5–15%. Thus, the *N*-isoboc phosphoamino acids in aqueous layer could be quantitatively and selectively extracted into diethyl ether containing 10% isopropanol in acid condition with NaCl saturation. In this procedure, the organic layers should be collected, taking care to avoid aqueous droplets. It was not necessary to completely draw the organic layer in each extraction.
5. The methylation of the ethereal extracts is successfully carried out by bubbling diazomethane. This reaction should be performed in a well-ventilated hood because diazomethane is explosive and toxic.
6. The derivative preparation can be performed within 30 min, and several samples can be treated simultaneously. The *N*-isoboc methyl ester derivatives of *O*-phosphoamino acids are very stable to moisture; therefore, no precaution to exclude moisture is necessary in their handling and storage. No decomposition was observed during GC analysis and even after standing in ethyl acetate for 3 wk at room temperature.

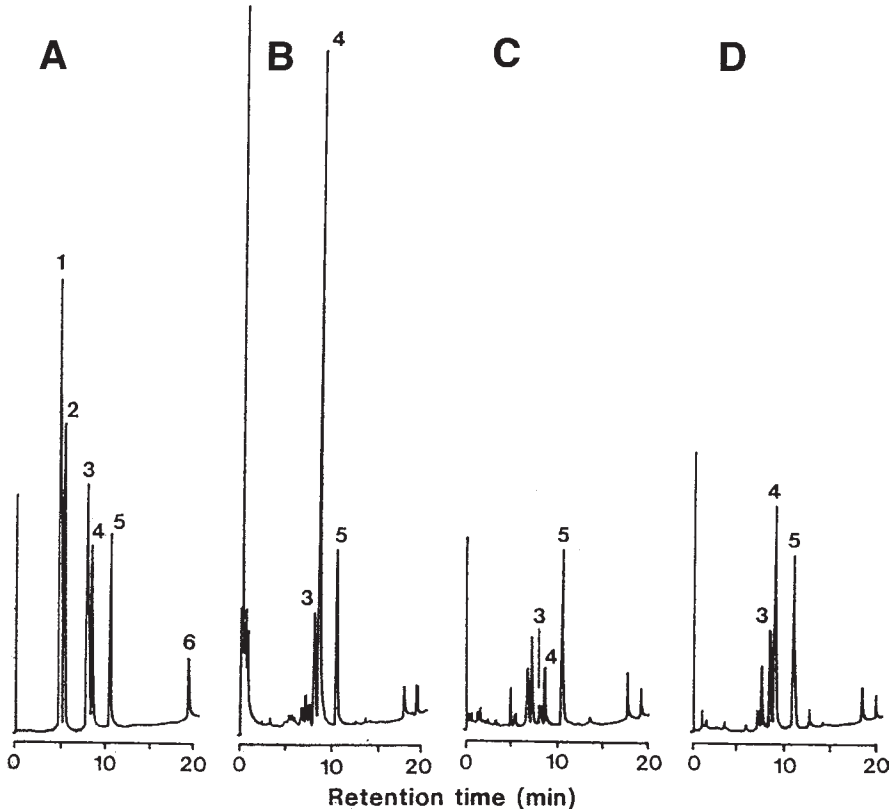


Fig. 4. Chromatograms obtained from standard and animal tissue samples for the free *O*-phosphoamino acid analysis by FPD-GC. (A) Standard (containing 0.25  $\mu\text{g}$  of each amino acid); (B) mackerel spleen (340 mg); (C) mouse brain (744 mg); (D) pig stomach (596 mg). GC conditions: column, DB-1701 (15 m  $\times$  0.53-mm id, 1.0- $\mu\text{m}$  film thickness); column temperature, programmed at 5 $^{\circ}\text{C}/\text{min}$  from 180 $^{\circ}\text{C}$  to 280 $^{\circ}\text{C}$ ; injection and detector temperatures, 290 $^{\circ}\text{C}$ ; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = 2-aminoethylphosphonic acid (AEP); 2 = *O*-phosphoethanolamine (PEA); 3 = *O*-phosphothreonine (P-Thr); 4 = *O*-phosphoserine (P-Ser); 5 = 2-amino-4-phosphonobutyric acid (APB) (IS); 6 = *O*-phosphotyrosine (P-Tyr).

7. The GC analysis was performed with FPD (526-nm interference filter), which is highly selective for phosphorus containing compounds, by using a megabore capillary column and direct (splitless) sample injection system. In this system, the sufficiently inactivated glass insert should be used to avoid tailing peaks of derivatives caused by the adsorption on injection port.
8. In preliminary test for several megabore capillary columns, DB-17, DB-1701, DB-210, and DB-5 gave good separation for *O*-phosphoamino acids.

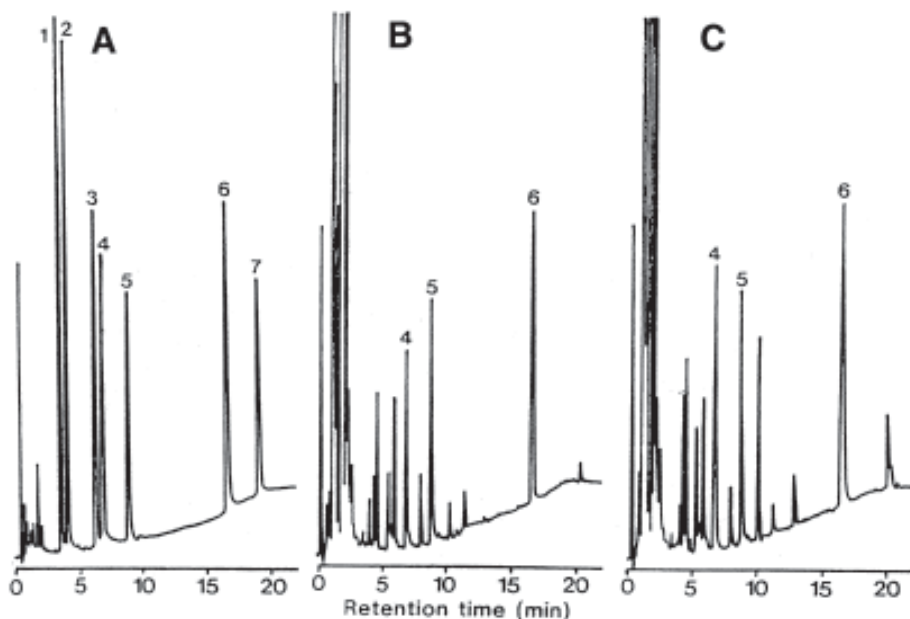


Fig. 5. Chromatograms obtained from standard and urine samples for the free *O*-phosphoamino acid analysis by FPD-GC. (A) Standard (containing 50 ng of each amino acid); (B) and (C) urine (3 mL) samples. GC conditions: column, DB-210 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, programmed at 3°C/min from 200°C to 260°C; injection and detector temperatures, 270°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = 2-aminoethylphosphonic acid (AEP); 2 = *O*-phosphoethanolamine (PEA); 3 = *O*-phosphothreonine (P-Thr); 4 = *O*-phosphoserine (P-Ser); 5 = 2-amino-4-phosphonobutyric acid (APB) (IS); 6 = 2-amino-8-phosphonooctanoic acid (APO) (IS); 7 = *O*-phosphotyrosine (P-Tyr) (from ref. 48 with permission).

9. For the free *O*-phosphoamino acid analysis, each amino acid gives a single and symmetrical peak from FPD-GC with a DB-1701, and are completely separated within 16 min from other biological phosphorus amino compounds, such as PEA and AEP (Fig. 4A) (43). Furthermore, the free *O*-phosphoamino acids can be also separated from these compounds using a DB-210 (Fig. 5A) (48). Other phosphate compounds, such as sugar phosphate and nucleic acid phosphate, and nonphosphorus amino acids were not detected by this method.
10. For the P-Ser/P-Thr and P-Tyr analyzes of phosphorylated protein samples, DB-1701 was used under different GC conditions, because these samples were separately treated by acid and base hydrolyzes. As shown in Figs. 6A and 7A, P-Ser/P-Thr and P-Tyr samples were separated within 12 and 8 min, respectively (44).

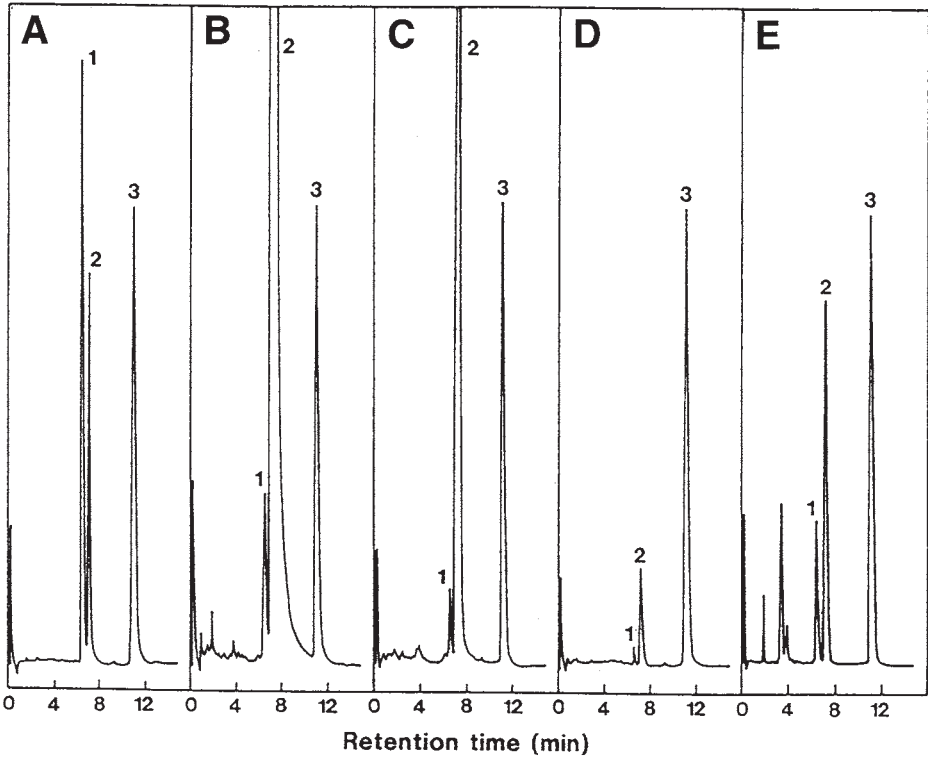


Fig. 6. Chromatograms obtained from a standard and acid hydrolysates of protein samples by FPD-GC. (A) Standard (containing 1 nmol of each compound); (B)  $\alpha$ -casein (0.5 mg); (C) phosvitin (0.01 mg); (D) histone (0.8 mg); (E) myelin (0.25 mg). GC conditions: column, DB-1701 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, 210°C; injection and detector temperatures, 260°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = *O*-phosphothreonine (P-Thr); 2 = *O*-phosphoserine (P-Ser); 3 = 2-amino-4-phosphonobutyric acid (APB) (IS) (from ref. 44 with permission).

11. DB-1701 and DB-5 proved to be the most satisfactory columns for the P-Ser/P-Thr and P-Tyr analyzes of urine hydrolysates, respectively. By using these columns, each *O*-phosphoamino acid elutes separately as single and symmetrical peaks within 10 min (Figs. 8A and 9A) (48).
12. The calibration curves for the free *O*-phosphoamino acids by FPD-GC were conducted using APB, which showed a similar behavior to other *O*-phosphoamino acids during the derivatization and was well separated from these amino acids on a chromatogram as the IS. A linear relationship was obtained at the range 0.01–2 mg with the regression lines for P-Ser, P-Thr and P-Tyr being  $y = 3.938x - 0.040$  ( $r = 0.9984$ ,  $n = 18$ ),  $y = 5.183x - 0.028$  ( $r = 0.9983$ ,  $n = 18$ ), and  $y = 1.761x -$

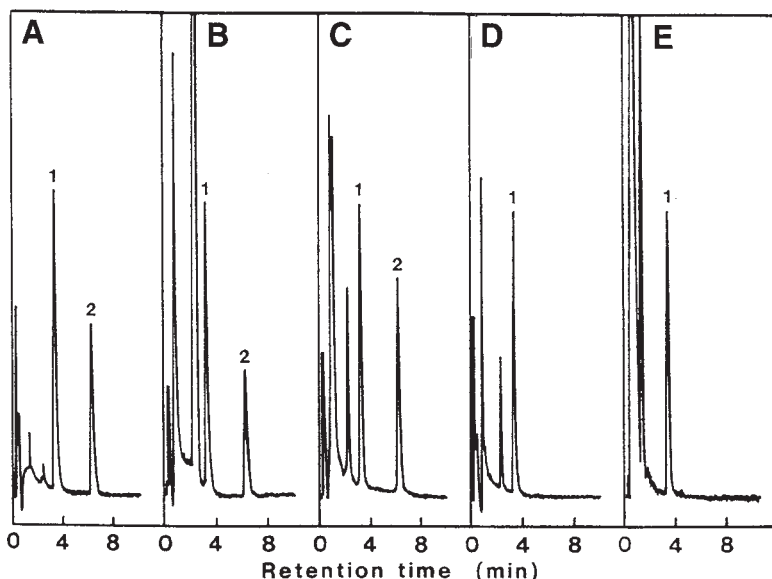


Fig. 7. Chromatograms obtained from a standard and base hydrolysates of protein samples by FPD-GC. (A) Standard (containing 0.2 nmol of each compound); (B)  $\alpha$ -casein (2.0 mg); (C) phosvitin (1.3 mg); (D) histone (2.7 mg); (E) myelin (1.0 mg). GC conditions: column, DB-1701 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, 270°C; injection and detector temperatures, 290°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = 2-amino-7-phosphonoheptanoic acid (APH) (IS); 2 = *O*-phosphotyrosine (P-Tyr) (from ref. 42 with permission).

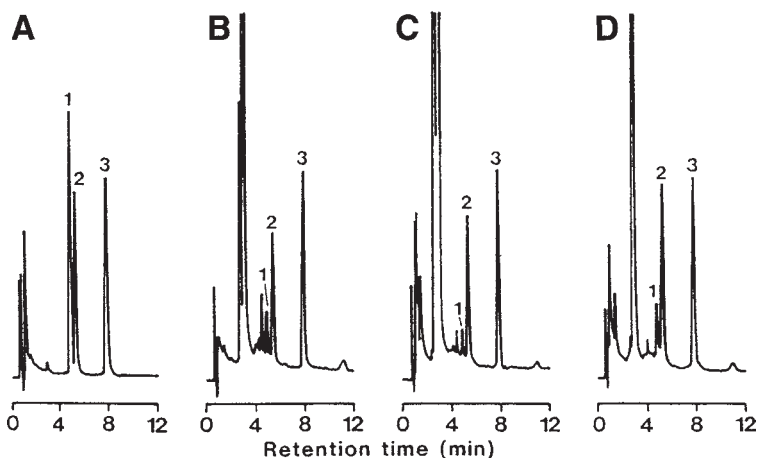


Fig. 8. Chromatograms obtained from a standard and acid hydrolysates of urine samples by FPD-GC. ; Standard (containing 100 ng of each compound); (B)–(D) urine (0.1 mL) hydrolysates. GC conditions: column, DB-1701 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, 220°C; injection and detector temperatures, 260°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = *O*-phosphothreonine (P-Thr); 2 = *O*-phosphoserine (P-Ser); 3 = 2-amino-4-phosphonobutyric acid (APB) (IS) (from ref. 48 with permission).

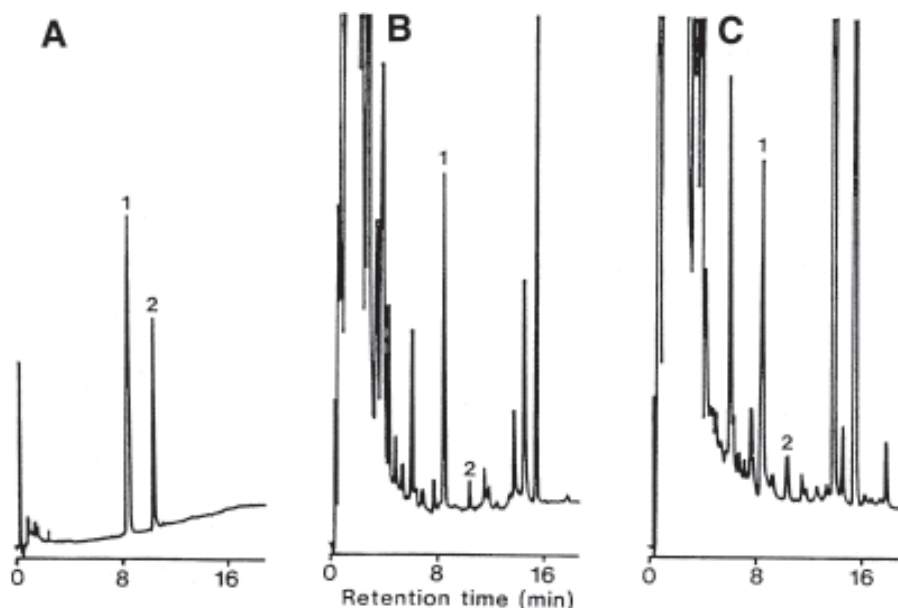


Fig. 9. Chromatograms obtained from a standard and base hydrolysates of urine samples by FPD-GC. (A) Standard (containing 50 ng of each compound); (B) and (C) urine (3 mL) hydrolysates. GC conditions: column, DB-5 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, programmed at 3°C/min from 230°C to 280°C; injection and detector temperatures, 290°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = 2-amino-8-phosphonooctanoic acid (APO) (IS); 2 = *O*-phosphotyrosine (P-Tyr) (from ref. 48 with permission).

