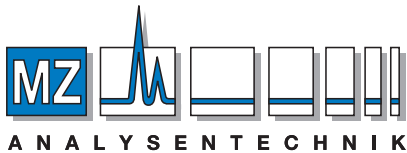




# Shimadzu HPLC Columns

Shimadzu  
High Performance Liquid Chromatograph



**AUTHORIZED DISTRIBUTOR**

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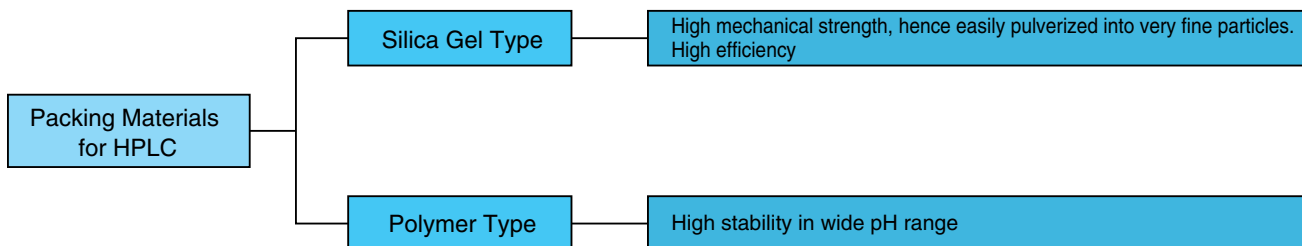
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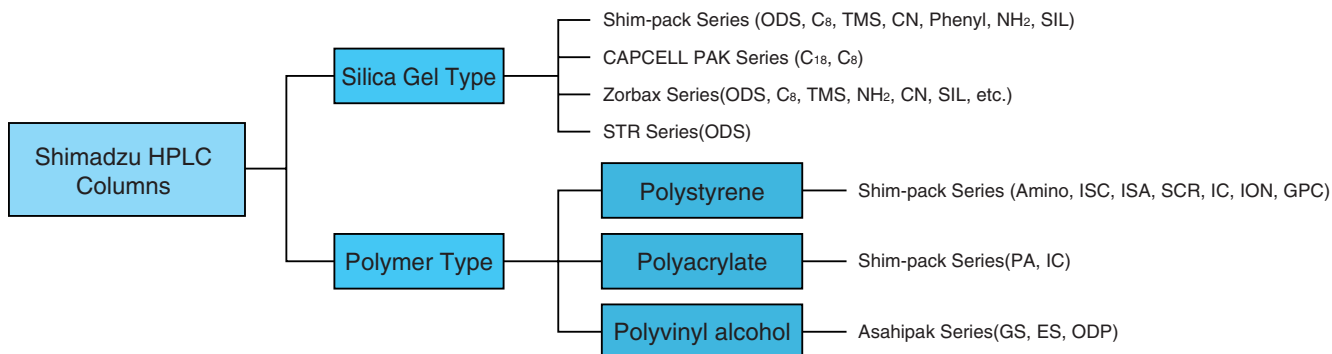
# 1 Classification of Shimadzu HPLC Columns

## Classification by Support Material

The solid supports of packing materials for HPLC are classified into the following two groups with the following features.

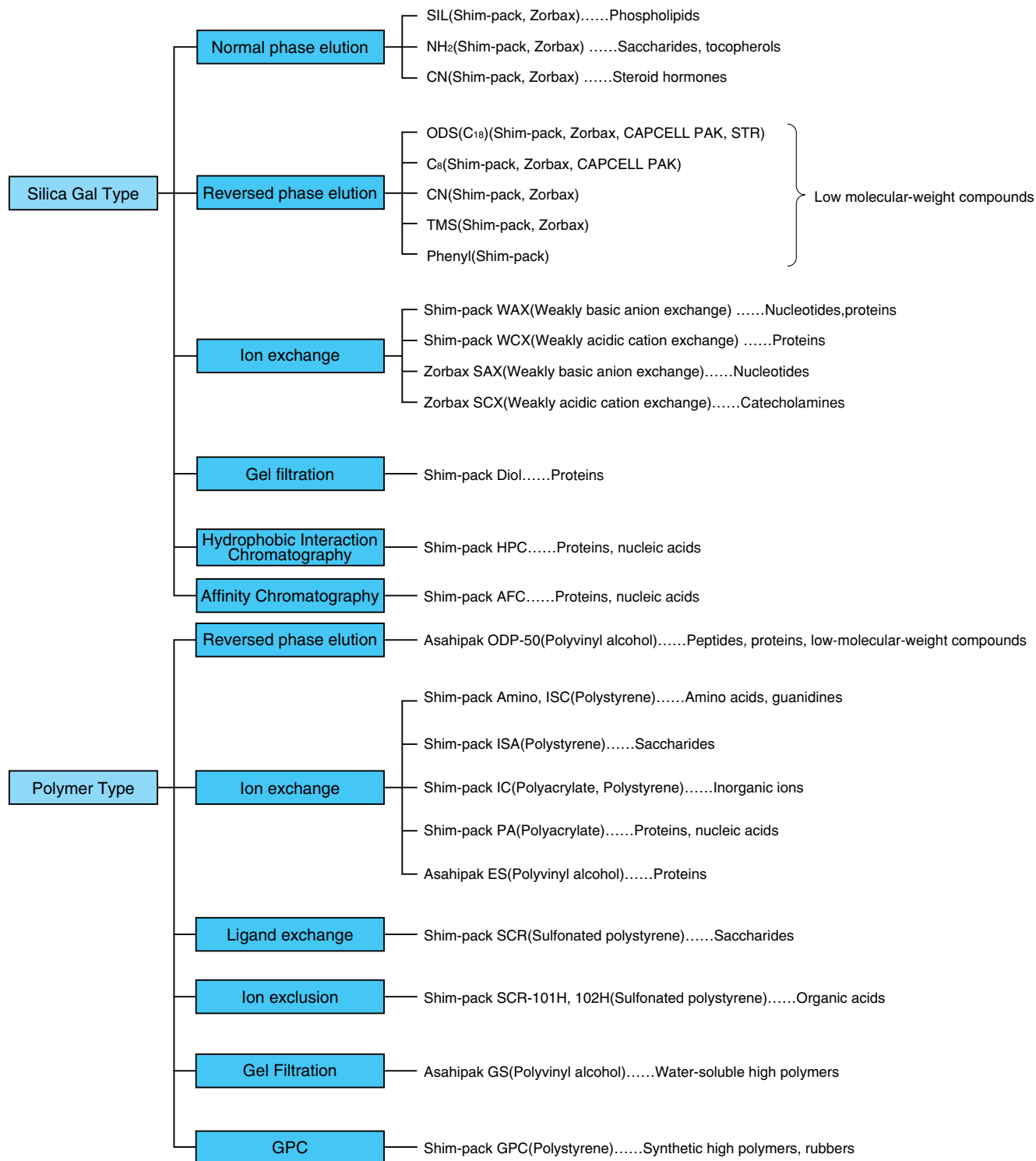


The Shimadzu HPLC columns can be classified into two groups as follows:



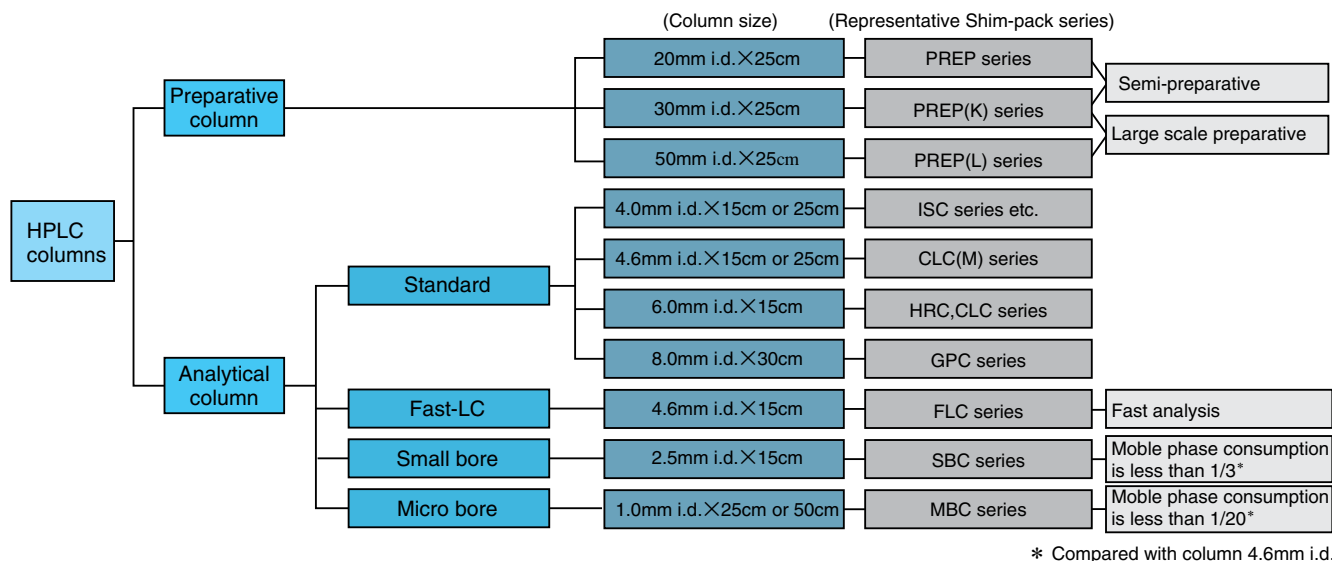
## Classification by Elution Mode

● The two groups are further classified as follows:



## Classification by Column Dimensions

- HPLC columns are classified as analytical columns and preparative columns according to the dimensions.
- Analytical columns are further classified into several groups.



- You can choose 20mm i.d., 30mm i.d., 50mm i.d. preparative columns in accordance with the scale of preparative. Each is 25cm long.
- As standard size analytical columns, 4~8mm i.d. and 15~30cm long columns are most widely used and are available in a wide variety.
- As other special sizes, analytical columns such as Fast-LC columns, Micro bore columns and small bore columns are available. (Refer to P.26)

### Micro bore columns

- Shimadzu MBC series micro bore columns are only 1.0mm in inner diameter.
- The mobile phase consumption is only about 1/20 that required in a 4.6mm i.d. column, because the same linear velocity can be obtained with about 1/20 flow rate.
- The sample injector, the detector flow-thru cell, and connecting pipes of a small capacity must be used.

### Small bore columns

- Shimadzu SBC series small bore columns are 2.5mm in inner diameter.
- The mobile phase consumption is about 1/3 compared with a 4.6mm i.d. column.
- Though this series columns are compatible with an ordinary HPLC, it is better to use small capacity units.

### Fast-LC columns

- Shimadzu FLC series fast LC columns are packed with 3 μm dia.
- Column packing is only 5cm long.
- They are compatible with an ordinary HPLC and permit very fast analysis.

## Classification by ODS Columns

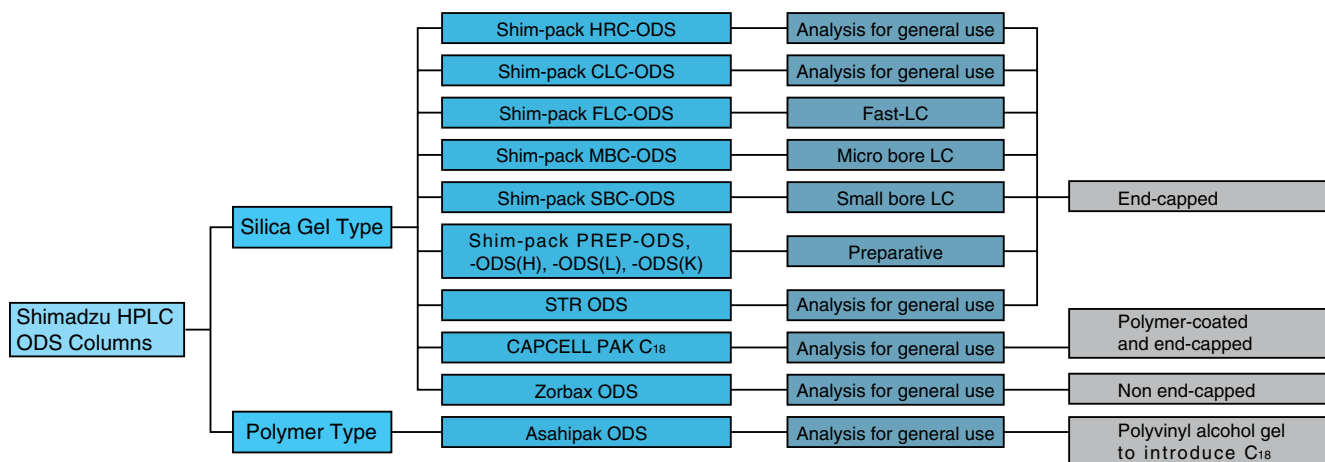
### ODS columns

ODS columns are packed with packing material having octadecyl group chemically-bonded on the surface and are most popularly used in HPLC.

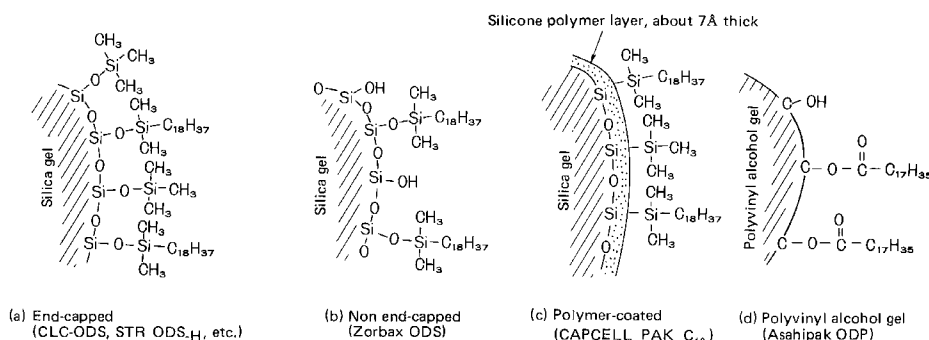
ODS columns can have different separation properties, according to the state of end-capping and the characteristics of the silica gel used as the solid support. They can also be different with the brand.