0.054 ( $r = 0.9972$ ,  $n = 18$ ), respectively, where  $y$  is the peak height ratio and  $x$  is the amount ( $\mu$ g) of each compound.

13. The derivatives provided excellent FPD responses and the minimum detectable amounts of P-Ser, P-Thr, and P-Tyr at a signal three times as high as the noise under our instrumental conditions were approx 50, 40, and 200 pg as injection amounts, respectively. The FPD-GC system described here was over 200 times more sensitive than the FID-GC system.
14. The calibration curves for the analysis of *O*-phosphoamino acids in protein hydrolysates were conducted using APB (for P-Ser/P-Thr analysis) and APH or APO (for P-Tyr analysis) as the IS (44,48). In each case, a linear relationship was obtained using different GC columns in the range 10–500 ng of each *O*-phosphoamino acid and the correlation coefficients were above 0.994 (48). The detection limits for P-Ser, P-Thr, and P-Tyr in these GC conditions were approx 40, 30, and 80 pg as injection amounts, respectively.
15. The method developed for free *O*-phosphoamino acids was successfully applied to animal tissue (43) and urine (48) samples. In preliminary experiments on mice,

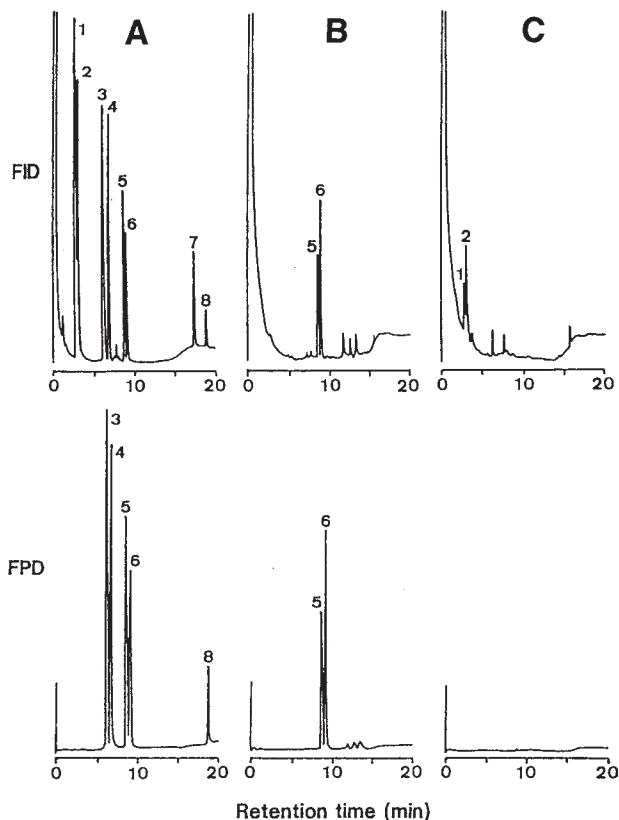


Fig. 10. Chromatograms obtained from (A) a standard solution, and (B) nonhydrolyzed and (C) hydrolyzed samples of porcine liver. The converted samples were analyzed by both FID-GC (top) and FPD-GC (bottom). GC conditions: column, DB-1701 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, isothermal at 180°C for 6 min, programmed at 20°C/min to 220°C, isothermal at 220°C for 6 min, programmed at 20°C/min to 270°C, and isothermal at 270°C for 5 min; injection and detector temperatures, 280°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = threonine (Thr); 2 = serine (Ser); 3 = 2-aminoethylphosphonic acid (AEP); 4 = *O*-phosphoethanolamine (PEA); 5 = *O*-phosphothreonine (P-Thr); 6 = *O*-phosphoserine (P-Ser); 7 = tyrosine (Tyr); 8 = *O*-phosphotyrosine (P-Tyr) (from **ref. 42** with permission).

no postmortem change of *O*-phosphoamino acids was observed after 24 h. *O*-phosphoamino acids in various organs were stable for at least 2 d under freezing at -20°C.

16. TCA is routinely used to precipitate proteins with no loss of phosphoryl groups (54). Free *O*-phosphoamino acids in tissue and urine samples were quantitatively extracted into the TCA soluble fraction.

**Table 1**  
**Recoveries of O-Phosphoamino Acids Added to Tissue and Urine Samples**

| Sample                         | O-Phospho-<br>amino acid | Added <sup>a</sup> | Amount found <sup>b</sup> |            | Recovery<br>(%) |
|--------------------------------|--------------------------|--------------------|---------------------------|------------|-----------------|
|                                |                          |                    | Nonaddition               | Addition   |                 |
| Pig tissue                     |                          |                    |                           |            |                 |
| Brain                          | P-Ser                    | 500                | 330 ± 20                  | 850 ± 14   | 104.0           |
|                                | P-Thr                    | 100                | 57 ± 2                    | 152 ± 3    | 95.0            |
|                                | P-Tyr                    | 100                | ND <sup>c</sup>           | 103 ± 4    | 103.0           |
| Heart                          | P-Ser                    | 500                | 161 ± 11                  | 671 ± 50   | 102.0           |
|                                | P-Thr                    | 100                | 18 ± 1                    | 117 ± 3    | 99.0            |
|                                | P-Tyr                    | 100                | ND                        | 97 ± 2     | 97.0            |
| Liver                          | P-Ser                    | 500                | 377 ± 13                  | 878 ± 64   | 100.2           |
|                                | P-Thr                    | 100                | 115 ± 2                   | 214 ± 4    | 99.0            |
|                                | P-Tyr                    | 100                | ND                        | 102 ± 3    | 102.0           |
| Kidney                         | P-Ser                    | 500                | 567 ± 27                  | 1085 ± 57  | 103.6           |
|                                | P-Thr                    | 100                | 46 ± 3                    | 146 ± 8    | 100.0           |
|                                | P-Tyr                    | 100                | ND                        | 103 ± 5    | 103.0           |
| Urine                          |                          |                    |                           |            |                 |
| A                              | P-Ser                    | 20                 | 26.8 ± 0.7                | 46.4 ± 2.1 | 98.0            |
|                                | P-Thr                    | 20                 | ND                        | 19.6 ± 0.6 | 98.0            |
|                                | P-Tyr                    | 20                 | ND                        | 19.0 ± 1.1 | 95.0            |
| B                              | P-Ser                    | 20                 | 12.3 ± 0.5                | 31.8 ± 0.9 | 97.5            |
|                                | P-Thr                    | 20                 | ND                        | 18.4 ± 0.7 | 92.0            |
|                                | P-Tyr                    | 20                 | ND                        | 18.7 ± 1.5 | 93.5            |
| Urine hydrolysate <sup>d</sup> |                          |                    |                           |            |                 |
| A                              | P-Ser                    | 500                | 1017 ± 42                 | 1504 ± 54  | 97.4            |
|                                | P-Thr                    | 500                | 133 ± 9                   | 593 ± 35   | 92.0            |
|                                | P-Tyr                    | 50                 | 2.4 ± 0.1                 | 47.7 ± 0.7 | 90.6            |
| B                              | P-Ser                    | 500                | 1636 ± 60                 | 2111 ± 118 | 95.0            |
|                                | P-Thr                    | 500                | 238 ± 7                   | 694 ± 25   | 91.2            |
|                                | P-Tyr                    | 50                 | 6.2 ± 0.4                 | 53.5 ± 1.7 | 94.6            |

<sup>a</sup>Tissue sample, ng/g; urine sample, ng/mL.

<sup>b</sup>Tissue sample, ng/g; urine sample, ng/mL; mean ± SD (n=3).

<sup>c</sup>Not detectable.

<sup>d</sup>P-Ser and P-Thr were added to acid hydrolysate and P-Tyr was added to base hydrolysate.

17. The TCA extract was applied on a Dowex 1 column in order to remove any coexisting substances. O-phosphoamino acids were adsorbed on the column and were eluted in 0.1 M HCl fraction. This column chromatographic procedure could be completed within 50 min, and the recoveries of P-Ser, P-Thr, and P-Tyr were 97–103%. **Figs. 4B–D (43)** and **5B,C (48)** show the chromatograms obtained from



**Table 2**  
***O*-Phosphoamino Acid Contents in Several Commercial Proteins**

| <i>O</i> -Phosphoamino acid (nmol/mg of protein) <sup>a</sup> |                    |                    |                    |
|---------------------------------------------------------------|--------------------|--------------------|--------------------|
| Protein                                                       | P-Ser <sup>b</sup> | P-Thr <sup>b</sup> | P-Tyr <sup>c</sup> |
| Albumin                                                       | ND <sup>d</sup>    | ND                 | ND                 |
| Ovalbumin                                                     | 7.74 ± 0.51        | ND                 | ND                 |
| α-Casein                                                      | 137.30 ± 14.20     | 0.69 ± 0.04        | 0.073 ± 0.004      |
| β-Casein                                                      | 104.28 ± 4.87      | 0.16 ± 0.01        | 0.013 ± 0.001      |
| κ-Casein                                                      | 67.26 ± 0.89       | 0.87 ± 0.01        | 0.071 ± 0.002      |
| Phosvitin                                                     | 848.14 ± 35.81     | 11.06 ± 0.75       | 0.182 ± 0.010      |
| Hemoglobin                                                    | 0.03 ± 0.002       | ND                 | ND                 |
| Cytochrome <i>c</i>                                           | 0.08 ± 0.002       | ND                 | ND                 |
| Protamine                                                     | 1.46 ± 0.07        | ND                 | ND                 |
| Histone                                                       | 0.30 ± 0.01        | 0.04 ± 0.002       | ND                 |
| Myelin basic protein                                          | 3.93 ± 0.28        | 1.00 ± 0.06        | ND                 |
| Alcohol dehydrogenase                                         | 0.07 ± 0.004       | ND                 | ND                 |
| Catalase                                                      | 0.17 ± 0.01        | 0.04 ± 0.001       | ND                 |
| Lysozyme                                                      | 0.05 ± 0.003       | ND                 | ND                 |
| Chymotrypsin                                                  | 0.13 ± 0.01        | ND                 | ND                 |

<sup>a</sup>Mean ± SD (*n* = 3).

<sup>b</sup>Analyzed after partial acid hydrolysis.

<sup>c</sup>Analyzed after partial base hydrolysis.

<sup>d</sup>Not detectable.

(From **ref. 44** with permission.)

animal tissue and urine samples, respectively. *O*-phosphoamino acids could be detected without any interference from coexisting substances by FPD-GC.

18. P-Ser and P-Thr were found in various animal tissues, but only P-Ser was found in urine samples. The occurrence of free P-Ser and P-Thr in these samples was confirmed by GC, GC-mass spectrometry (GC-MS), and thin-layer chromatography of nonhydrolyzed and hydrolyzed samples (42). As shown in **Fig. 10**, peaks corresponding to P-Ser and P-Thr were observed in the chromatograms obtained by FID-GC (top), and FPD-GC (bottom) analyzes of the porcine liver sample. But these peaks disappeared after acid hydrolysis of sample with 6 *M* HCl at 110°C for 48 h, and the peaks corresponding to Ser and Thr newly appeared instead (**Fig. 10**). As shown in **Table 1**, the recoveries of P-Ser, P-Thr, and P-Tyr added to tissue and urine samples were 92–104%, the relative standard deviations being 1.6–8.0% (*n* = 3).
19. The method developed for protein-bound *O*-phosphoamino acids was successfully applied to purified proteins (44), phosphorylated proteins by protein kinases (45), animal tissue (46) and urine (48) samples. For the presence of *O*-phosphoamino acid residues in protein samples, several commercial proteins were analyzed by FPD-GC. As shown in **Figs. 6B–D** and **7B–D**, *O*-phosphoamino

acids in protein hydrolysates were selectively detected without any interference from other amino acids (44). This method is reproducible and directly applicable for the analysis of the extent of both in vivo and in vitro phosphorylation without radiolabeling.

20. The proteins containing *O*-phosphoamino acids were recovered by better than 99% in the pellet with the TCA treatment without loss of phosphoryl groups (54). Phospholipids, such as phosphatidylcholine and phosphatidylserine, were removed from the protein-bound fraction by washing the pellet with ether.
21. The results of *O*-phosphoamino acid analyzes of the protein samples after partial acid and base hydrolyzes are shown in **Table 2**. P-Ser was detected in all the samples investigated in this study, except for BSA, and P-Thr and P-Tyr were also detected in casein and phosvitin. This method was successfully applied to *O*-phosphoamino acid analysis of the proteins phosphorylated by protein kinases (45).
22. Although the recovery of *O*-phosphoamino acids from the protein-bound fraction under the best conditions for acid hydrolysis was low (approx 35–40%) (43), the method could reproducibly measure endogenously phosphorylated amino acids in the tissue and urine samples without radiolabeling.
23. As shown in **Table 1**, the recoveries of *O*-phosphoamino acids added to urine hydrolysates were 91–97%, the relative standard deviations being 1.5–6.8% ( $n = 3$ ).
24. **Figs. 8B–D** and **9B,C** show the chromatograms obtained from acid and base hydrolysates of urine samples by FPD-GC. The P-Ser, P-Thr, and P-Tyr peaks obtained from urine hydrolysates were confirmed by GC-MS analysis (47). By using this method, we demonstrated that P-Tyr levels in mouse urine and liver increase during liver regeneration after partial hepatectomy (49).
25. The *O*-phosphate linkages of Ser and Thr residues are defined by their stability to acids and lability to bases. Thus, acid hydrolysis has been routinely used to study the identity and amounts of P-Ser and P-Thr in peptides and proteins. For the acid hydrolysis of phosvitin (containing P-Ser, P-Thr, and P-Tyr residues) with the HCl vapor phase using a Pico-Tag workstation, the maximal recoveries of P-Ser and P-Thr were obtained at 110°C for 2–3 h (**Fig. 11**), these conditions being in agreement with those reported previously (20). Under these conditions, P-Ser was detected as over 70 times more than P-Thr in phosvitin, whereas P-Tyr was not detected at all.
26. Under the best conditions for acid hydrolysis, the recovery of P-Ser from phosvitin was calculated to be  $35.7 \pm 1.5\%$  ( $n = 3$ ) by measuring the total phosphate content. The recovery of P-Ser from  $\alpha$ -casein was  $39.5 \pm 4.1\%$  ( $n = 3$ ) under the same conditions. These low recoveries seem to have been caused by both incomplete peptide hydrolysis and destruction of the liberated P-Ser to Ser. The typical recovery for proteins has been reported to be in the range 20–30% (19). Our results presented here are generally somewhat higher, the recoveries may show some dependence on the neighboring amino acid residues (55).
27. The phosphate linkage of the Tyr residue is far more resistant to base than to acid hydrolysis. P-Tyr tolerated the base hydrolysis fairly well in 5 *M* KOH at 155°C

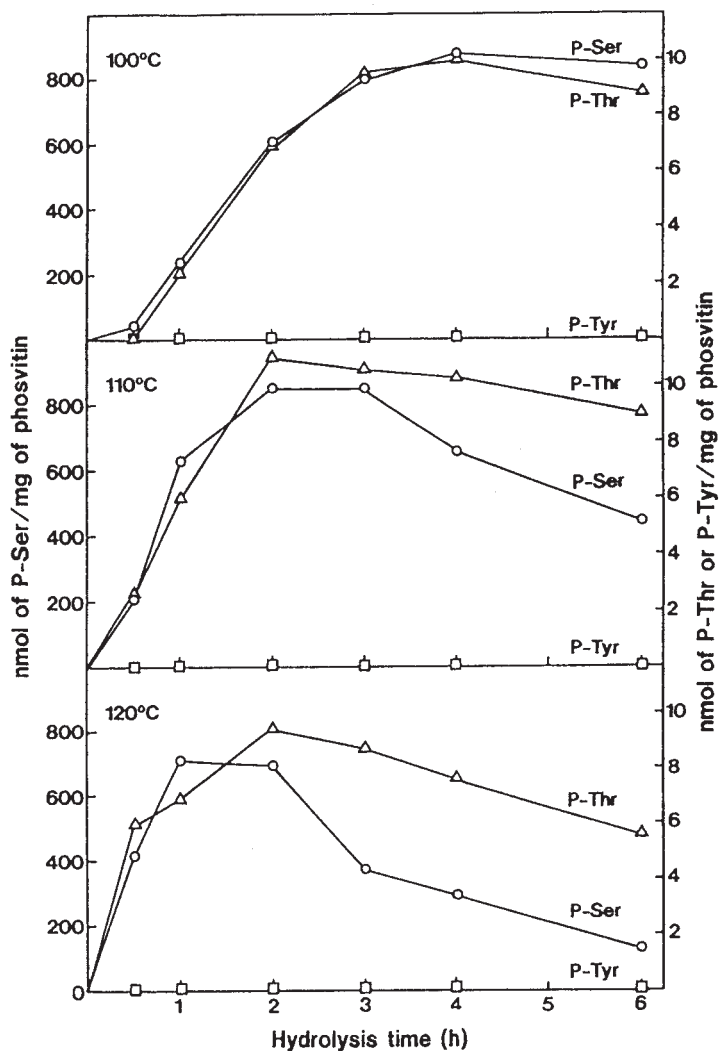


Fig. 11. Release of *O*-phosphoamino acids from phosvitin by acid hydrolysis.

for 30–35 min (17,54). In our experiment, the maximal recovery of P-Tyr from phosvitin was obtained in 5 M KOH at 130°C for 0.5–4 h (Fig. 12).