The delicate differences in separation properties can be utilized to achieve some difficult analysis.

Shimadzu provides various types of ODS columns to ensure a wide field of application.



### Method to Introduce C<sub>18</sub> Groups



### Selection of ODS Columns

As standard analytical columns, we provide Shim-pack HRC-ODS, CLC-ODS, STR ODS, Zorbax ODS, CAPCELL PAC C<sub>18</sub>, Asahipak ODP.

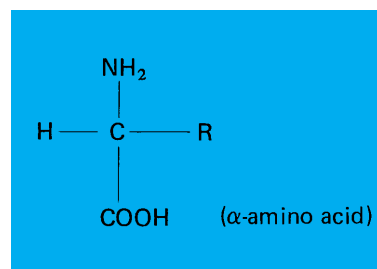
The following table is a rough guide for selecting ODS columns.

Column name	Features	Mobile phase pH range
Shim-pack HRC-ODS Shim-pack CLC-ODS STR ODS	End-capped to minimize the influence of residual silanol groups and eliminate tailing of basic compounds.	2~7.5
Zorbax ODS	Utilizes the influence of residual silanol compounds.	2~7.5
CAPCELL PAK C <sub>18</sub>	Silica gel solid support is coated with silicone polymer to improve stability in basic mobile phase.	2~10
Asahipak ODP-50	Applicable to mobile phase up to pH13. The selectivity is different from that of silica gel types.	2~13

## 2 Guide for Selection of Operational Conditions

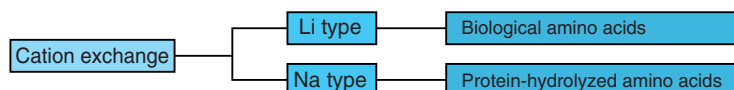
### Amino Acids

Amino acids are amphoteric compounds having carboxyl groups and amino groups. Amino acids are classified as acidic, neutral, and basic according to the number of these groups. Also they are classified as aliphatic type and aromatic type, depending on the group at the position of R.



#### Separation mode

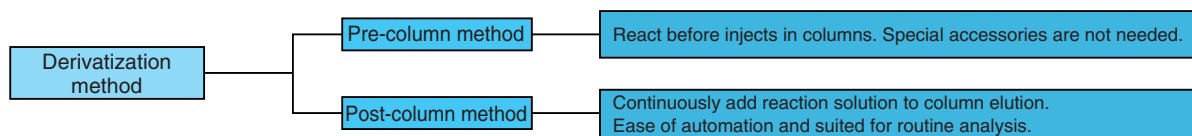
- The cation exchange chromatography is most popular.
- It is classified into two methods, Li type and Na type.



- The Li type is recommended for analysis of biological amino acids. It is also used for separating glutamine and glutamic acid.
- Reversed phase elution mode can separate particular type of amino acids but cannot separate all free amino acids.
- It is quite effective for separating amino acids derivatized by precolumn derivatization.

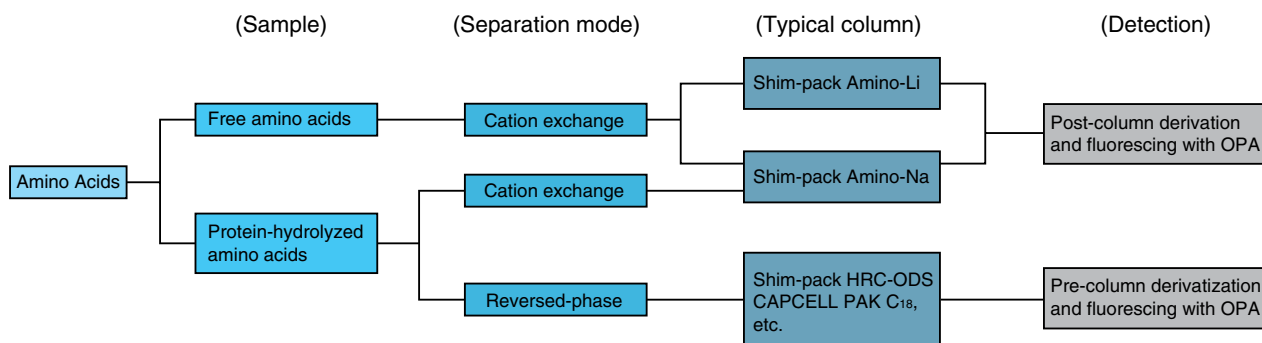
#### Detection Method

- The UV absorption method which utilizes the absorption of carboxyl groups at 200~210nm is not widely used though it is useful for detecting some type of amino acids.
- Pre-column derivatization and post-column derivatization methods are often used for detecting amino acids.



- Pre-column derivatization method:  
The sample solution is derivatized before entering the analytical column. Combination of the pre-column derivatization method and the reversed phase elution mode provides rapid analysis of amino acids produced in hydrolysis of proteins. The reagents for pre-column derivatization such as OPA(ortho-phthalaldehyde) are commercially available.
- Post-column derivatization method:  
The derivatizing reagent is continuously added to the column effluent. This method has the noteworthy advantage that the entire operation can be easily automatized. Combination of the post-column derivatization method and the cation exchange method permits analysis of wide variety of amino acids ranging from free amino acids to amino acids produced in hydrolysis of proteins. The representative reagents used in the post-column derivation method are OPA (for fluorescing) and ninhydrin.

#### Column



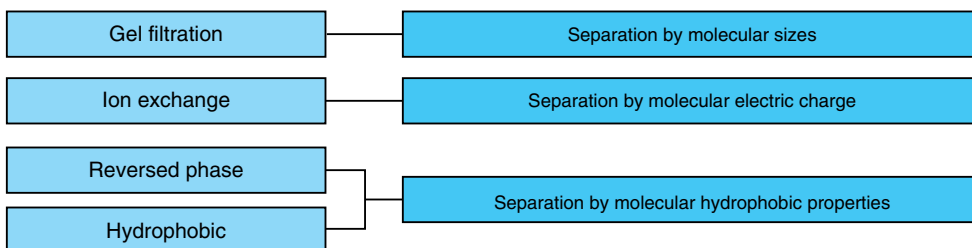


## Peptides and Proteins

Peptides and proteins consist of amino acids which are combined by acid-amide bonding. Their ionic and hydrophobic properties are different depending on the type and the number of constituting amino acids. Proteins have high-degree structures with various combinations.

### Separation mode

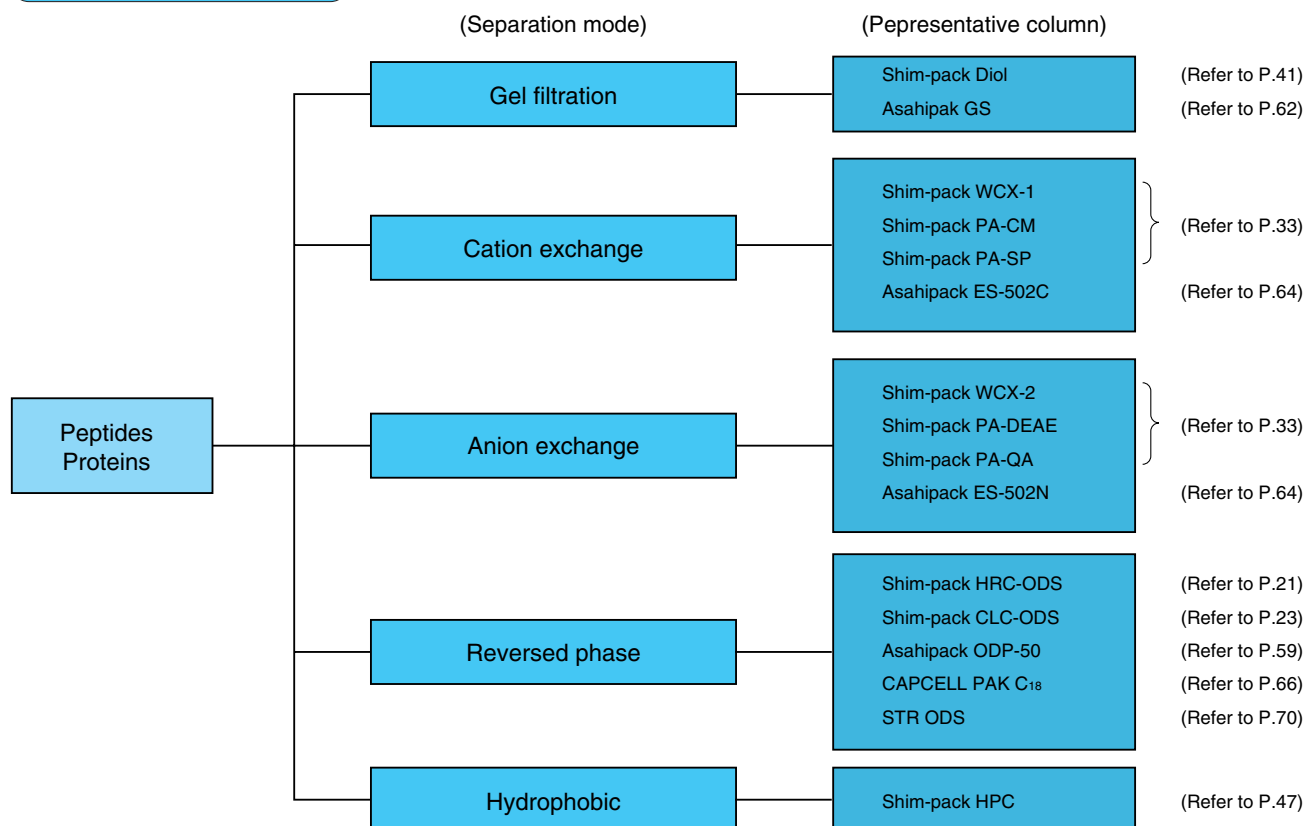
- Peptides and proteins are generally analyzed by one of the following methods.



### Detection Method

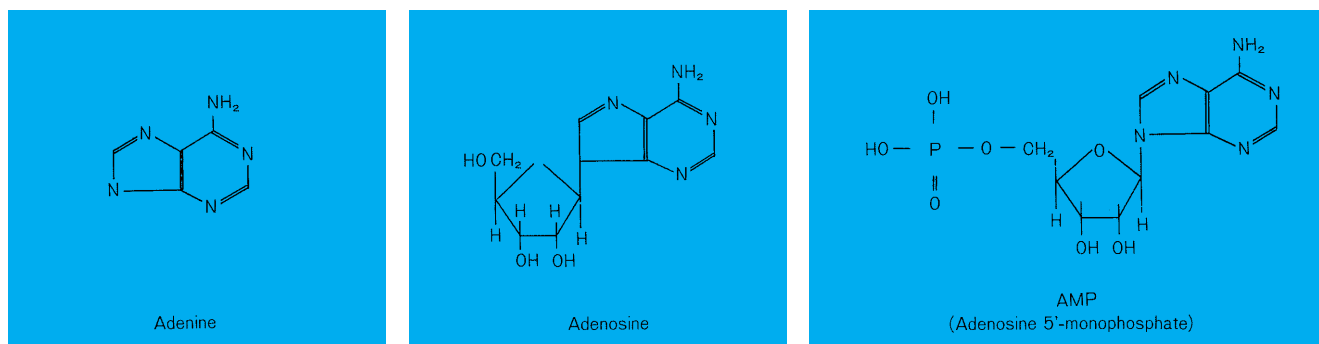
- The UV spectrophotometric method is generally used. Detection at 280nm is effective in most cases.
- Detection at about 210nm is used for detection of peptides consisting of aliphatic amino acids and detection of low-concentration peptides and proteins.

### Column



## Nucleic Acids

Nucleic acids play important roles in organism. They are classified into the following three groups:



Nucleic acid base

Nucleoside  
(Nucleic acid base + sugar)

Nucleotide  
(Nucleoside + phosphorus)