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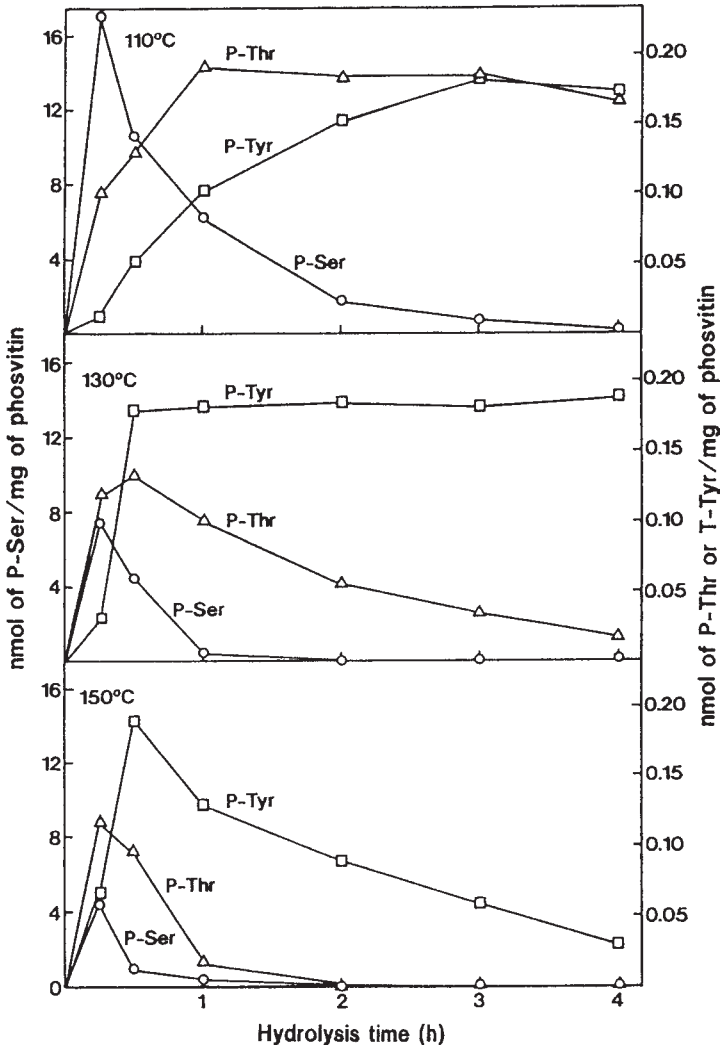


Fig. 12. Release of *O*-phosphoamino acids from phosvitin by base hydrolysis.

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## Determination of Sulfur Amino Acids, Glutathione, and Related Amino Thiols in Biological Samples by Gas Chromatography with Flame Photometric Detection

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### 1. Introduction

The determination of sulfur amino acids, glutathione (GSH), and related amino thiols has been carried out by isotachopheresis (*1*), amino acid analyzer (AAA) (*2,3*), gas chromatography (GC) (*4–8*), high-performance liquid chromatography (HPLC) (*9–32*), GC-mass spectrometry (GC-MS) (*33,34*), liquid chromatography-mass spectrometry (LC-MS) (*35*), and capillary zone electrophoresis (CZE) (*36–38*). However, isotachopheresis and AAA methods are nonselective for sulfur amino acids and lack sensitivity. GC methods based on the conversion into trimethylsilyl (*4,5*), neopentylidene (*6*), and N-trifluoroacetyl *n*-butyl ester (*7*) derivatives lack sensitivity, gives a tailing peak and requires anhydrous derivatization conditions. Although GC method based on the preparation of N-heptafluorobutyryl isobutyl or ethyl esters (*8*) is selective and sensitive by flame photometric detection (FPD), this method is not applied to the analysis of biological samples. As the HPLC methods, the ultraviolet (UV) (*9–12*), and the postcolumn UV derivatization with 4,4'-dithiopyridine (*13*) and 5,5'-dithiobis (2-nitrobenzoic acid) (*14*), the precolumn fluorescence derivatization with 4-(aminosulfonyl)- or ammonium-7-fluoro-2,1,3-benzoxadiazole-4-sulfonate (*15,16*), methyl 4-(6-methoxynaphthalene-2-yl)-4-oxo-2-butenate (*17*), N-(1-pyrenyl)maleimide (*18*), monobromobimane (*19–23*), *o*-phthaldialdehyde (*24,25*), 9-fluorenylmethyloxycarbonyl chloride (*26*), and 2-chloro-1-methylpyridinium (*27*), and electrochemi-

cal detection (28–32) have been reported. Many of these HPLC methods were highly sensitive, but some of these methods lack specificity and require clean-up of the sample to remove the excess reagent and coexisting substances. GC-MS methods based on the conversion into *tert*-butyldimethylsilyl (33) and *N*(O,S)-propoxycarbonyl propyl ester (34) derivatives, and LC-MS method were highly sensitive and specific, but these methods require expensive equipment. Furthermore, CZE methods with electrochemical detection are capable of achieving higher separation efficiency, use less organic solvents, and require small amounts of samples in comparison with HPLC, but these methods are not applied enough to the analysis of biological samples. Current methods for the determination of sulfur amino acids, GSH, and related aminothiols have also been described in detail in refs. 39–43.

Recently, we have developed a selective and sensitive method for the determination of sulfur amino acids by FPD-GC using a DB-17 capillary column (15 m × 0.53-mm id), in which these compounds were analyzed as their *N*(S)-isopropoxycarbonyl (isoPOC) methyl ester derivatives (44). By using this method, we demonstrated that the contents of these amino acids in urine (44) and plasma (45) samples could be rapidly and simply determined. Furthermore, we have developed a selective and sensitive method for the determination of GSH and related aminothiols such as cysteine (Cys), cysteinylglycine (CysGly), and  $\gamma$ -glutamylcysteine ( $\gamma$ -GluCys) by FPD-GC, using a short capillary column (5 m × 0.53-mm id) of crosslinked DB-1 (46). By using this method, we demonstrated that the contents of these aminothiols in blood (46) and tissue (47) samples could be rapidly and simply determined. In the aforementioned methods, other sulfur containing amino acids such as taurine and related aminosulfonic acids could not be analyzed, but these compounds could be analyzed as their *N*-isoBOC di-*n*-butylamide derivatives by GC (48–52). In this chapter, selective and sensitive method for the determination of sulfur amino acids, GSH, and related aminothiols, except for aminosulfonic acids by FPD-GC, is described on the basis of the above results, and optimum conditions and typical problems encountered in the development and application of the method are discussed.

The derivatization process of sulfur amino acids, GSH, and related aminothiols and structures of main compounds are shown in Fig. 1. *N*(S)-isoPOC methyl esters of these compounds can be easily, rapidly, and quantitatively prepared by the reaction with isopropyl chloroformate (isoPCF) by shaking in aqueous alkaline media, followed by esterification with hydrogen chloride-methanol (HCl-MeOH). The oxidized and protein-bound forms of these compounds were reduced to free thiols by adding sodium borohydride (NaBH<sub>4</sub>) prior to derivatization. The main advantage of this method is that these amino acids can be easily converted in an aqueous medium without any

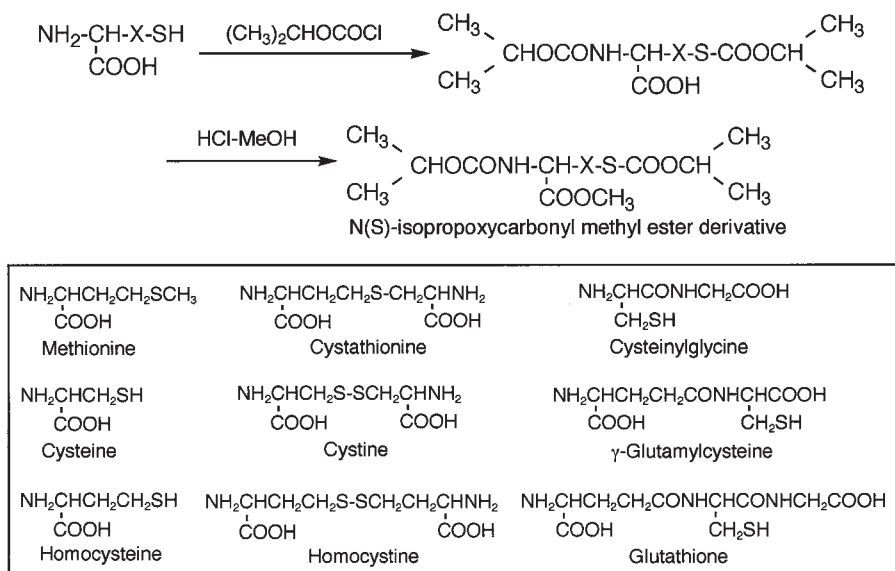


Fig. 1. Derivatization process and structures of sulfur amino acids, glutathione, and related aminothiols.

further clean-up procedure and the derivatives are stable to moisture. Furthermore, the derivatives can be quantitatively and reproducibly resolved as single peaks using a capillary column, and provide an excellent FPD response.

## 2. Materials

### 2.1. Equipment

1. Shimadzu 12A gas chromatograph equipped with a hydrogen flame ionization detector and a flame photometric detector (S-filter).
2. For the analysis of sulfur amino acids, a fused-silica capillary column (15 m  $\times$  0.53-mm id, 1.0- $\mu\text{m}$  film thickness) of crosslinked DB-17 (50% phenyl-50% methylpolysiloxane, J & W, Folsom, CA).
3. For the analysis of GSH and related aminothiols, a fused-silica capillary column (5 m  $\times$  0.53-mm id, 1.5- $\mu\text{m}$  film thickness) of crosslinked DB-1 (methylpolysiloxane, J & W).
4. Model LK-21 ultradisperser (Yamato Kagaku, Tokyo, Japan).
5. Pasteur capillary pipet (Iwaki glass no. IK-PAS-5P).

### 2.2. Reagents

1. Amino acids and peptides: L-methionine (Met), ethionine (Eth), L-thioprolin (TPro), L-cysteine (Cys), D,L-homocysteine (HCys), L-cystine (Cyt), D,L-ho-

mocystine (HCyt), and djenkolic acid (Dje) were purchased from Nacalai Tesque (Kyoto, Japan). L-Cystathionine (CTH), L-methionine sulfone (Met-S), S-methyl-L-cysteine (SM-Cys), S-carboxymethyl L-cysteine (SC-Cys) and cysteinylglycine (CysGly) were purchased from Sigma (St. Louis, MO). Glutathione (GSH) and  $\gamma$ -glutamylcysteine ( $\gamma$ -GluCys) were purchased from Kohjin Co., Ltd. (Tokyo, Japan). Glutathione disulphide (GSSG) was purchased from Wako Pure Chemical Industries (Osaka, Japan). All of the nonsulfur protein amino acids used were purchased from Ajinomoto (Tokyo, Japan).

2. Internal standards: S-2-Aminoethyl-L-cysteine (AE-Cys) and lanthionine (LTH) as internal standard (IS) were purchased from Sigma.
3. Derivatizing reagents: isopropyl chloroformate (isoPCF) was obtained from Wako Pure Chemicals Industries (Osaka, Japan). Hydrogen chloride in methanol (HCl-MeOH) obtained from Tokyo Kasei Kogyo (Tokyo, Japan) was diluted with methanol at a concentration of 1 M.
4. Other materials: Sodium borohydride ( $\text{NaBH}_4$ ) and dithioerythritol (DTE) were obtained from Nacalai Tesque. Peroxide-free diethyl ether was obtained from Dojindo Laboratories (Kumamoto, Japan). Distilled water was used after freshly purification with a Model Milli-Q Jr. water purifier (Millipore, Bedford, MA). All other chemicals were of analytical-reagent grade.

### 2.3. Solutions

1. Standard sulfur amino acid solutions: three standard stock solutions (each 2 mM), one containing the 13 sulfur amino acids except for GSH, CysGly, and  $\gamma$ -GluCys, the second containing Cys, CysGly, and  $\gamma$ -GluCys, and the third containing nonsulfur protein amino acids; are prepared in 0.05 M HCl.
2. Working standard solutions: each working standard solution is made up freshly as required by dilution of the stock solution with 0.01 M HCl. When these standard solutions are stored at 4°C, they are stable for at least 2 wk.
3. The GSH standard solution: GSH is freshly dissolved in distilled water at the required concentration on each day, because GSH is partially hydrolyzed to  $\gamma$ -GluCys during storage.
4. The internal standard solutions: AE-Cys and LTH are dissolved in 0.05 M HCl to make a stock solution at a concentration of 2 mM. Store at 4°C.
5.  $\text{NaBH}_4$  solution:  $\text{NaBH}_4$  is dissolved in 0.1 M NaOH at a concentration of 100 mg/mL solution. This solution is stable at 4°C for at least 1 wk.
6. DTE solution: DTE is dissolved in distilled water at a concentration of 0.5 mM.

## 3. Methods

### 3.1. Derivatization

Derivatization procedure of sulfur amino acids, GSH and related aminothiols is shown in Fig. 2 (see Note 1).

1. For the analysis of 13 sulfur amino acids, pipet an aliquot of the sample containing 0.5–10 nmol of each amino acid and 0.1 mL of 2  $\mu$ M AE-Cys (IS) (if necessary) into a 10-mL Pyrex glass reaction tube with a PTFE-lined screw cap.

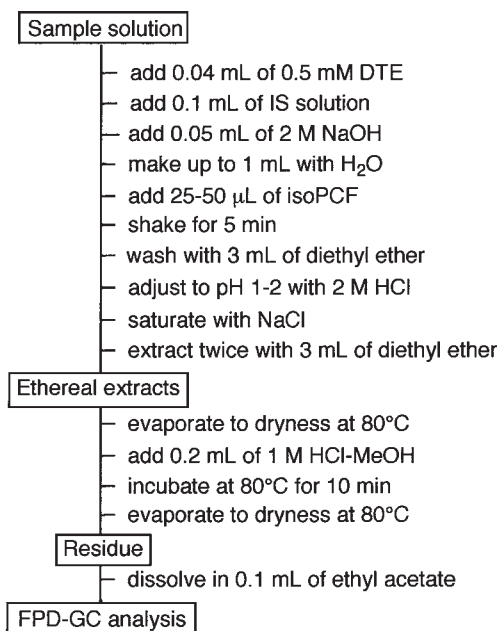


Fig. 2. Schematic flow diagram of the derivatization procedure.

2. For the analysis of GSH and related aminothiols, pipet an aliquot of the sample containing 0.2–25 nmol of each compounds and 0.1 mL of 10  $\mu$ M LTH (IS) (if necessary) into a reaction tube.
3. To these solutions, add 40  $\mu$ L of 0.5 mM DTE (*see Note 2*) and 0.05 mL of 2 M NaOH, and make the total volume up to 1 mL with distilled water.
4. Immediately add 25–50  $\mu$ L of isoPCF and shake the mixture at 300 cycles per minute (up and down) for 5 min at room temperature (*see Note 3*).
5. Extract the reaction mixture with 3 mL of peroxide-free diethyl ether (*see Note 2*), to remove the excess of reagent, with vigorous shaking for 5–10 s by hand (*see Note 4*).
6. Centrifuge at 2000g for 30 s then discard the ethereal extract.
7. Acidify the aqueous layer to pH 1–2 with 2 M HCl and saturate with NaCl.
8. Extract the mixture twice with 3 mL of peroxide-free diethyl ether with vigorous shaking for 5–10 s by hand (*see Note 5*).
9. Repeat centrifugation at 2000g for 30 s, and transfer the ether layers into another tube by means of a Pasteur capillary pipet.
10. Evaporate the ethereal extracts to dryness at 80°C, and to the residue add 0.2 mL of 1 M HCl-MeOH. Incubate the mixture at 80°C for 10 min (*see Note 6*).
11. Evaporate the residual solvent to dryness at 80°C under a stream of dry air. Re-dissolve the residue on the walls of the tube in 0.1–0.2 mL of ethyl acetate and inject 0.2–1  $\mu$ L of this solution into the gas chromatograph (*see Note 7*).

### 3.2. Gas Chromatography

1. GC analysis was carried out with a Shimadzu 12A gas chromatograph equipped with a flame ionization detector and a flame photometric detector (S-filter) (*see Note 8*).
2. For the analysis of sulfur amino acids, a fused-silica capillary column (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness) of crosslinked DB-17 is used. The operating conditions are as follows: column temperature program at 5°C/min from 130°C to 270°C or at 10°C/min from 150°C to 270°C; injection and detector temperature, 280°C; nitrogen flow rate, 10 mL/min (*see Note 9*).
3. For the analysis of GSH and related aminothiols, a fused-silica capillary column (5 m  $\times$  0.53-mm id, 1.5- $\mu$ m film thickness) of crosslinked DB-1 is used. The operating conditions are as follows: column temperature program at 16°C/min from 130°C to 290°C; injection and detector temperature, 300°C; nitrogen flow rate, 12 mL/min (*see Notes 10 and 11*).
4. Measure the peak heights of each compound and the IS. Calculate the peak height ratios against the IS to construct calibration curves (*see Notes 12–15*).

### 3.3. Sample Preparation (*see Note 16*)

#### 3.3.1. Preparation of Urine Samples (44)

1. Collect early morning urine samples from healthy volunteers and processed immediately or store at  $-20^{\circ}\text{C}$  until used.
2. Urine samples (0.02–0.1 mL) are used directly for derivatization and FPD-GC analysis (*see Note 17*).

#### 3.3.2. Preparation of Blood Samples (46)

1. Collect venous blood samples from healthy volunteers in 9 vol of 1 mM ethylenediamine tetraacetate (EDTA) and mix (*see Note 18*).
2. Immediately process the 10% (v/v) clear hemolysates or store at  $-20^{\circ}\text{C}$  until used.
3. To determine the free GSH, Cys, CysGly, and  $\gamma$ -GluCys contents, 0.1 mL of 10% hemolysate is directly used as the sample for derivatization.
4. To obtain total (free and bound, reduced, and oxidized) GSH, Cys, CysGly, and  $\gamma$ -GluCys, chemically reduce a blood sample with  $\text{NaBH}_4$  (*see Note 19*). To 0.05 mL of 10% hemolysate add 0.1 mL of 10  $\mu\text{M}$  LTH (IS) and 0.2 mL of 100 mg/mL  $\text{NaBH}_4$ , and then make the total volume up to 1 mL with distilled water. After adding 1 drop of *n*-hexanol (*see Note 20*), the mixture is incubated at  $100^{\circ}\text{C}$  for 10 min (*see Note 21*). After cooling, the reaction mixture is used as the sample for derivatization.

#### 3.3.3. Preparation of Tissue Samples (47)

The outline of the method for determining the different forms of GSH and related aminothiols in tissue samples is shown in **Fig. 3** (*see Note 22*). Five male ddY mice 6 wk old ( $28.0 \pm 0.9$  g) were used in the experiments.

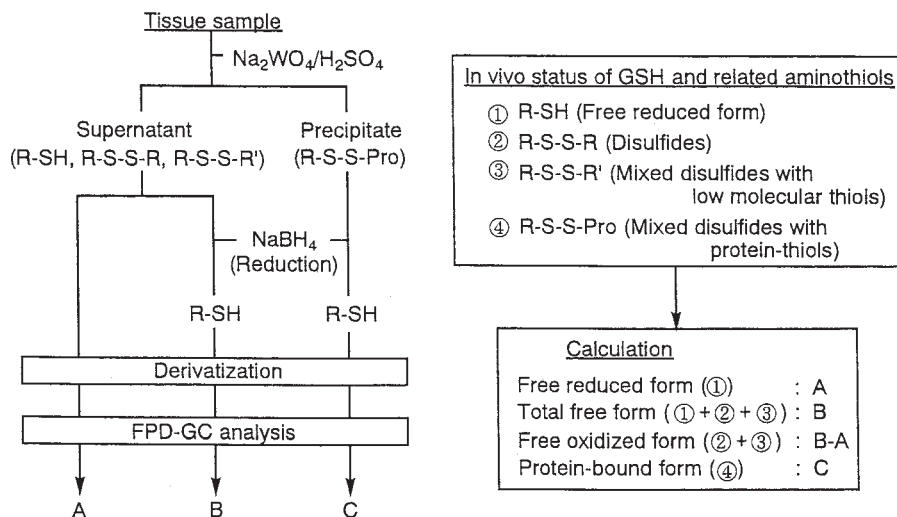


Fig. 3. Outline of method for the determination of different forms of glutathione and related aminothiols in tissue samples.

1. Immediately after dissection, each organ should be removed and stored at  $-20^{\circ}\text{C}$  until used.
2. Homogenize an aliquot of each pooled tissue (0.2–0.4 g) with 2.4 mL of 0.05 M H<sub>2</sub>SO<sub>4</sub> and 1.2 mL of 2.5% Na<sub>2</sub>WO<sub>4</sub> with an LK-21 ultradisperser (Yamato Kagaku, Tokyo, Japan) after adding one drop of *n*-hexanol.
3. Centrifuge at 2000g for 5 min, then reextract the precipitate with the same volume of 0.05 M H<sub>2</sub>SO<sub>4</sub> and 2.5% Na<sub>2</sub>WO<sub>4</sub> as above.
4. Combine the supernatants and make up to 10 mL with distilled water. This solution is then ready to use to determine the free and free oxidized forms.
5. For the analysis of free GSH and other aminothiols, 0.1 mL of a free fraction is directly derivatized.
6. For the analysis of total free GSH and other aminothiols (thiols + disulfides + mixed disulfides with other low-molecular-weight thiols), add 0.1 mL of the free fraction to 0.2 mL of 100 mg/mL of NaBH<sub>4</sub> in a 0.01 M NaOH solution, and incubate the mixture at 60°C for 5 min. After cooling, the reaction mixture is used as the sample for derivatization.
7. On the other hand, the precipitate is dissolved in 5 mL of 0.01 M NaOH and made up to 10 mL with distilled water, before being used to determine the protein-bound form. For the analysis of protein-bound GSH and other aminothiols, add 0.5 mL of the protein-bound fraction to 0.2 mL of 100 mg/mL of NaBH<sub>4</sub> in a 0.01 M NaOH solution, and incubate the mixture at 60°C for 5 min. After cooling, the reaction mixture is used as the sample for derivatization.

#### 4. Notes

1. IsoPCF and HCl-MeOH were used as derivatizing reagents for the amino and sulfhydryl groups, and for the carboxyl group, respectively (**Fig. 1**).
2. In order to prevent the oxidation of sulfur function during derivatization, DTE was added to reaction mixture and peroxide-free diethyl ether was used as an extraction solvent.
3. The *N(S)*-isopropoxycarbonylation of sulfur amino acids, GSH, and related aminothiols with isoPCF proceeded rapidly and quantitatively in aqueous alkaline media (**44,46**). This reaction was completed within 5 min by shaking at room temperature.
4. The reaction mixture was then washed with diethyl ether under alkaline condition in order to remove the excess reagent. This procedure also serves to exclude amines and phenols both of which often coexist with amino acids in biological samples, as they are derivatized to the corresponding *N*- or *O*-isoPOC derivatives, which are soluble in organic solvents under the same conditions as aforementioned.
5. The resulting *N(S)*-isoPOC derivatives in aqueous layer were quantitatively and selectively extracted into diethyl ether after acidification to pH 1.0–2.0. In this procedure, the ether layers should be collected, taking care to avoid aqueous droplets. It was not necessary to completely draw the ether layer in each extraction.
6. Although the methylation of the carboxyl group with diazomethane was simple (**53,54**), the interfering peaks originating in the diazomethane generating reagents were observed under our FPD-GC conditions. The methylation of *N(S)*-isoPOC derivatives with HCl-MeOH was completed within 5 min at 80°C, and no interfering peak was observed.
7. The derivative preparation was accomplished within 30 min, and several samples could be treated simultaneously. The *N(S)*-isoPOC methyl ester derivatives of sulfur amino acids and related compounds were stable under normal laboratory conditions and no decomposition was observed during GC analysis.
8. The GC analysis was performed with FPD (394-nm interference filter), which is highly selective for sulfur containing compounds, by using a megabore capillary column and direct (splitless) sample injection system. In this system, the sufficiently inactivated glass insert should be used to avoid tailing peaks of derivatives caused by the adsorption on injection port.
9. Nonsulfur amino acids were also derivatized and detected with flame ionization detection (FID) (**Fig. 4A**), but these amino acids were not detected at all with FPD (**Fig. 4C**). On the other hand, sulfur amino acids were detected with both FID (**Fig. 4B**) and FPD (**Fig. 4D**). As shown in **Fig. 4**, 13 sulfur amino acids and IS could be completely resolved as single and symmetrical peaks within 32 min on a DB-17 capillary column (15 m × 0.53-mm id, 1.0- $\mu$ m film thickness) (**44**).
10. The separation of biologically important sulfur amino acids was achieved within 18 min on a DB-17 capillary column (**Fig. 5A**). However, the derivatives of GSH and  $\gamma$ -GluCys could not be eluted with this capillary column because of high molecular mass and high boiling point of these derivatives.



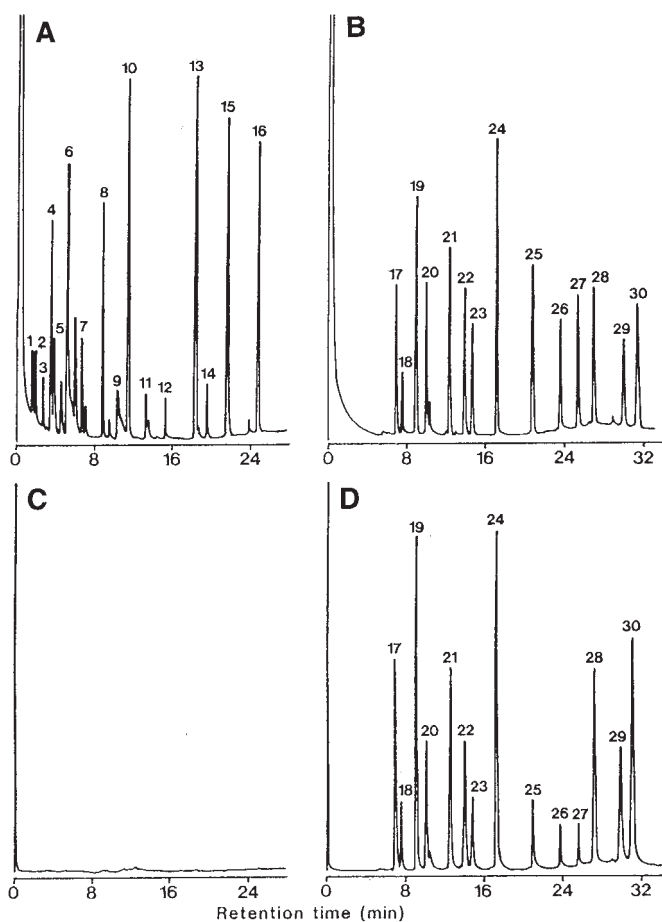


Fig. 4. Standard chromatograms of the *N(O,S)*-isoprpxycarbonyl methyl ester derivatives of nonsulfur and sulfur amino acids. **(A)** Nonsulfur amino acids (containing 200 nmol of each amino acid); **(B)** sulfur amino acids (containing 200 nmol of each amino acid); **(C)** nonsulfur amino acids (containing 50 nmol of each amino acid); **(D)** sulfur amino acids (containing 1 nmol of each amino acid). The derivatized samples were analyzed by FID-GC (**A** and **B**) and FPD-GC (**C** and **D**). GC conditions: column, DB-17 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, programmed at 5°C/min from 130°C to 270°C and then held for 5 min; injection and detector temperatures, 280°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = alanine; 2 = glycine; 3 = valine; 4 = leucine; 5 = isoleucine; 6 = serine + threonine + proline; 7 = aspartic acid; 8 = glutamic acid; 9 = hydroxyproline; 10 = phenylalanine; 11 = asparagine; 12 = glutamine; 13 = lysine; 14 = histidine; 15 = tyrosine; 16 = tryptophan; 17 = S-methylcysteine (SM-Cys); 18 = thioproline (TPro); 19 = methionine (Met); 20 = ethionine (Eth); 21 = cysteine (Cys); 22 = S-carboxymethylcysteine (SC-Cys); 23 = homocysteine (HCys); 24 = methionine sulfone (Met-S); 25 = S-2-aminoethylcysteine (AE-Cys) (IS); 26 = lanthionine (LTH); 27 = cystathionine (CTH); 28 = cystine (Cyt); 29 = djenkolic acid (Dje); 30 = homocystine (HCyt) (from **ref. 44**, with permission).

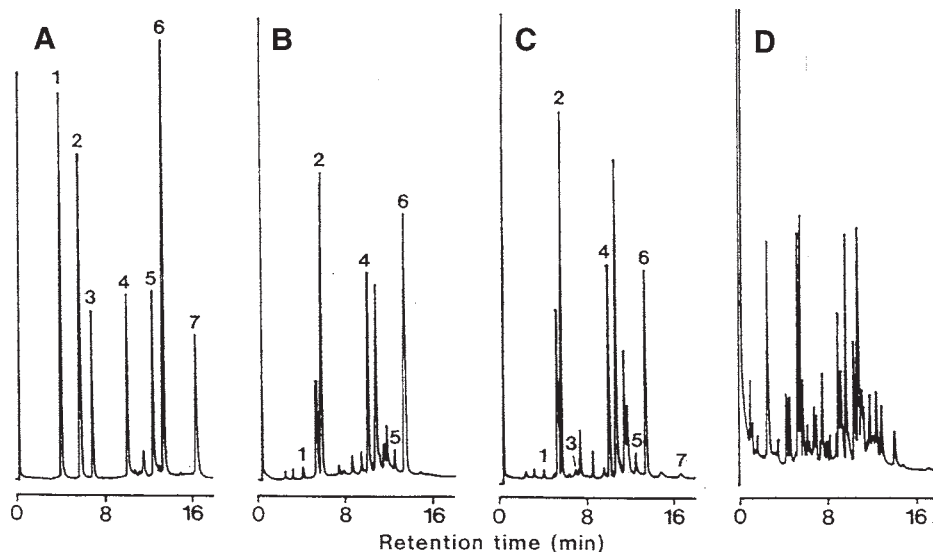


Fig. 5. Chromatograms obtained from standard and urine samples. (A) Standard (containing 2 nmol of each amino acid); (B) urine A (20  $\mu$ L); (C) urine B (20  $\mu$ L); (D) urine B (0.8 mL). The derivatized samples were analyzed by FPD-GC (A–C) and FID-GC (D). GC conditions: column, DB-17 (15 m  $\times$  0.53-mm id, 1.0- $\mu$ m film thickness); column temperature, programmed at 10°C/min from 150°C to 270°C and then held for 5 min; injection and detector temperatures, 280°C; nitrogen carrier gas flow rate, 10 mL/min. Peaks: 1 = methionine (Met); 2 = cysteine (Cys); 3 = homocysteine (HCys); 4 = S-2-aminoethylcysteine (AE-Cys) (IS); 5 = cystathionine (CTH); 6 = cystine (Cyt); 7 = homocystine (HCyt) (from ref. 44, with permission).

11. In order to solve above problem, a column was cut short. Of different length of several columns tested, DB-1 (5 m  $\times$  0.53-mm id, 1.5- $\mu$ m film thickness) proved to be the most satisfactory column for the purpose. By using this column, Cys, CysGly,  $\gamma$ -GluCys, GSH, and IS were separately eluted as single and symmetrical peaks within 10 min (Fig. 6A) (46). However, GSSG, disulfide form of GSH, could not be eluted even with this column.
12. The calibration curves for sulfur amino acids by FPD-GC were conducted using AE-Cys, which showed a similar behavior to other sulfur amino acids during the derivatization and was well separated from these amino acids on a chromatogram as the IS (44,45). A linear relationship was obtained from double logarithmic plots in the range 0.5–10 nmol of sulfur amino acids, and correlation coefficients were above 0.998 (Table 1).
13. The derivatives of sulfur amino acids provided excellent FPD responses and the minimum detectable amounts of these amino acids at a signal three times as high as the noise under our instrumental conditions were approx 0.5–1.0 pmol as injec-

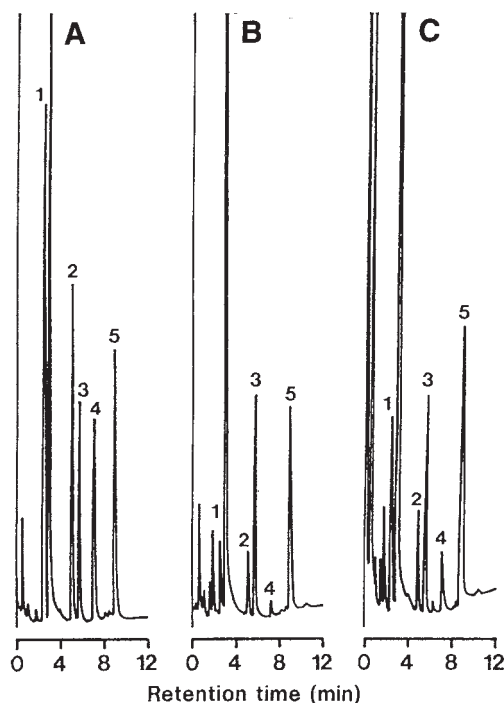


Fig. 6. Chromatograms obtained from standard and whole blood samples. (A) Standard (containing 5 nmol of glutathione and 1 nmol of other aminothiols); (B) blood (10 mL, nonreduction); (C) blood (5 mL, reduction). GC conditions: column, DB-1 (5 m  $\times$  0.53-mm id, 1.5- $\mu$ m film thickness); column temperature, programmed at 16°C/min from 130°C to 290°C; injection and detection temperature, 300°C; nitrogen flow rate, 12 mL/min. Peaks: 1 = cysteine (Cys), 2 = cysteinylglycine (CysGly), 3 = lanthionine (LTH) (IS), 4 =  $\gamma$ -glutamylcysteine ( $\gamma$ -GluCys), 5 = glutathione (GSH) (from ref. 46, with permission).

tion amounts. The FPD-GC system described here was over 20 times more sensitive than the FID-GC system.