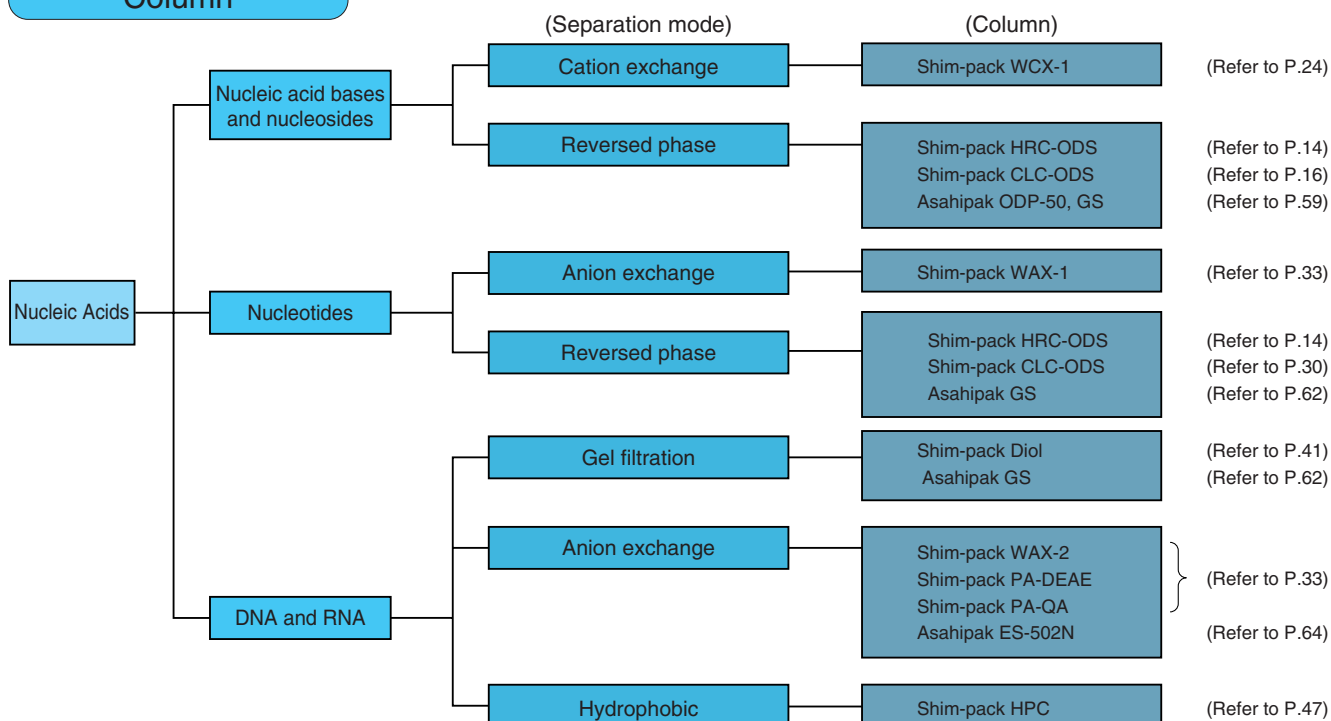
### Separation mode

- Nucleic acid bases and nucleosides are analyzed in the cation exchange mode or the reversed phase elution mode.
- Nucleotides are analyzed in the anion exchange mode in most cases. The reversed phase elution is also useful with suitable selection of mobile phases.
- In analysis of high molecular weight samples such as oligonucleotides, DNA, and RNA, gel filtration mode, hydrophobic mode, and ion exchange mode are selectively used.

### Detection Method

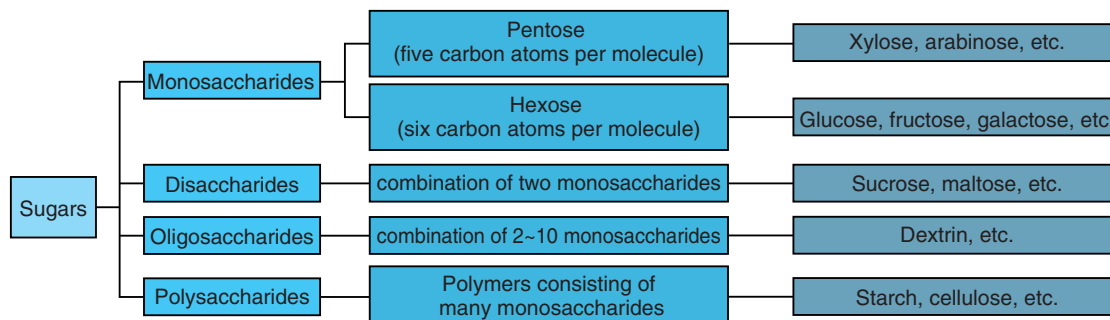
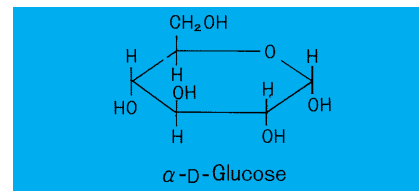
- Nucleic acids and related compounds generally have strong ultraviolet absorption, therefore UV spectrophotometric detection (250~260nm) is used in most cases.

### Column



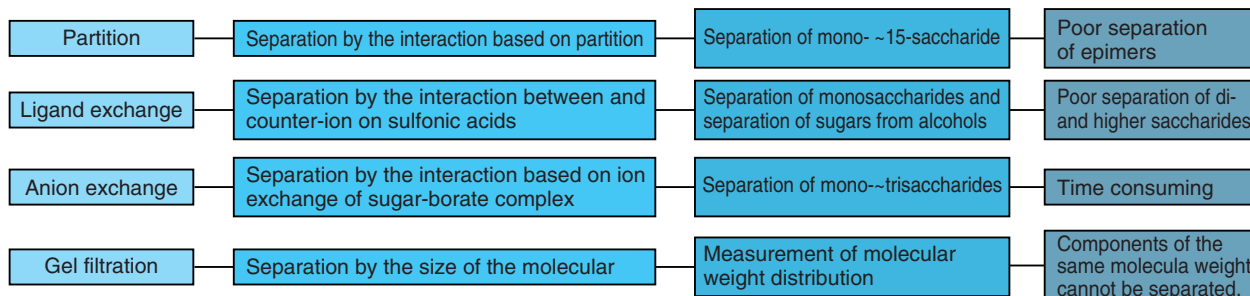
# Sugars

Sugars such as glucose and sucrose are very common and are available in a great variety.



## Separation mode

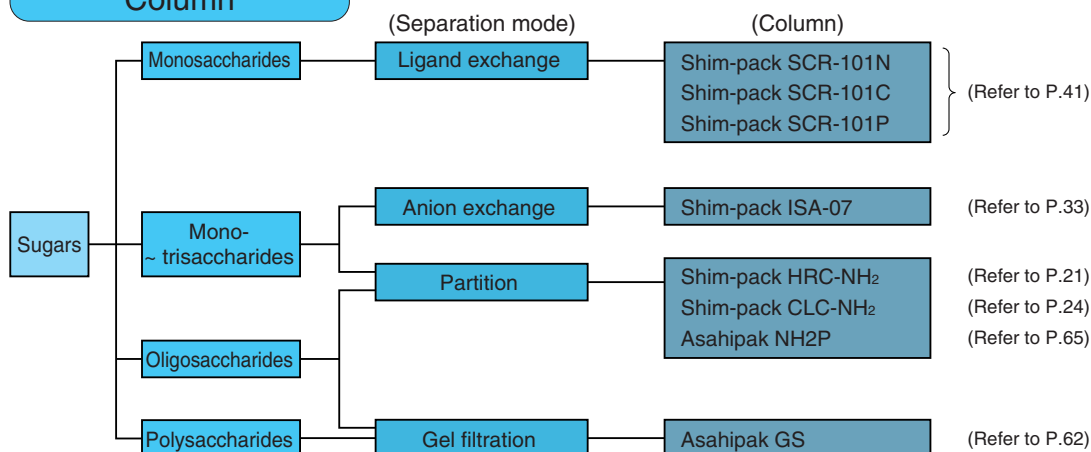
● Sugars are generally analyzed by one of the following methods.



## Detection Method

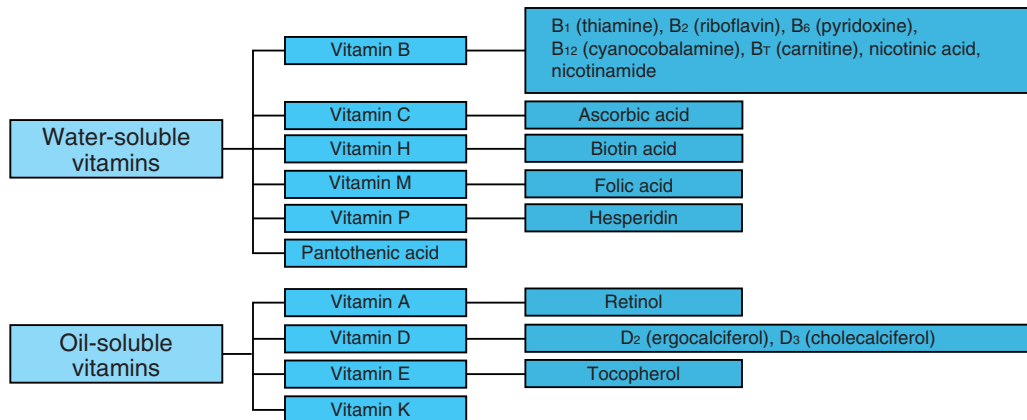
- A UV spectrophotometric detector is not used because sugars absorb radiation only around 190nm.
- A refractive index detector is generally used in the analysis of sugars.
- When sugar content is low and impurity content is high, as in the case of brewage products and natural materials, the post-column derivatization method is used.

## Column



# Vitamins

Vitamins are classified as water-soluble type and oil-soluble type.

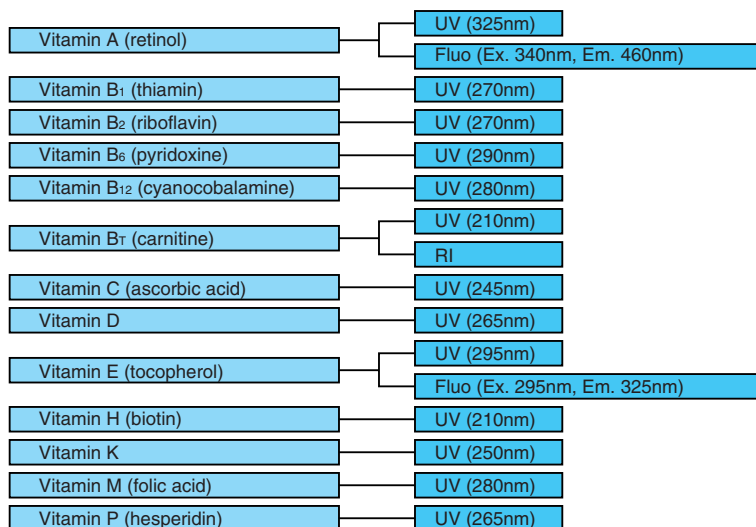


## Separation mode

- Both water-soluble and oil-soluble vitamins are analyzed in the reversed phase elution mode.
- Some oil-soluble vitamins are analyzed in the partition mode.

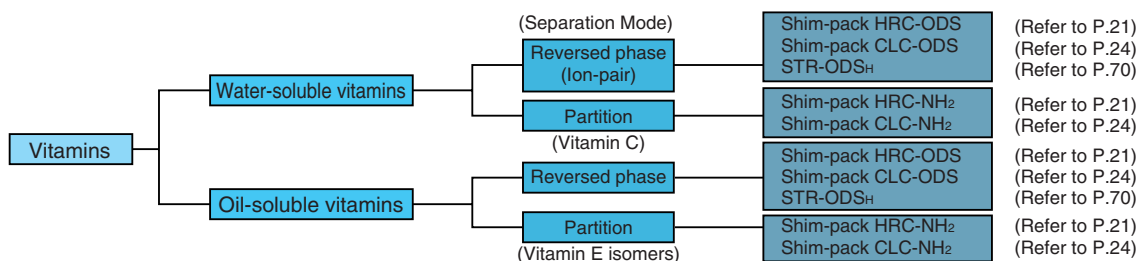
## Detection Method

- The UV spectrophotometric method is generally used: the spectrofluorometric method is used for some vitamins.



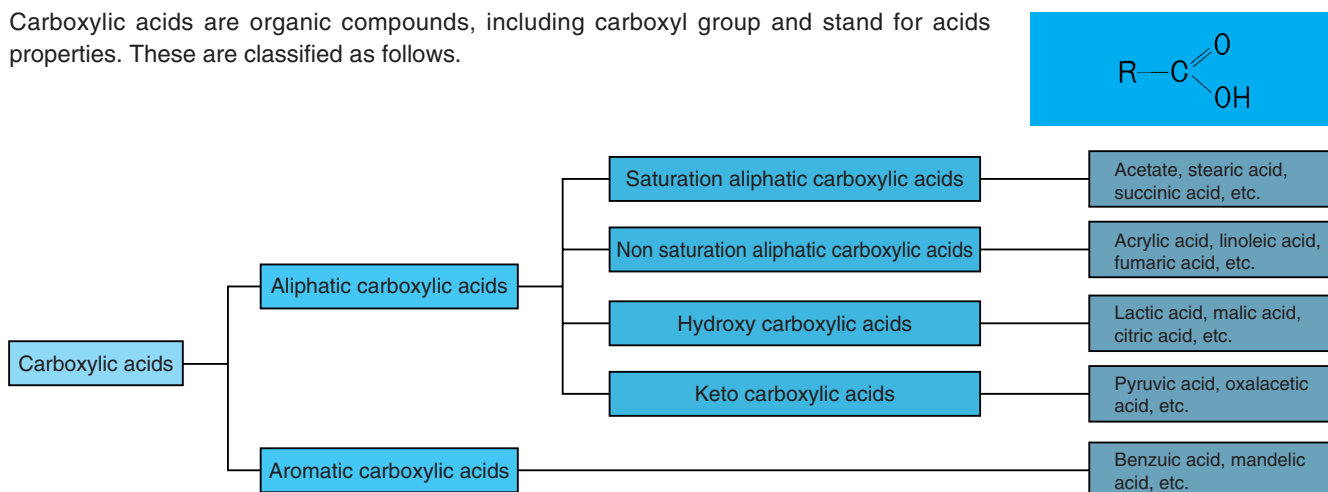
Note: The above wavelength values are approximate; they can be influenced by the state of mobile phase.

## Column



## Carboxylic acids

Carboxylic acids are organic compounds, including carboxyl group and stand for acids properties. These are classified as follows.



Carboxylic acids are also classified into mono, di, poly (more than 3) carboxylic acids in accordance with the number of carboxyl group.