- The calibration curves for GSH and related aminothiols by FPD-GC were conducted using LTH, which showed a similar behavior to other aminothiols during the derivatization and was well separated from these aminothiols on a chromatogram as the IS (46,47). A linear relationship was obtained from double-logarithmic plots in the range 1–25 nmol for GSH and 0.2–5 nmol for other aminothiols, and correlation coefficients were above 0.999 (Table 2).
- The minimum detectable amount of GSH and related aminothiols to give a signal three times as high as the noise under our FPD-GC conditions were approx 0.2–5 pmol as injection amounts.

**Table 1**  
**Linear Regression, Detection Limits, and Recoveries from Urine Samples on the Determination of Sulfur Amino Acids**

| Sulfur amino acid | Regression line <sup>a</sup> |           | Correlation coefficient | Detection limit (pmol) | Urine A                   |                                              |              | Urine B                   |                                              |              |
|-------------------|------------------------------|-----------|-------------------------|------------------------|---------------------------|----------------------------------------------|--------------|---------------------------|----------------------------------------------|--------------|
|                   | Slope                        | Intercept |                         |                        | Amount found Non-addition | (nmol/mL) <sup>b</sup> Addition <sup>c</sup> | Recovery (%) | Amount found Non-addition | (nmol/mL) <sup>b</sup> Addition <sup>c</sup> | Recovery (%) |
| Met               | 1.822                        | -0.222    | 0.9983                  | 0.5                    | 12.9 ± 0.6                | 111.0 ± 6.2                                  | 98.1         | 8.0 ± 0.5                 | 93.0 ± 0.3                                   | 85.0         |
| Cys               | 1.777                        | -0.308    | 0.9989                  | 0.5                    | 33.9 ± 1.2                | 136.5 ± 1.3                                  | 102.6        | 69.0 ± 1.5                | 165.0 ± 4.5                                  | 96.0         |
| HCys              | 1.703                        | -0.514    | 0.9995                  | 0.5                    | ND <sup>d</sup>           | 90.0 ± 2.2                                   | 90.9         | ND                        | 92.5 ± 2.5                                   | 92.5         |
| CTH               | 1.781                        | -0.548    | 0.9998                  | 1.0                    | 22.1 ± 0.9                | 122.0 ± 1.2                                  | 99.9         | 22.0 ± 0.3                | 114.5 ± 2.5                                  | 92.5         |
| Cyt               | 1.834                        | -0.188    | 0.9998                  | 1.0                    | 45.0 ± 1.2                | 145.0 ± 0.9                                  | 100.0        | 61.5 ± 0.3                | 159.0 ± 3.5                                  | 97.5         |
| HCyt              | 1.859                        | -0.059    | 0.9986                  | 1.0                    | ND                        | 113.0 ± 1.3                                  | 113.0        | ND                        | 107.5 ± 1.5                                  | 107.5        |

<sup>a</sup> $\log y = a \log x + b$  :  $y$ , peak height ratio against the IS;  $x$ , amount of each amino acid (nmol);  $a$ , slope;  $b$ , intercept. Range: 0.5–10 nmol.

<sup>b</sup>Mean ± SD ( $n = 3$ ).

<sup>c</sup>Addition: 100 nmol/mL.

<sup>d</sup>Not detectable.

**Table 2**  
**Linear Regression, Detection Limits, and Recoveries from Blood Samples on the Determination of GSH and Related Amino Thiols**

| Amino thiol | Regression line <sup>a</sup> |           |                         | Detection limit (pmol) | Non-reduced blood |                        |              | Reduced blood |                        |              |
|-------------|------------------------------|-----------|-------------------------|------------------------|-------------------|------------------------|--------------|---------------|------------------------|--------------|
|             | Slope                        | Intercept | Correlation coefficient |                        | Amount found      | (nmol/mL) <sup>b</sup> | Recovery (%) | Amount found  | (nmol/mL) <sup>b</sup> | Recovery (%) |
| Cys         | 1.418                        | 0.367     | 0.9997                  | 0.2                    | 19.0 ± 1.4        | 111.2 ± 3.8            | 92.2         | 83.3 ± 2.2    | 260.7 ± 8.3            | 88.7         |
| CysGly      | 1.419                        | 0.186     | 0.9992                  | 0.3                    | 19.5 ± 0.9        | 110.2 ± 8.3            | 90.7         | 55.4 ± 2.6    | 248.9 ± 13.1           | 96.8         |
| γ-GluCys    | 1.419                        | -0.029    | 0.9990                  | 0.5                    | 14.3 ± 1.0        | 104.6 ± 3.2            | 90.3         | 72.6 ± 4.8    | 286.0 ± 7.7            | 106.7        |
| GSH         | 1.677                        | -1.154    | 0.9993                  | 5.0                    | 464 ± 34          | 954.0 ± 42             | 98.0         | 958 ± 55      | 1977 ± 121             | 101.9        |

<sup>a</sup> $\log y = a \log x + b$  :  $y$ , peak height ratio against the IS;  $x$ , amount of each amino acid (nmol);  $a$ , slope;  $b$ , intercept. Range: 1–25 nmol for GSH, 0.2–5 nmol for the other amino thiols.

<sup>b</sup>Mean ± SD ( $n = 3$ ).

<sup>c</sup>Addition: 500 nmol/mL for GSH, 100 nmol/mL for the other amino thiols.

<sup>d</sup>Addition: 1000 nmol/mL for GSH, 200 nmol/mL for the other amino thiols.

16. The method developed for sulfur amino acids was successfully applied to small urine (0.02–0.1 mL) (**44**) and plasma (0.1 mL) (**45**) samples without prior clean-up of the sample. As shown in **Fig. 5B,C**, unknown peaks were observed in front of the Cys peak and in the near of the IS peak, but Met, Cys, CTH, Cyt, and trace quantities of HCys and HCyt were detected without any interference from coexisting substances when the sample was analyzed by FPD-GC.
17. It was difficult to determine sulfur amino acids in urine by FID-GC (**Fig. 5D**). As shown in **Table 1**, the overall recoveries of sulfur amino acids added to urine samples were 85–113% and the relative standard deviations were 0.3–6.4% ( $n = 3$ ).
18. The method developed for GSH and related aminothiols was successfully applied to blood sample, in which whole blood was used because it should reflect more accurately the total GSH status of blood including GSH bound to plasma proteins, and erythrocytes contain almost all of the GSH (**55**). Hemolysis of red blood cell was carried out in solution of EDTA to prevent oxidation of thiol function by any metallic elements. The hemolysate of small blood (5–10  $\mu$ L) could be used without prior clean-up of the sample (**46**). **Figure 6B,C** show typical chromatograms obtained from nonreduced and reduced blood samples, respectively. Free and total GSH and related aminothiols in whole blood sample could be analyzed without any interference from coexisting substances. As shown in **Table 2**, the overall recoveries of GSH and related aminothiols added to whole blood samples were 88–107% and the relative standard deviations were 2.6–7.5% ( $n = 4$ ).
19. In order to determine the redox status of sulfur amino acids (**45**), GSH, and related aminothiols (**46,47**), disulfides in the sample must be reduced to the corresponding thiols prior to derivatization.  $\text{NaBH}_4$  efficiently converts all these aminothiols present as free thiols and mixed disulfides with other low-molecular mass thiols or protein-thiols to thiol forms (**20–22**).
20. Formation of gas and foaming during the  $\text{NaBH}_4$  reduction of biological samples could be reduced by adding one drop of *n*-hexanol, surface active agent. The reduced sample could be directly derivatized after cooling.
21. In order to confirm the reduction conditions, some disulfide compounds such as Cyt, HCyt, and GSSG were reduced with  $\text{NaBH}_4$ . The reduction of these compounds were accomplished within 10 min at 100°C by using 20 mg  $\text{NaBH}_4$  in aqueous alkaline medium.
22. In order to determine the concentration of the different forms of GSH and related aminothiols in tissue samples, the fractionation of the tissue samples was investigated according to the previous method (**56**). As shown in **Fig. 3**, the sample was separated into its free and protein-bound fractions by precipitation with  $\text{Na}_2\text{WO}_4\text{-H}_2\text{SO}_4$ . No free GSH or other aminothiols were apparent in the protein-bound fraction. **Figure 7B–D** shows typical chromatograms obtained from the free fraction (-reduction), free fraction (+reduction), and protein-bound fraction (+reduction), respectively. Each form of GSH and related aminothiols in a tissue sample could be analyzed without any interference from coexisting substances. The overall recoveries of GSH and related aminothiols added to the free and protein-bound

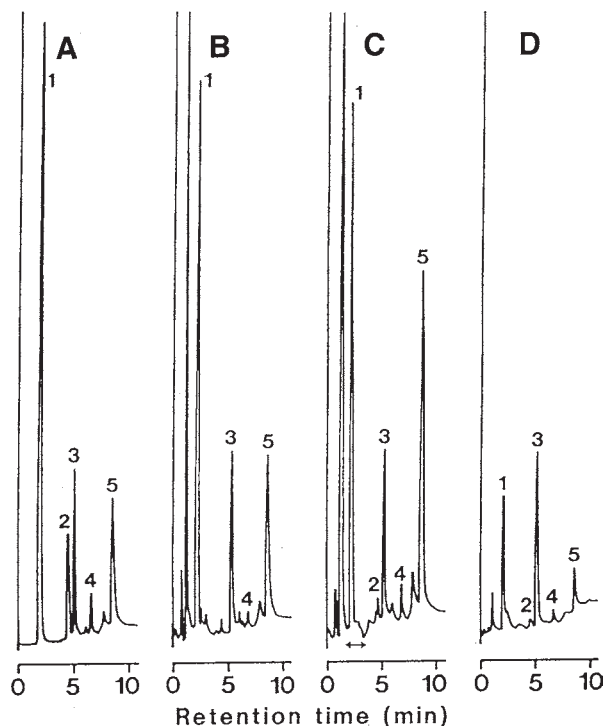


Fig. 7. Gas chromatograms obtained from standard solution and free and protein-bound fractions of mouse liver. (A) Standard containing 0.25 nmol of cysteine, 0.5 nmol of cysteinylglycine and  $\gamma$ -glutamylcysteine and 5 nmol of glutathione; (B) free fraction (-reduction); (C) free fraction (+reduction); (D) protein-bound fraction (+reduction). GC conditions and peak no.: see Fig. 6. The double arrow on the time-scale indicates the time during which the recorder response was reduced to half (from ref. 47 with permission).

fractions of mouse liver were 91.2–109.6%, and the relative standard deviation was 1.1–10.7% ( $n = 4$ ). Oxidized GSH and related aminothiols (disulfides and mixed disulfides with low-molecular weight thiols) were measured by subtracting their free reduced contents from their total free contents.

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## Capillary Electrophoretic Determination of 4-Hydroxyproline

Qingyi Chu and Michael Zeece

### 1. Introduction

Capillary electrophoresis (CE) represents a relatively new separation technology that has gained acceptance in a wide variety of applications. A discussion of the basic theory of CE separations is beyond the scope of this presentation, but is well addressed in a number of recent texts (1–5). In principle, differential migration of analytes in a potential field is achieved as a result of individual differences in mass/charge ratios. In CE, substantial advantage in separation speed, efficiency, and resolution is derived from the very-high field strengths (e.g., 150–300 V/cm) typically used. In addition, high field strengths induce the effect of electroosmosis in the column. The walls of the silica column have a negative charge and attract hydrated counter ions from the buffer. When power is applied to the system, the positively charged ions with their associated water molecules, migrate toward the cathode with substantial velocity. This results in the flow of water termed electroosmosis. The flow proceeds from the anode to the cathode and serves as a pump. Because the electroosmotic water flow is much greater than the velocity of the analytes, all components are swept to the cathode.

Most columns used in CE have an internal diameter of 50–100  $\mu\text{m}$  and can be 1 m long. High field potentials can be used because the column has a large surface area to internal volume ratio that facilitates rapid dissipation of heat. Extremely small volume samples are required for CE analysis. For example, a typical amino acid separation, such as those presented here, employ 10–20 nL sample injections. The small volume of sample injection however, also presents a challenge for detection. In order to achieve good sensitivities with UV detection, it is necessary to label the amino acids with phenylisothiocyanate

(PITC) prior to analysis. This reaction converts the amino acids to their phenylthiohydantoin (PTH) derivatives.

The mode of CE separation used for the amino acid analysis described here employs the anionic detergent sodium dodecyl sulfate (SDS). CE performed in the presence of SDS is often termed a micellar electrokinetic chromatography (MEKC) separation (6). When SDS concentrations are above 8–10 mM, the molecules form micelles in which the hydrophobic tails are oriented toward the center and sulfate groups are on the surface. The micelles have a strong negative charge at the pHs employed in most MEKC separations and migrate toward the anode. Their velocity, however, is less than that of the electroosmotic flow and all components are carried to the cathode. The SDS micelle thus becomes a pseudostationary phase with which analytes can interact. The resulting partitioning provides additional selectivity and the technique has been extensively used for the separation and determination of amino acids and other compounds (7).

## 2. Materials

### 2.1. Equipment

1. CE units are available from several well-known vendors and the method described here was performed with an ISCO (Lincoln, NE) model 3850 capillary electropherograph. Samples are loaded into the capillary by vacuum injection and detection of separated analytes is performed at 254 nm.
2. A CAESAR (Version 4.01, 1994, Roman Scientific) from Scientific Resources (Eatontown, NJ) was used to record and analyze data.
3. A Centri-Vap (Fisher Scientific, Pittsburgh, PA) or similar device for drying small volumes in microfuge tubes.
4. An amine coated capillary (50- $\mu\text{m}$  id Polymicro Technology, Phoenix AZ).

### 2.2. Reagents

Highest purity reagents were used throughout the experiments.

1. Phenyl isothiocyanate was obtained from Pierce (Rockford, IL).
2. Sodium hydroxide.
3. Hydrochloric acid.
4. Triethylamine.
5. Diethanolamine.
6. Malonic acid .
7. Chloro(chloro.methyl)dimethylsilane.
8. Phthalic dicarboxaldehyde.
9. 3- and 4-Hyp.
10. Cis and trans 4-Hyp.
11. Methanol.
12. Sodium phosphate (monobasic).

13. SDS.
14. Thymine (as internal standard).
15. Water used in these experiments was obtained by reverse osmosis (18 mOhm).

### 2.3. Amine Capillary Preparation

An amine coated capillary was used in these separations (50- $\mu\text{m}$  id Polymicro Technology, Phoenix, AZ). The length to the detector was 40 cm and total column length was 70 cm. This column can be purchased from commercial vendors or can be made in the lab (*see Note 1*). The procedure for coating the capillary with diethanolamine was originally described by Kuhn and Hoffstetter-Kuhn (8) and is summarized as follows.

1. Using a 1-mL syringe fitted with an adapter, fill 1 m of fused silica capillary with 1 M NaOH and let stand at room temperature (20°C) for 1 h.
2. Rinse the capillary twice with water.
3. Rinse the capillary twice with methanol followed by two rinses with acetone
4. Fill the capillary with 10% (w/v) chloro(chloromethyl)-dimethylsilane in methanol and hold for 30 min at room temperature.
5. Cover the ends of the silane-filled capillary with tape and incubate at 100°C for 3 h.
6. Flush the column with first with methanol, then with water.
7. Fill the capillary with 3.0 M diethanolamine HCl dissolved in methanol, and incubate overnight at room temperature.

## 3. Methods

### 3.1. PTH Labeling of Hyp Standards

Standards are necessary for identification of the Hyp peak in the separation and for development of standard curves. The preparation of these standards as PTH derivatives of Hyp isomers was performed essentially as described by Chu et al. (9) and summarized below.

1. Dissolve 6.6 mg of Hyp in 1.0 mL of coupling buffer. This solution contains methanol-triethylamine-water in 7:1:1 (v/v) ratio. A 1.5 or 2.0 mL microfuge tube works well for holding this mixture.
2. Flush the tube with nitrogen gas and add 10  $\mu\text{L}$  of PITC per 100  $\mu\text{L}$  of sample.
3. Incubate the reaction for 10 min at room temperature and then dry in a Centri-Vap (Fisher Scientific).
4. The samples are stored dry at -70°C and are dissolved in 100  $\mu\text{L}$  acetonitrile just prior to electrophoresis (*see Note 2*).