### Separation mode

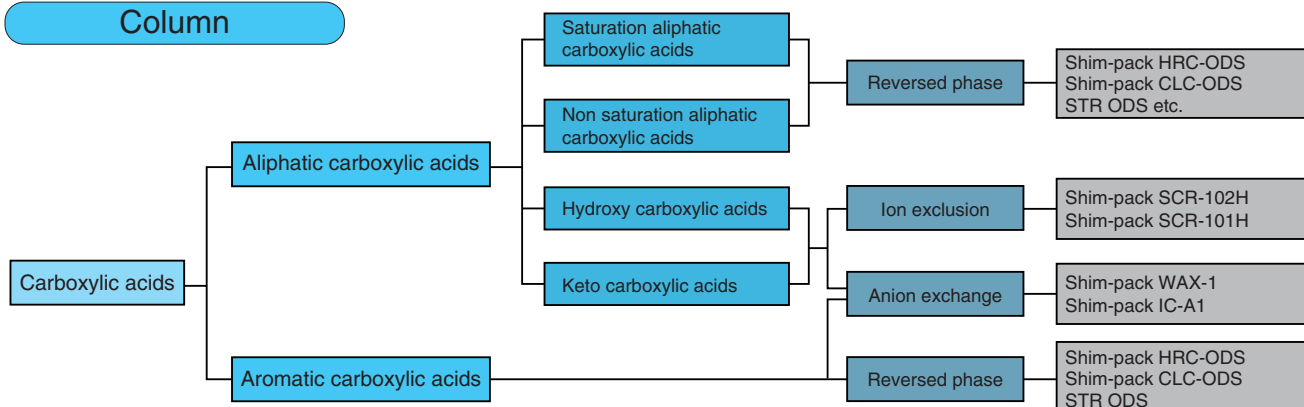
Ion exclusion chromatography, anion ion exchange chromatography and reversed phase chromatography can be used in accordance with the dissociation and the side chain of carboxylic acid.

- Ion exclusion chromatography is suitable for the analysis of high aqueous carboxylic acids and used widely to analyze food and cultivation solutions.
- Anion exchange chromatography is a basic mode of carboxylic acids having the properties of acids and has an advantage of separating from neutral components.
- Reversed phase chromatography is suitable for the separation of fatty acid and aromatic carboxylic acids.

### Detection Method

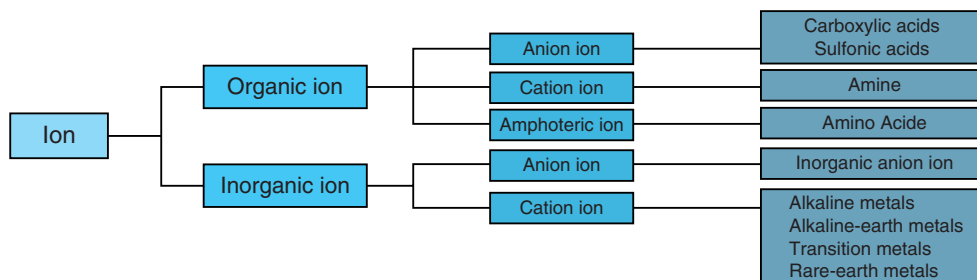
- Carboxylic acids have slight absorbency at around 205nm, therefore UV spectrophotometric detector can be used. As far as aromatic carboxylic acids and aliphatic carboxylic acids are concerned, these absorbencies are relatively strong so that can be detected with a high sensitivity.
- However, the absorbency coefficients of carboxylic acids are generally small and the selectivity of the detection are poor. As a result, there are cases where it is difficult to analyze some samples. Therefore, pre-column derivatization using label to carboxyl group might be used.
- Post-column conductivity detector making use of separating carboxyl group makes it possible to improve selectivity.

### Column



# Ion

Ions are classified as follows. Mainly inorganic ion and low molecule organic ion are the object of ion chromatography analysis.

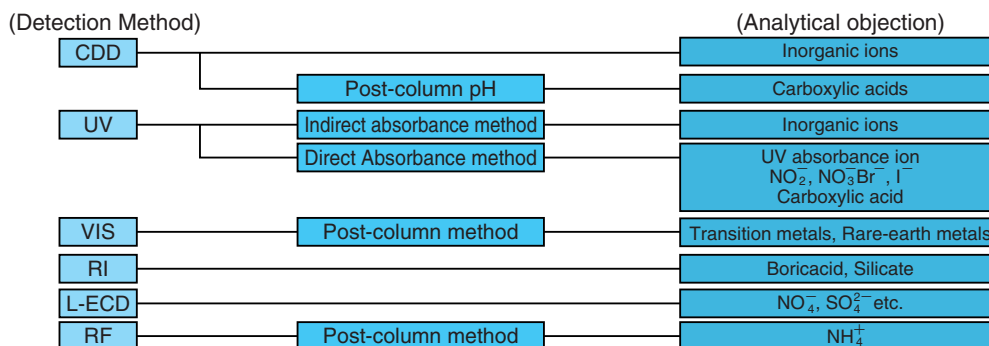


## Separation mode

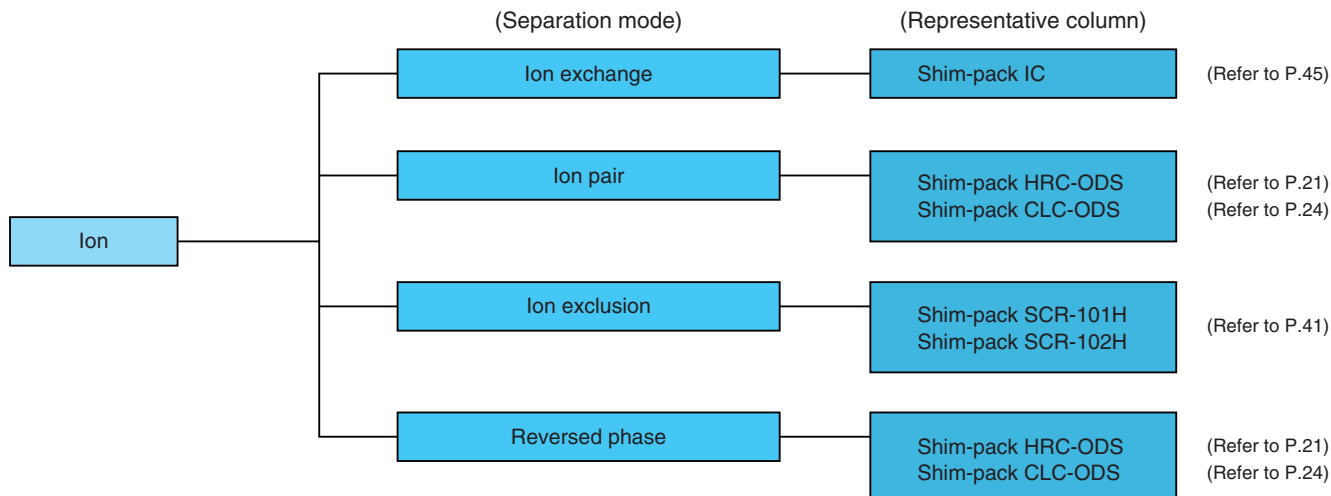
Ion exchange is a main method, but sometimes ion pair and reversed phase method are used.

## Detection Method

- In IC (Ion chromatography) analysis, generally conductivity detector can be used with a high sensitive.
- In special ions, many selectable detector can be used to make use of each chemical property.