### 3.2. Preparation of Muscle Samples

Hyp determination can be made from isolated collagen fractions or whole muscle tissues samples. The procedures for preparing these materials for labeling is summarized as follows (*see Note 3*).

### 3.2.1. Isolated Collagen Fractions

Perimysial or other types of collagen can be isolated from skeletal muscle tissue using the well-established procedures described by Light and Champion (10). Following isolation, the collagen samples are prepared as follows (see **Note 4**).

1. Desiccate the material by washing three times in cold (4°C) acetone. For each wash, the sample is suspended in 4 vol of cold acetone, stirred by hand and then centrifuged at 2000g for 10 min. The supernatant is discarded each time. Be sure to use polyethylene type tubes, because clear plastics such as polycarbonate, are attacked by the solvent.
2. Dry the sample in a hood under a stream of nitrogen.
3. Add 50 mg of collagen to a suitable hydrolysis vial.
4. Add 5.0 mL of 6 M HCl, draw a vacuum, and heat for 24 h at 110°C.
5. Centrifuge or filter the sample after hydrolysis to remove any insolubles.
6. Withdraw 100- $\mu$ L aliquots (equivalent to approx 1.0 mg collagen sample) and place in a microfuge tube.
7. Dry the samples in a Centri-Vap. The dried samples are now ready for labeling, but can also be stored at -70°C.

### 3.2.2. Whole Muscle

To analyze whole muscle tissue for Hyp content, the tissue must first be dried and then hydrolyzed. The preparation of muscle samples is as follows (see **Note 4**).

1. Grind or finely dice 10 g of muscle.
2. Add 100–200 mg of ground muscle to a mortar and pestle prechilled with liquid nitrogen. Add liquid nitrogen slowly and grind to a powder. Keep the sample frozen hard during grinding with subsequent additions of liquid nitrogen.
3. Lyophilize the sample.
4. Add 65 mg of dried muscle powder in a suitable hydrolysis vial.
5. Add 5 mL of 6 M HCl, draw a vacuum and heat for 24 h at 110°C.
6. Centrifuge or filter the sample after hydrolysis to remove any insolubles.
7. Withdraw 150- $\mu$ L aliquots (equivalent to about 2.0 mg muscle) and place in a microfuge tube.
8. Dry the samples in a Centri-Vap. The dried samples are now ready for labeling, but can also be stored at -70°C.

### 3.3. Labeling of Amino Acids from Hydrolyzed Samples

The strategy for labeling amino acids from hydrolyzed samples is based on a modification of the procedure described by Yaegaki et al. (11). Amino acids from hydrolyzed tissues are labeled using a two-step procedure that involves initial reaction with *o*-phthalaldehyde (OPA) to react with primary amines fol-

lowed by their precipitation under acidic conditions. In the second step, amino acids (Pro and Hyp) are reacted with PITC. The procedure is summarized as follows (*see Note 5*).

1. Add 100  $\mu\text{L}$  of 0.5 M  $\text{Na}_2\text{CO}_3$ ,  $\text{NaHCO}_3$  pH 9.5, to the dried aliquot of hydrolyzed sample. The sample should contain approx 1.0 mg of hydrolyzed material.
2. Add 100  $\mu\text{L}$  (2 mg) of 20 mg/mL OPA reagent, mix, and let stand at room temperature for 15 min.
3. Precipitate the reaction products by adding of 900  $\mu\text{L}$  of 1.0 M  $\text{NaH}_2\text{PO}_4$  pH 2.5. Be sure to mix the sample well (e.g., in a vortex mixer).
4. Centrifuge at 10,000g for 5 min and filter the supernatant through 0.45- $\mu\text{m}$  spin-filters (Millipore, Bedford, MA)
5. Dry 100  $\mu\text{L}$  of filtrate in the Centri-Vap.
6. Add 150  $\mu\text{L}$  coupling buffer (described in **Subheading 3.1**).
7. Flush the vial with nitrogen and add 10  $\mu\text{L}$  of PITC.
8. Incubate the mixture at room temperature for 10 min and Centri-Vap to dryness.
9. The dried samples are ready for CE analysis or can be stored at  $-70^\circ\text{C}$ .

### 3.4. CE Analysis

Samples are ready for analysis after the appropriate preparation steps have been performed. The procedure for CE for identification and determination of Hyp in these samples is described below (*see Note 6*).

1. Add 100  $\mu\text{L}$  50% acetonitrile to the tube containing the dried and labeled amino acid sample (prepared in **Subheading 3.3**). Mix well by vortexing.
2. Centrifuge at 10,000g for 2 min.
3. Withdraw 30  $\mu\text{L}$  of the sample supernatant and add 10  $\mu\text{L}$  2.0 mM thymine (internal standard).
4. Equilibrate the column filling it with the run buffer (50 mM Na-malonate, 75 mM SDS, pH 5.0).
5. The sample is then loaded to the CE using either vacuum or pressure injection following the CE manufacturer's instructions. Generally, loads will be in the order of 5 to 50 nL. The equipment manufacturer should provide information regarding determination of load volumes.
6. Set the instrument voltage at 15 kV and start the run.
7. Record the data for all peaks.
8. Identify the thymine and Hyp peaks in the separation from the procedure described below.

### 3.5. Quantitation of Hyp in Samples

In order to determine the amount of Hyp accurately in unknown samples, it is first necessary to identify the peak in the mixture using coinjection and then use an internal standard method for its quantitation. The procedures described here were developed for determination of Hyp in muscle tissue samples (9). It



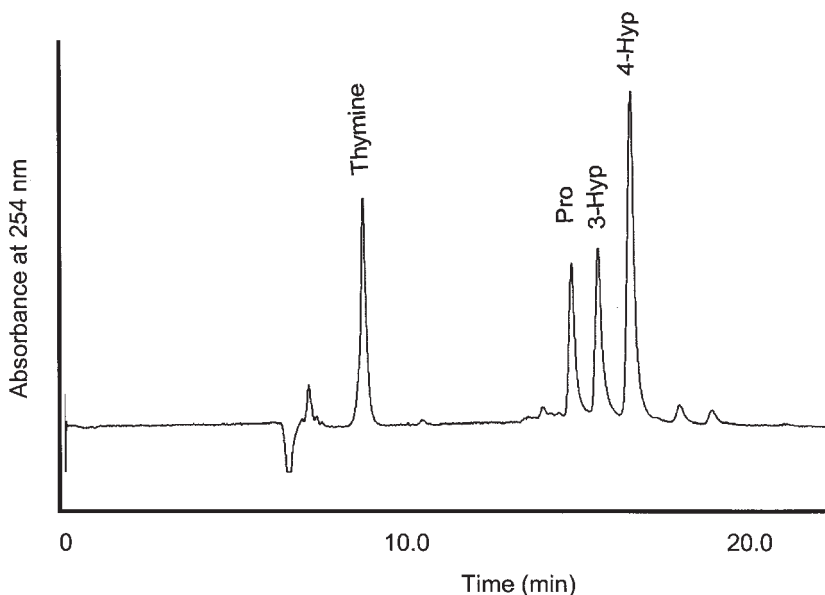


Fig. 1. Electropherogram of PTH-Proline, 3- and 4-Hydroxyproline standards. Amino acids were labeled with PITC as described in **Subheading 3.1.** to obtain the PTH derivatives. Approximately 20 nL of a mixture containing 2.0 mM each of PTH-Proline, 3- and 4-Hydroxyproline was separated by capillary electrophoresis in 50 mM Na-malonate, 75 mM SDS, pH 5.0 at 15 kV. The capillary was amine coated, 50  $\mu$ m in id and 70 cm in length (40 cm to the detector). The migration of the components are identified in the figure.

is important for accurate quantitation that the Hyp peak be reasonably separated from other peaks in the sample. Thus it may be necessary to adjust parameters such as voltage or buffer composition when analyzing samples prepared from other tissues.

### 3.5.1. Identification of Hyp in the Separation (Electropherogram)

In order to identify the Hyp peak in a sample, the migration time must first be determined using standards (prepared in **Subheading 3.1.**) (*see Note 7*). A sample electropherogram illustrating the separation of proline, 3-hydroxyproline and 4-hydroxyproline is shown in **Fig. 1**. The procedure is as follows.

1. Dilute the appropriate PTH-Hyp isomer to 2.0 mM in acetonitrile.
2. Add 10  $\mu$ L of 2.0 mM thymine per 30  $\mu$ L of Hyp solution.
3. Mix and perform CE separation as described in **Subheading 3.4.** The conditions for separation of the standard (s) must be the same as those used for samples.

### 3.5.2. Coinjection of Hyp

1. Obtain the electropherogram for the unknown samples by performing the separation as described in **Subheading 3.4.**
2. Add 10  $\mu\text{L}$  2.0 mM Hyp to the sample and run the separation again.
3. Compare the two electropherograms. The peak with substantially increased area and height corresponds to the Hyp peak.

### 3.5.3. Standard Curve

1. Prepare a set of at least 6 Hyp dilutions covering the range of 0.05 to 2.0 mM.
2. Add 10  $\mu\text{L}$  2.0 mM thymine to 30  $\mu\text{L}$  of each standard concentration and mix.
3. Perform the separations as described in **Subheading 3.4.** using a constant injection volume (e.g., 10 nL).
4. Determine the **Hyp peak area ratio** for each standard concentration by dividing the peak area for Hyp by that of the thymine peak.
5. Plot the **Hyp peak area ratio** vs the Hyp concentration for each point. Use a linear regression analysis of the data points. The slope of this line (Hyp concentration/peak area ratio) represents the **Conversion Factor**. The concentration of Hyp in the unknown sample is calculated from the equation below.

$$\text{Hyp peak area ratio in unk} \times \text{Conversion Factor} \\ (\text{Hyp conc/peak area}) = \text{Hyp Conc}$$

6. To determine the amount (moles) of Hyp in the unknown sample, the total volume of the hydrolyzed aliquot must be used. For example, assuming that an aliquot corresponding to 1.0 mg of tissue was hydrolyzed, labeled, and finally dissolved in a final volume of 100  $\mu\text{L}$  then the total amount of Hyp in that sample is calculated as follows:

$$\text{Hyp Conc} \times \text{injection volume (nL)} \times \text{dilution factor} \\ (100 \mu\text{L}/10 \text{ nL}) = \text{moles Hyp}$$

Because the amount of tissue hydrolyzed is known, the Hyp content can also be expressed as moles of Hyp per mg tissue.

## 4. Notes

1. Amine capillary preparation: Since the time of our original investigations, the commercial availability of amine coated capillaries has significantly increased. The quality and consistency of prepared capillaries is well worth the cost. Thus, unless the investigator is very familiar with this technology, it is recommended that amine coated capillary be obtained through a commercial vendor.
2. PTH Labeling of Hyp Standards: The labeling reaction of Hyp isomers or Pro with PITC is rather easy to do. Precautions include flushing the reaction mixture with nitrogen and protecting it from light. UV light accelerates the degradation of PTH derivatives. Storage should be in a light protected vial at  $-20^\circ$  to  $-70^\circ\text{C}$ . The shelf life of PTH-Hyp was only 2–3 wk. Additional peaks representing breakdown products were observed in CE separations with increasing storage.

3. This method describes procedures for application of CE to the determination of Hyp isomers and Pro in muscle tissue. It is also applicable to the analysis of these amino acids in other tissues. Additional considerations that might be helpful in performing Hyp analysis are addressed below.
4. Whole muscle samples or isolated collagen fractions can be analyzed by this procedure. The protocol has a number of steps, but is faster and requires less sample than other (HPLC) methods. The sensitivity of this method was found to be sufficient such that determinations could be made in as little as 1 mg of sample (9).
5. A critical step for determination of Hyp in muscle samples involves labeling of amino acids in hydrolyzed sample (**Subheading 3.3.**). This is a two-step procedure in which amino acids (all except Hyp and Pro) are reacted with OPA. The OPA amino acids are then precipitated with 1 M  $\text{NaH}_2\text{PO}_4$ , pH 2.5. The insoluble amino acid derivatives are then removed by centrifugation; the supernatant containing imino acids (Hyp and Pro) is subsequently reacted in the second step with PITC. However, precipitation of OPA derivatized amino acids may not be complete with  $\text{NaH}_2\text{PO}_4$  in some samples. Alternatively, we have found that substitution of OPA with Naphthalenedicarboxaldehyde results in more efficient precipitation with  $\text{NaH}_2\text{PO}_4$  and greater solubility for Hyp and Pro.
6. The CE separation and determination of Hyp and Pro in biological samples is relatively straightforward and faster than many HPLC approaches because the smaller sample requirements result in faster processing time. For example, this procedure contains a number of steps in which drying of the sample is necessary. Because the CE procedure requires only 100–150  $\mu\text{L}$  hydrolysis aliquots, the time required to process the sample is substantially reduced. We have also successfully performed Hyp analysis with 1–5 mg samples using microwave hydrolysis containing 200–500  $\mu\text{L}$  of HCl, to further reduce the processing time.
7. A critical factor in the quantitation of Hyp in biological samples by CE is the requirement for an internal standard. CE uses very small sample volumes (nL) and thus slight differences in the injection volume can result in large variations for Hyp peak areas. The use of internal standard peak area ratio method eliminates the errors caused by injection differences. The thymine peak also provides a reference point (migration time) for assessing the consistency of electropherograms.

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## Total Plasma Homocysteine Analysis by HPLC with SBD-F Precolumn Derivatization

Isabella Fermo and Rita Paroni

### 1. Introduction

Over the last two decades, various studies have shown that moderate and persistent hyperhomocysteinemia is implicated in the development of atherosclerosis, which is responsible for 50% of all mortality and morbidity in Western countries. Considering that the most traditional risk factors for heart disease and stroke, such as plasma lipids, cigarette smoking, hypertension, obesity, and diabetes, only account for 50% of cardiovascular disease (1,2), one can understand the reason why homocysteine (Hcy) measurement is included in the list of tests for investigating the causes of atherosclerosis and thrombosis.

One of the major problems encountered in studies on the potential atherogenic role of Hcy was the development of an accurate and simple assay, capable of screening, in a normal population, subjects having a congenital predisposition to occlusive vascular disease. Several approaches have been described in literature for measuring total plasma homocysteine (tHcy), which is defined as the sum of free and protein-bound homocysteine, homocystine, and homocysteine-cysteine mixed disulfide. These procedures involve, after a reduction step, the use of gas-chromatography-mass spectrometry (GC-MS) (3), radioenzymic assay (4), and high-performance liquid chromatography (HPLC). The latter, which is the most widely applied, may be coupled with (1) spectrophotometric detection in the visible range (amino acid analyzer) after postcolumn ninhydrin derivatization (5); (2) direct electrochemical detection (6); or (3) spectrofluorimetric detection after precolumn derivatization with different labeling agents such as monobromobimane (mBrB) (7), *o*-phthalaldehyde (OPA) (8), or halogensulfonyl benzofurazans (ABD-F and SBD-F) (9,10).

Among the HPLC procedures for tHcy analysis in plasma, methods based on SBD-F (ammonium-7-fluorobenzo-2-oxa-1,3-diazole-4-sulphonate) reaction appear to be attractive for routine use in clinical chemistry laboratories. SBD-Hcy adduct is very stable (1 wk at 4°C) (*see Note 1*) (*11*) and SBD-F being a thiol-specific labeling molecule, the derivatization yields a very clean chromatographic profile in comparison to mBrB, which produces interfering fluorescence byproducts (*7*), or to OPA, which reacts unspecifically with all the amino groups of the sample (*8*). We have described here a procedure that involves reduction of disulfide bonds by sodium borohydride (NaBH<sub>4</sub>), use of mercaptopropionylglycine as internal standard (*12*), precolumn derivatization with SBD-F, and isocratic analysis by reversed-phase (RP) HPLC with fluorescence detection.

## 2. Materials

### 2.1. Equipment

1. Polypropylene micro tubes (2 mL) with screw-cap (Sarstedt, Numbrecht, Germany).
2. Microfuge model 11 (Beckman, Palo Alto, CA).
3. Block heater (Asal, Milano, Italy) or oven.
4. HPLC monopump model 116 with a solvent selector valve (Beckman, Palo Alto, CA) (*see Note 2*).
5. Fluorimetric detector model RF 551 (Shimadzu, Kyoto, Japan).
6. Autosampler model 507 (optional device) (Beckman).
7. Column: Ultrasphere ODS (150 × 4.6-mm id, 5 μm) (Beckman).
8. Guard-column: Lichrospher 100 RP-18 (40 × 4-mm id, 5 μm) (Merck, Darmstadt, Germany).

### 2.2. Reagents

1. NaBH<sub>4</sub>.
2. Dimethyl sulfoxide (DMSO).
3. 7-fluorobenzo-2-oxa-1,3-diazole-4-sulphonate (SBD-F).
4. Boric acid.
5. ethylenediaminetetracetic acid (EDTA).
6. L-homocystine (Hcy-Hcy).
7. Sodium acetate.
8. Sodium hydroxide (NaOH).
9. 70% perchloric acid (PCA).
10. Methanol (CH<sub>3</sub>OH).
11. Acetonitrile (CH<sub>3</sub>CN).
12. Glacial acetic acid.
13. Mercaptopropionylglycine (IS).
14. Bidistilled water (H<sub>2</sub>O).

### 2.3. Solutions

1. Reducing solution: Prepare 3 mol/L  $\text{NaBH}_4$  solution by dissolving 0.285 mg of  $\text{NaBH}_4$  in 2.5 mL of  $\text{NaOH}$  0.1 mol/L, then mix with DMSO (2:1, v/v) and store in a glass vial at 4°C. Prepare this solution fresh every week.
2. Deproteinizing solution: Prepare 0.6 mol/L PCA solution by mixing 25 mL PCA (70%) with 0.186 g EDTA and diluting to 500 mL with  $\text{H}_2\text{O}$ . This solution is indefinitely stable when stored at 4°C (see **Note 3**).
3. Buffer solution: Prepare 0.2 mol/L boric acid solution with 4 mmol/L EDTA, pH 9.5 by dissolving 12.37 g boric acid and 1.49 g EDTA in 1 L  $\text{H}_2\text{O}$ . Warm to dissolve. Adjust to pH 9.5 with concentrated  $\text{NaOH}$  while still warm. Cool to room temperature before use and store this solution at room temperature.
4. Neutralizing solution: Prepare  $\text{NaOH}$  2 mol/L by dissolving 80 g  $\text{NaOH}$  in 1 L  $\text{H}_2\text{O}$ .
5. Derivatizing solution: Prepare 4.2 mmol/L SBD-F solution by mixing 1 mg SBD-F with 1 mL of buffer solution. Store at 4°C in a dark vial for 1 d.
6. Internal standard solution: Mix 1 mg of IS with 1 mL  $\text{HCl}$  0.1 mol/L and 9 mL  $\text{H}_2\text{O}$ . Store 200- $\mu\text{L}$  aliquots at -20°C up to 6 mo. To use, dissolve each aliquot with 600  $\mu\text{L}$  solution 3 diluted 1:1 with  $\text{H}_2\text{O}$ . Store at 4°C for 1 wk.
7. Homocystine standard solution: Prepare solution 200  $\mu\text{mol/L}$  by dissolving in a 100 mL volumetric flask 5.4 mg of Hcy-Hcy in 5 mL  $\text{HCl}$  0.1 mol/L and diluting to 100 mL with  $\text{H}_2\text{O}$ . Freeze 300- $\mu\text{L}$  aliquots at -20°C for 6 mo. Use each aliquot only once and then discard.
8. Stock HPLC eluents: Prepare sodium acetate solution 0.2 mol/L by dissolving 27.22 g sodium acetate trihydrate in 1 L  $\text{H}_2\text{O}$ . Dilute 11.5 mL glacial acetic acid with 1 L  $\text{H}_2\text{O}$  and adjust to pH 4.0 with the sodium acetate solution ( $\approx$ 3:1) to obtain 0.2 M sodium acetate buffer. Store all these solutions at 4°C.
9. Working HPLC eluents: Mobile phase A: add 20 mL of  $\text{CH}_3\text{OH}$  to 980 mL of buffer 8 (2%  $\text{CH}_3\text{OH}$ ). Mobile phase B: add 300 mL of  $\text{CH}_3\text{OH}$  to 700 mL of solution 8 (30%  $\text{CH}_3\text{OH}$ ).

## 3. Methods

### 3.1. Preanalytical Steps

Collect whole blood specimens in sterile Vacutainer<sup>®</sup> tubes containing sodium citrate (Becton Dickinson, Rutherford, NJ). Keep the samples on ice until centrifugation at 3000g for 10 min at 4°C, then immediately freeze the plasma specimens at -20°C until analysis (see **Note 4**).

### 3.2. Procedure (see Flow Chart 1)

1. Reduction step. Mix 100  $\mu\text{L}$  plasma sample with 50  $\mu\text{L}$  of internal standard solution and 20  $\mu\text{L}$  of  $\text{NaBH}_4$  solution (reducing solution) in a polypropylene tube with screw-cap. Incubate at 50°C for 30 min (see **Note 5**).

### Flow Chart 1 SBD-F Precolumn Derivatization Method

---

|                                |   |                                                                                                          |
|--------------------------------|---|----------------------------------------------------------------------------------------------------------|
| • Plasma collection            | ⇒ | Anticoagulant: sodium citrate, eparin, or EDTA<br>- Blood sample in ice<br>- Prompt separation of plasma |
| • IS addition                  | ⇒ | 100 $\mu\text{L}$ mercaptopropionylglycine solution                                                      |
| • Reduction of disulfide bonds | ⇒ | by 20 $\mu\text{L}$ 3 mol/L $\text{NaBH}_4$ solution (30 min at 50°C)                                    |
| • Plasma deproteinization      | ⇒ | by 100 $\mu\text{L}$ 0.6 mol/L PCA in 1 mmol/L EDTA                                                      |
| • Plasma neutralization        | ⇒ | by 20 $\mu\text{L}$ 2 mol/L NaOH                                                                         |
| • Derivatization               | ⇒ | by 50 $\mu\text{L}$ SBD-F solution (60 min at 60°C)                                                      |
| • HPLC separation              | ⇒ | isocratic elution (14 min); reversed-phase C18 column                                                    |
| • Fluorimetric detection       | ⇒ | $\lambda_{\text{ex}} = 385 \text{ nm}$ ; $\lambda_{\text{em}} = 515 \text{ nm}$                          |

---

2. Deproteinization step. Add 100  $\mu\text{L}$  0.6 mol/L PCA (solution 2). Vortex, then centrifuge at 12000g for 10 min.
3. Derivatization step. Transfer 100  $\mu\text{L}$  of supernatant in an Eppendorf tube and add 200  $\mu\text{L}$  of 0.2 mol/L borate buffer pH 9.5 (buffer solution), 20  $\mu\text{L}$  of 2 mol/L NaOH (neutralizing solution), and 50  $\mu\text{L}$  of SBD-F solution (derivatizing solution). Vortex and then incubate the samples at 60°C for 1 h. Remove and allow vials to cool to room temperature before injecting (*see Note 6*).
4. Transfere derivatized samples into the autosampler vials (optional device) and inject 10  $\mu\text{L}$  into the HPLC (*see Notes 7 and 8*).

### 3.3. Calibration Curve

1. For each batch of samples to be run on HPLC, dilute the 200  $\mu\text{mol/L}$  homocysteine stock solution to get the final concentrations: 100, 50, and 25  $\mu\text{mol/L}$  (use solution 3 diluted 1:1 with  $\text{H}_2\text{O}$ ).
2. Spike 100  $\mu\text{L}$  of pooled plasma with 10  $\mu\text{L}$  of each of these hcy-hcy standard solutions to obtain a calibration curve in the 5, 10, 20, 40  $\mu\text{mol/L}$  range (Hcy molar equivalents) (*see Notes 7 and 9*). Plot the relative fluorescence intensities of SBF-Hcy/IS ( $y$ ) as a function of Hcy concentrations ( $x$ ), and perform the least square analysis to obtain the regression equation.
3. The unknown tHcy concentration in a plasma sample is determined by this calculation:

$$[\text{tHcy}]_{\text{sample}} \mu\text{mol/L} = (\text{Hcy peak area} : \text{IS peak Area})_{\text{sample}} / \text{slope of calibration curve}$$



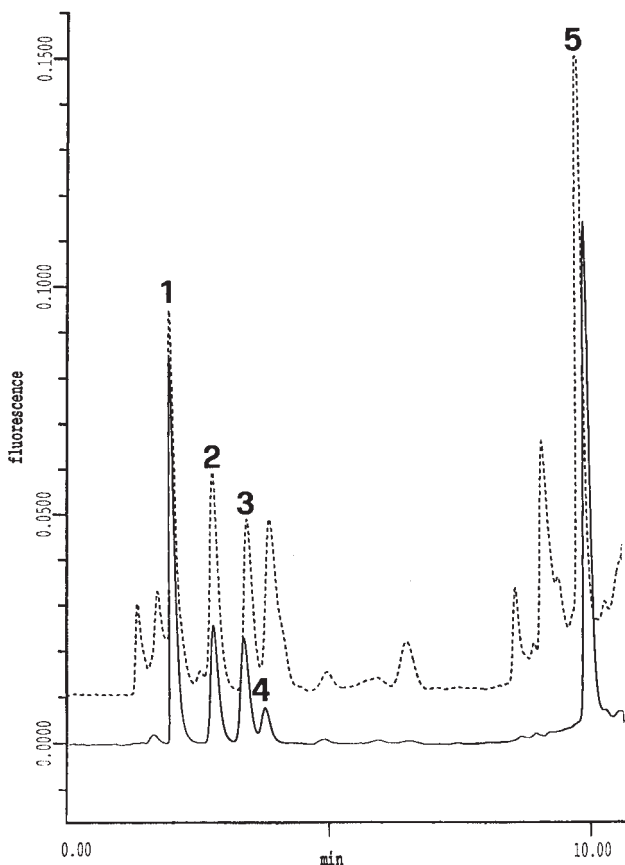


Fig. 1. Chromatograms of: (—) plasma sample with tHcy level of 23.0  $\mu\text{mol/L}$ ; (---) Bio-Rad Liquicheck™ Homocysteine Control-2 (lot. no. 18312, Bio-Rad Diagnostic Group, Irvine, CA). The separation was carried out on a Beckman Ultrasphere ODS column ( $150 \times 4.6\text{-mm}$  id;  $5\ \mu\text{m}$ ). (peak 1 = cysteine; peak 2 = homocysteine; peak 3 = cysteinylglycine; peak 4 = glutathione; peak 5 = mercaptopropionylglycine).

### 3.4. Chromatographic Analysis

1. Fluorimetric detector:  $\lambda_{\text{ex}} = 385\ \text{nm}$ ;  $\lambda_{\text{em}} = 515\ \text{nm}$ .
2. Run time: 14 min.
3. Equilibrate the column with 100% mobile phase A at 1.2 mL/min and 0.2 min after the analysis started, switch the selector valve to 100% eluent B.
4. At 4 min increase flow rate to 1.4 mL/min over 0.5 min, and at 4.5 min, switch back the valve to 100% eluent A.
5. At 12 min, decrease the flow rate to 1.2 mL/min in 0.5 min and at 14 min you are ready for the next injection.

## HPLC Run Program

| Time<br>(min) | Flow<br>(mL/min) | Duration<br>(min) | Mobile<br>phase |
|---------------|------------------|-------------------|-----------------|
| 0             | 1.2              | —                 | A               |
| 0.2           | 1.2              | —                 | B               |
| 4.0           | 1.4              | 0.5               | B               |
| 4.5           | 1.4              | —                 | A               |
| 12            | 1.2              | 0.5               | A               |
| 14            | STOP             | —                 | A               |

The chromatogram in **Fig. 1** shows the following peaks: cysteine (1), homocysteine (2), cysteinylglycine (3), glutathione (4), and mercaptopyrionylglycine (5).

#### 4. Notes

1. The fluorescence SBD-Hcy adduct, being photosensitive, has to be stored in the dark where is stable for 1 wk at 4°C (*11*).
2. This method requires the use of a simple monopump HPLC, but with a solvent selector valve. If the solvent switching is not available, the IS will be eluted later and the analysis will last too long. As an alternative, the method can be performed without the addition of IS, and the chromatographic separation will be performed using only the mobile phase A at flow rate 1.2 mL/min. In this case, we have to be careful, above all, during the reduction step and the use of a surface active agent is recommended. In addition, we suggest running the plasma calibration curve at least three times per day in order to check the performance of own detector system.
3. If the deproteinizing solution shows an amount of precipitate on the bottom of the glass vial, we are probably using out-of-date reagents. The reductant must to be completely dissolved in the vial before its use.
4. We adopted sodium citrate as anticoagulant as it is commonly used by the Coagulation Service of our Institute; no differences in tHcy plasma levels were found after EDTA or heparin treatment (*8*). It is important to know that differences in blood-sample handling can cause significantly different results in tHcy measurement. During whole blood storage at room temperature, Hcy concentration tends to increase over time because of the protracted production and release of Hcy by blood cells. This process is slowed down when blood samples are left on ice. There are no problems for plasma samples because tHcy in this biological fluid is stable for at least 4 d at room temperature, for several weeks at 4°C and for years at -20°C (*7*).
5. Sodium borohydride is a potent reductant, but because this reaction involves the formation of gas and sample foaming, the use of surface active agents such as amyl alcohol is also suggested. In our method, this addition is avoided thanks to (1) the use of 2-mL conical bottom tubes with screw-caps and (2) the addition of an internal standard. Pay attention during PCA deproteinization also: as the addition of an acid sparks off sample foaming. For this reason, we prefer to utilize tubes with an internal volume of 2 mL, at least.

6. Derivatization reaction can be directly performed in the autosampler vials if these tubes are heat-resistant and suitable for the block heater (incubation equipment).
7. The lowest limit for reliable quantification (LOQ) of the method is  $0.85 \mu\text{mol/L}$ , corresponding to  $1.7 \text{ pmol}$  injected (**12**), and the analysis is carried out by utilizing a simple fluorometric detector on line with the HPLC apparatus.
8. The normal ranges for plasma tHcy differ from one laboratory to another, and these discrepancies may be related to the procedure or technique used or the criteria chosen to select the control subjects. However, tHcy plasma levels between  $5.0$  and  $15.0 \mu\text{mol/L}$  are usually considered as normal (**13**). By applying this procedure, we found the following reference values:  $7.65 \pm 2.36$  and  $8.9 \pm 2.14 \mu\text{mol/L}$  for females ( $n = 40$ ) and males ( $n = 40$ ), respectively (**8**).
9. Plasma calibration curve must be used instead of aqueous calibrators because the latter give a linear equation with a different slope ( $-20\%$ ), by showing highlighting the notable influence of the matrix on SBD-F thiol derivatization (**9,12**). We can use Hcy standard instead of Hcy-Hcy, but, in this case, care must be taken with the purity of the Hcy powder (**14**). All standards must be stored at  $-20^\circ\text{C}$ .

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## Determination of Early Glycation Products by Mass Spectrometry and Quantification of Glycation Mediated Protein Crosslinks by the Incorporation of [<sup>14</sup>C]lysine into Proteins

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### 1. Introduction

The Maillard reaction, popularly known as nonenzymatic glycosylation (NEG) or glycation, is a complex chemical reaction and occurs in vivo between reactive aldose or ketose sugars and protein-bound free amino groups (1). NEG has been implicated in diabetic or age-related complications (2), Alzheimer's disease (3), and also in cataract formation (4). In vivo, any protein with free amino groups can react with reducing sugars via the Maillard reaction. However, the extent of damage caused by NEG is amplified in diabetic tissues (because of the elevated levels of blood sugar levels) and also in proteins with a long half-life like collagen and lens crystallin proteins (1,2). Therefore, determination of early glycation products and protein crosslinks produced because of glycation will be a significant aspect to investigate the extent of damage caused by NEG in vivo.

As shown in **Fig. 1**, the Maillard reaction involves the initial interaction of a reducing sugar (like glucose in **Fig. 1**) with  $\alpha$ -/ $\epsilon$ - amino groups of protein-bound amino acids, with the  $\epsilon$ - amino group on lysine as the preferential site. An initial labile Schiff base formed during the reaction undergoes Amadori rearrangement and yields a stable ketoamine derivative. Following dehydration and subsequent chemical modification, some of the Amadori products accumulate with time as advanced glycation end products (AGEs). Though it has been proposed that AGEs have a significant impact in age-related or diabetic

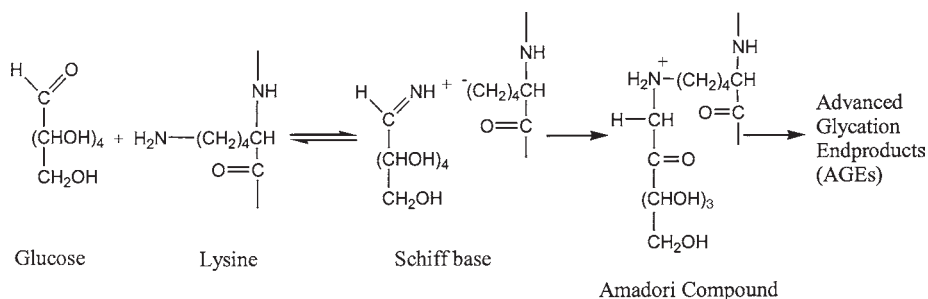


Fig. 1. Reaction scheme for the Maillard reaction (nonenzymatic glycation). Glucose is shown as the model sugar and lysine is shown as the model amino acid.

complications (1–3), it is still a debate to understand the mechanism involved in the development of glucose dependent diabetic complications. This is partly because of the slow reaction property of glucose with model proteins in vitro (5). Previously, we have shown that the oxidation products of ascorbic acid (which is relatively higher than glucose in human lens) can glycate and crosslink lens proteins at an accelerated rate compared to glucose under similar in vitro conditions (6–8), suggesting a possible role for ascorbate mediated glycation in vivo.

Characterization and quantification of glycation crosslinks has been the subject of study for decades. Several methods have been developed in the past to determine early and advanced glycation products. These methods include borohydride reduction (9), thiobarbituric acid assay (10), the fructosamine assay (11), periodate assay (12), size-exclusion chromatography (13), ion-exchange chromatography (14), boronate-affinity chromatography (15), and immuno detection (16). With the exception of immunological methods, almost all the glycation assay methods are based on chemical methods, and are nonspecific. Additionally, in vitro assay methods do not guarantee the detection of similar compounds in vivo. For example, some of the reagents used in borohydride reduction, thiobarbituric acid assay, acid/base hydrolysis, or periodination can nonspecifically interact with nonglycated products in the assay system and generate faulty reaction products that are counted as glycated products. Other examples are the fructosamine assay and phenylboronate assay, which are based on the detection of serum albumin reacted with glucose (aldose sugar). Both these assay methods failed to detect serum albumin glycated by fructose (ketose sugar) in diabetics (17). Although measurement of fluorescent compounds like pentosidine (18) in glycated proteins is an alternate choice, not all AGEs (e.g., carboxymethyl lysine, ref. 19) are fluorescent in nature. Another drawback in estimation of fluorescent AGEs is the highly crosslinked or insol-

uble nature of glycated proteins in diabetic or aged tissues. Therefore, alternate assay methods to detect and quantitate the early and advanced glycation end products represent a needed and valid approach to estimate the damage caused by nonenzymatic glycation *in vivo*.

Recently, we have shown that mass spectrometry (MS) can be effectively used to detect and determine the early glycation products formed in a model glycation system (20). MS has successfully been used to precisely determine glycated proteins or peptides in model glycation systems (21–23) (*see Note 1*). Although sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) is a widely used method to observe crosslinked proteins, high molecular-weight proteins (>100 kDa) that are formed because of glycation are poorly resolved on polyacrylamide gels. These high molecular-weight proteins compress as a single band on the top of the gel and scan as a single peak by densitometric gel scanners. In order to quantitate glycation crosslinks, we have developed an *in vitro* assay method based on the incorporation of [<sup>14</sup>C]lysine into proteins during glycation (24). Because AGE inhibitors like aminoguanidine and semicarbazide inhibit the incorporation of [<sup>14</sup>C]lysine into proteins (24), this assay can be used to compare the crosslinking ability of various sugars, and it can also be used to test the ability of specific amino acids to act as donors for crosslinking to protein (*see Note 2*).

In this chapter, we describe a method to determine early glycation products by MS and also a method to quantitate the glycation crosslinks in model proteins by the incorporation of [<sup>14</sup>C]lysine into proteins.

## 2. Materials

### 2.1. Equipment

1. High-performance liquid chromatography (HPLC) system + ultraviolet (UV) detector.
2. Vydac C18 HPLC column (10 × 250 mm) (Vydac, The Separations Group, Hesperia, CA).
3. Nylon filters (Gelman Scientific, Ann Arbor, MI).
4. 1.5-mL sterile amber vials (Pierce Chemicals, Rockford, IL).
5. Metal ion-free Eppendorf tubes (Perfecto Scientific, Atascadero, CA).
6. FAB-MS (Varian, Netherlands).
7. ESI-MS (Varian).

### 2.2. Reagents

1. Diethylenetriaminepentaacetic acid (DTPA), [<sup>1-12</sup>C]L-threose and other biochemicals can be obtained from Sigma Chemical Co. (St. Louis, MO). [<sup>1-13</sup>C]L-threose can be obtained from Omicron Chemicals, South Bend, IN, and  $\alpha$ -N-t-Boc-lys-ala-ala peptide can be obtained from Bachem, Inc. (Philadelphia, PA).

2. Radioactive amino acids like [ $^{14}\text{C}$ ]lysine (specific activity 165 mCi/mmol) and [ $^{14}\text{C}$ ]leucine (specific activity 315 mCi/mmol) can be obtained from NEN Research Products (Boston, MA) or ICN Biomedicals Inc. (Costa Mesa, CA). Ultrapure lysozyme is from AMRESCO, Solon, OH. Model sugars like glucose, ascorbic acid, dehydroascorbic acid, or L-threose and RNase can be obtained from Sigma Chemical Co. Trichloroacetic acid (TCA) can be obtained from Fisher Scientific (Pittsburgh, PA).
3. Metal-free deionized water should be used to prepare buffers and other reagents.

### 2.3. Solutions

1. Phosphate-DTPA buffer: 0.1 M sodium phosphate buffer, pH 7.0, containing 0.1 mM DTPA.
2. Stock A: 1 M Boc-KAA peptide in phosphate -DTPA buffer.
3. Stock B: Threose in phosphate -DTPA buffer.
4. Solvent A: 0.1% TFA.

## 3. Methods

### 3.1. Glycation of *t*-Boc-lys-ala-ala (Boc-KAA) Peptide by Threose [ $^{12}\text{C}$ and $1\text{-}^{13}\text{C}$ ]

1. Prepare 0.1 M sodium phosphate buffer, pH 7.0 containing 0.1 mM DTPA as the chelator (referred as phosphate-DTPA buffer) in deionized water. Prepare stock solutions of 1 M Boc-KAA peptide and threose in phosphate -DTPA buffer just before use.
2. Prepare a reaction mixture containing peptide and threose (1:10 on a molar basis) by mixing 20  $\mu\text{L}$  of 1 M peptide and 200  $\mu\text{L}$  of 1 M threose in 1 mL of phosphate-DTPA buffer. Also prepare control reaction mixtures without peptide and without sugar in 1 mL of phosphate-DTPA buffer.
3. Immediately after the addition of reactants, filter the reaction mixtures through 0.2- $\mu\text{m}$  sterile nylon filters (Gelman Scientific) into sterile amber vials of 1.5-mL capacity (Pierce Chemicals). Use a laminar flow hood for this purpose.
4. Incubate the vials with reaction mixtures in an incubator maintained at 37°C.
5. To observe the reaction products that accumulate with time, withdraw aliquots (equivalent to approx 100  $\mu\text{g}$  peptide) at different intervals (0–72 h) (*see Note 3*) and freeze the aliquots immediately at  $-80^\circ\text{C}$ .

### 3.2. Preparation of Samples for Mass Spectrometry

1. Thaw aliquots frozen at  $-80^\circ\text{C}$  (**step 5, Subheading 3.1.**) at room temperature and inject on a semipreparative Vydac C18 RPHPLC column (10  $\times$  250 mm), which has been preequilibrated with 0.1%TFA in water (solvent A) (*see Note 4*). Set the absorbance on UV detector at 210 nm to monitor the absorbance of the peptide and reaction products.
2. Elute reaction products that are not bound to HPLC column in 5 column volumes of solvent A. This is followed by the elution of reaction products that are bound



to HPLC column in approx 10 column volumes of solvent B (60% acetonitrile and 0.1% TFA in water).

3. For accurate mass determination, collect the reaction products in metal-free (*see Note 5*) Eppendorf tubes (1.5-mL/tube/min) and combine the fractions with maximum absorbance at 210 nm (*see Note 6*).

### 3.3. FABMS Analysis of Modified Peptides

1. Dissolve the dried samples (**step 3, Subheading 3.2.**) in 5–10  $\mu\text{L}$  of methanol-water and add 90% formic acid (Spectrapure) to a final concentration of 22% (*see Note 7*). Remove a 1.0- $\mu\text{L}$  aliquot and mix immediately with a drop of a 1-thioglycerol and place on the MS instrument probe (*see Note 8*).
2. Insert the probe into a mass spectrometer and record spectra over a  $m/z$  range of 300–2000 at a resolution of 2000 (*see Notes 9 and 10*).

### 3.4. Electrospray Ionization Mass Spectrometry (ESIMS) of Boc-KAA Peptide Modified During Glycation

1. Dissolve the samples obtained from HPLC (**step 3, Subheading 3.2.**) in a solution of 50:50 acetonitrile:water with 0.1% TFA (*see Note 7*) and deliver to the analyzer at a flow rate of 5  $\mu\text{L}/\text{min}$  in the same solution (*see Note 11*).
2. Record the spectra over a  $m/z$  range 300–2000 and analyze the data with massLinX software or any other appropriate software.

### 3.5. Quantification of Glycation Crosslinks in Proteins by the Incorporation of [ $^{14}\text{C}$ ]Lysine into Proteins During Glycation

We describe below an assay method to determine the glycation-dependent crosslinks formed between [ $^{14}\text{C}$ ]lysine and lysozyme in the presence of a model sugar, L-threose (*see Note 12*). To measure the reactivity of the  $\alpha$ -amino group alone with lysozyme/RNase during glycation, [ $^{14}\text{C}$ ]leucine can be used instead of [ $^{14}\text{C}$ ]lysine in the reaction mixture. Adequate care should be taken to handle and dispose of any radioactive waste according to NRC guidelines.

1. Prepare 0.1 *M* sodium phosphate buffer, pH 7.0, containing 0.1 *mM* DTPA as the chelator (referred as phosphate-DTPA buffer) in deionized water. Just before use, prepare stock solutions of lysozyme or RNase (25 mg/mL) and L-threose or reactive sugar of choice (1 *M*) in phosphate-DTPA buffer.
2. Dilute the stock solution of [ $^{14}\text{C}$ ]lysine to a working stock of 50  $\mu\text{Ci}/100 \mu\text{L}$  in deionized water and keep on ice.
3. A typical 1-mL reaction mixture is prepared as described below (*see Note 13*).

|                              |                                                                |                                            |                                                |
|------------------------------|----------------------------------------------------------------|--------------------------------------------|------------------------------------------------|
| Lysozyme/RNase<br>(25 mg/mL) | [ $^{14}\text{C}$ ]lysine<br>50 $\mu\text{Ci}/100 \mu\text{L}$ | Phosphate-DTPA buffer<br>770 $\mu\text{L}$ | L-threose<br>(1 <i>M</i> )<br>20 $\mu\text{L}$ |
|------------------------------|----------------------------------------------------------------|--------------------------------------------|------------------------------------------------|

4. Immediately after the addition of L-threose, sterile filter each reaction mixture through a 0.2- $\mu\text{m}$  nylon filter into an amber vial (1.5-mL capacity) and incubate in an incubator maintained at 37°C.

5. Withdraw aliquots in duplicate at different intervals from 0–72 h and immediately freeze the aliquots at  $-80^{\circ}\text{C}$  until use.

### **3.6. Measurement of Radioactivity in the Aliquots Withdrawn from Reaction Mixtures**

Because free lysine readily adheres to proteins and methods like dialysis or size-exclusion chromatography do not completely remove lysine nonspecifically bound to proteins, we routinely used the filter disk method of Mans and Novelli (26) to determine the protein-bound radioactivity. According to this method, proteins will be precipitated by cold TCA on a 3 MM Whatman filter paper disk and any unbound or low molecular-weight compounds will be washed off prior to determine the protein-bound radioactivity. The proteins bind to the fiber of the filter disk and remain on the filter paper during the washing procedure with TCA.

1. Aliquots frozen at  $-80^{\circ}\text{C}$  should be thawed prior to use. Spot 10 mL of each aliquot (in triplicate) on 3 MM Whatman filter paper disks and air dry the filter paper disks under a fume hood (filter paper disks can conveniently be suspended with pins on a Styrofoam box lid to air dry).
2. After the reagents have been absorbed into the disks, drop the filter paper disks with pins still inserted into a solution of ice-cold 10% TCA while stirring with a magnetic stirrer and wait 20 min (*see Note 14*). A perforated 250-mL plastic beaker (a cork borer can be used to make holes in the plastic beaker) can be used to hold the filter paper disks and place the plastic beaker in a 500-mL glass beaker containing 10% TCA.
3. Transfer the plastic beaker with filter paper disks to a glass beaker containing 5% TCA (room temperature) and stir for 20 min. Following this, transfer the plastic beaker with filter paper disks to a glass beaker containing 5% TCA maintained at about  $75^{\circ}\text{C}$  and stir for 15 min. Hot TCA assures the measurement of incorporation into protein only.
4. Transfer the plastic beaker with filter paper disks to a mixture of ethanol-anhydrous ether (2:1) under stirring and wait for 15 min. This step is required to remove residual TCA from the filter paper disks.
5. Transfer the filter paper disks carefully to a plastic weighing boat or to a sheet of aluminium foil and air-dry the filter paper disks in the fume hood or under a heat lamp.
6. Remove the pins, transfer the dried filter paper disks to a vial containing radioactive scintillation cocktail (e.g., Ultima gold, Packard, Meriden, CT) and count radioactivity in a liquid scintillation counter. Calculate the amount of radioactivity present on each disk and determine the amount incorporated into lysozyme or RNase at different intervals during glycation reaction.

## **4. Notes**

1. Determination of molecular weights of glycosylated peptides by either fast atom bombardment mass spectrometry (FABMS) or by electrospray ionization mass spec-

trometry (ESIMS) is an effective method of establishing whether a peptide has become glycated, how many sugar moieties are attached to the peptide, and whether there is evidence of crosslinks. Typical mass accuracy for determinations by both FABMS and ESIMS is about  $\pm 0.2$  Daltons. Although both techniques can be used, each has its advantages and disadvantages. FABMS is customarily used for peptides with molecular masses of 250–4000. The mass range for ESIMS is much larger. ESIMS can be used to determine molecular masses of both peptides and proteins.

2. Although [ $^{14}\text{C}$ ]lysine can be used to measure glycation dependent crosslinking of [ $^{14}\text{C}$ ]lysine to model proteins, N- $\alpha$ -acetyl[ $^{14}\text{C}$ ]lysine can be used to estimate the glycation crosslinks formed because of the specific interaction of  $\epsilon$ -amino group of lysine with proteins during glycation (27). To measure specific lysine-amino acid crosslinks, homopolymers of amino acids like polylysine, polyarginine or polyhistidine can also be used.
3. L-threose is a highly reactive tetrose sugar and its reaction with free amino groups is generally completed in only a few days (25). Therefore, L-threose is selected as a model sugar in the experiments. If threose is substituted with sugars like glucose or fructose, aliquots should be withdrawn at weekly intervals or until the reaction products reach to a plateau.
4. FABMS or ESIMS require that the sample should be free of nonvolatile salts containing cations such as sodium and potassium. These cations can replace the protons on some peptides, leading to many additional charged states. For this reason, it is recommended that the last step in the isolation of the peptides be RP-HPLC. Dialysis usually does not adequately remove interfering salts. Because of the extreme sensitivity to sodium ions, samples should be collected and stored in plastic tubes after the final RP desalting.
5. Because sodium ion impurities in samples injected on MS can lead to inaccurate mass spectra, care should be taken to select metal ion-free Eppendorf tubes. We routinely used plastic tubes obtained from Perfector Scientific, Atascadero, CA.
6. The most important information for someone submitting samples for mass spectrometric analysis is how much sample is required for analysis, and that the samples should not contain too many components and be free of salts.
7. For both FABMS and ESIMS, the sample is usually dissolved in an acidic solvent, such as 0.1% TFA or acetic acid before introduction into the mass spectrometer.
8. FABMS analysis is performed by mixing the dissolved sample with a matrix such as glycerol/thioglycerol or nitrobenzyl alcohol, and placing this mixture on the probe that is inserted into the instrument. The peaks caused by the matrix contribute to higher background signals, making this technique less sensitive than ESIMS, particularly at masses below 300 Daltons.
9. We have used a range between 300–2000, because the peptide we used in this study has a molecular mass of 389. An appropriate range should be selected based on the size and mass of protein or peptide in use.
10. One advantage of FABMS is that, in the positive ion mode, a monoprotonated peptide is obtained, so the spectrum shows an  $\text{MH}^+$  ion, which is the molecular

mass of the peptide plus one Dalton. The molecular mass of the peptide is readily apparent. In contrast, ionization of the sample in ESIMS produces several charged states of each peptide, resulting in a spectrum showing peaks with the mass/charge ratio of each of the charged states.

11. For ESIMS, the dissolved sample is introduced into the mass spectrometer as a continuous flow of solution. Although the amount of sample required depends very much on the sample characteristics, about one nanomole of peptide is typical for a FABMS analysis, whereas 100 picomole is usually sufficient for ESIMS. Detection limits 1000-fold lower may be achieved in special cases.
12. One can also use radioactive-labeled proteins instead of [ $^{14}\text{C}$ ]lysine to measure the incorporation of labeled sugars into proteins.
13. Always set up a reaction without sugar to correct for any nonspecific interaction of [ $^{14}\text{C}$ ]lysine with proteins. Also minimize the volume (50–100  $\mu\text{L}$ ) of samples with [ $^{14}\text{C}$ ]lysine to apply on the filter paper disks.
14. To show that no carryover of [ $^{14}\text{C}$ ]lysine from one disk to others in between washings of the disks, include plain filter paper disks during the TCA wash procedure.

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