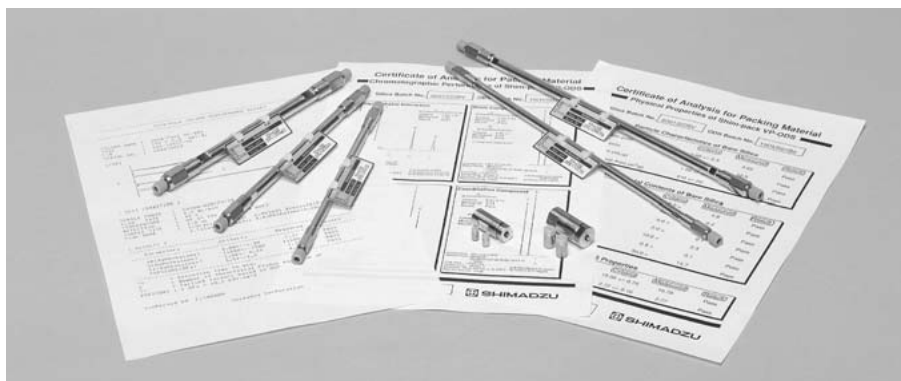
## Column



# 3 Shimadzu HPLC Columns

## Shim-pack VP-ODS for GLP/GMP compliance

For development or validation of analytical method, manufacturing uniformity of columns is increasing. Shim-pack VP-ODS has been developed to meet such expectations.



### 〈VP-ODS Series〉

#### Analytical Column Shim-pack VP-ODS

Cat. No.	I.D.×length (mm)
228-34937-91	4.6×150
228-34937-92	4.6×250
228-34937-93	6.0×150
228-34937-94	2.0×150
228-34937-95	2.0×250
228-34937-96	4.6×150, a set of 3pcs.
228-34937-97	2.0×150, a set of 3pcs.

\* An analytical column set consists of three columns whose packings (Silica-based) are from different production batches.

#### Guard Column Shim-pack VP-ODS

Cat. No.	I.D.×length (mm)
228-34938-91	4.6×10 Incl. exchangeable cartridge 2pcs.
228-34938-92	A holder for I.D. 4.6mm
228-34938-93	2.0×5 Incl. exchangeable cartridge 2pcs.
228-34938-94	A holder for I.D. 2.0mm

\* Guard column consists of a cartridge and holder. Holders can be used repeatedly.

#### Short Columns for Analytical/Preparative

Cat. No.	I.D.×length (mm)	Remarks
228-36849-91	4.6×50	5 μm diameter,
228-36849-92	20×50	12nm (120Å) pore
228-36849-93	20×100	porous silica ODS

### Excellent manufacturing uniformity

To minimize column-to-column performance deviation of ODS columns, silica-bases, surface treatment and packing procedures are strictly controlled respectively and only the products that passed the quality criteria are delivered to customers. For development or validation of method, it would be efficient to run the test with a set of three columns with packings of different batches.

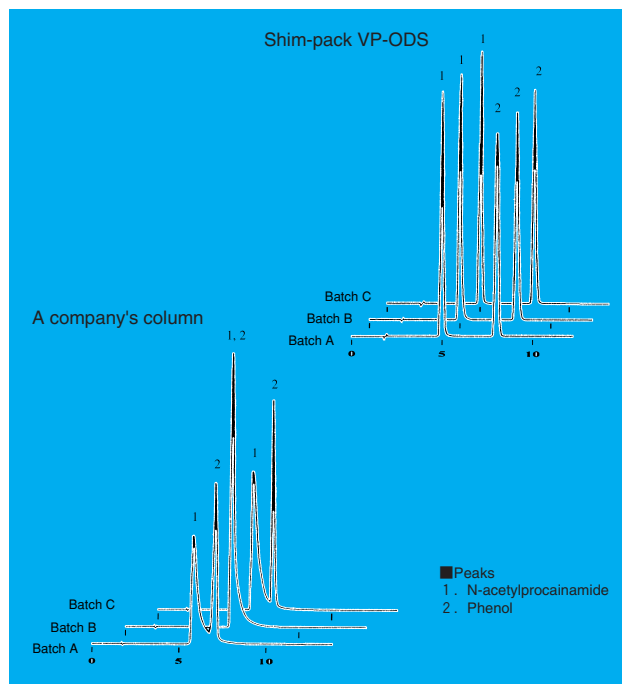
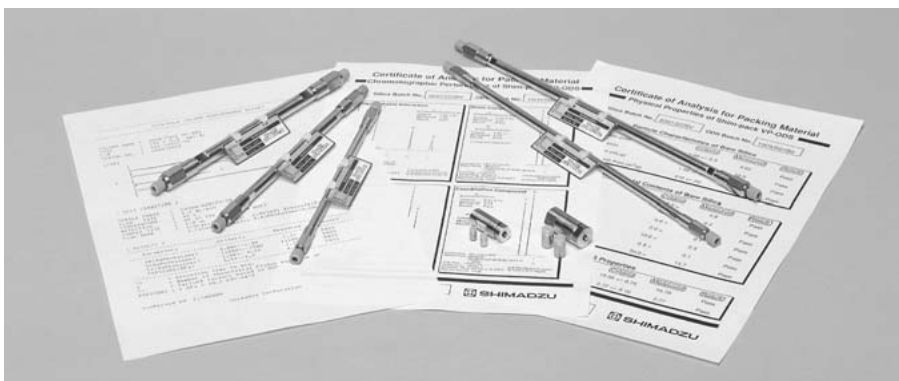


Fig. 1 Comparison of lot-to-lot repeatability (Silica-based material)

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228-34938-93	2.0×5 Incl. exchangeable cartridge 2pcs.
228-34938-94	A holder for I.D. 2.0mm

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228-36849-92	20×50	12nm (120Å) pore
228-36849-93	20×100	porous silica ODS

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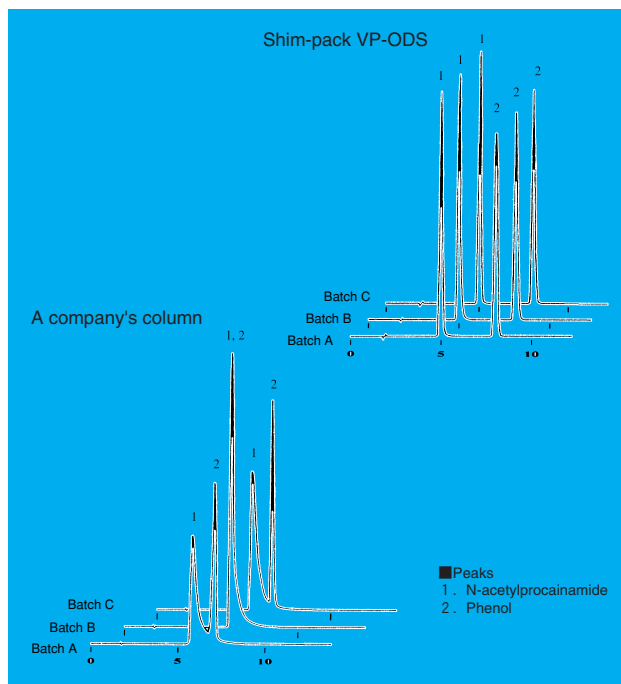
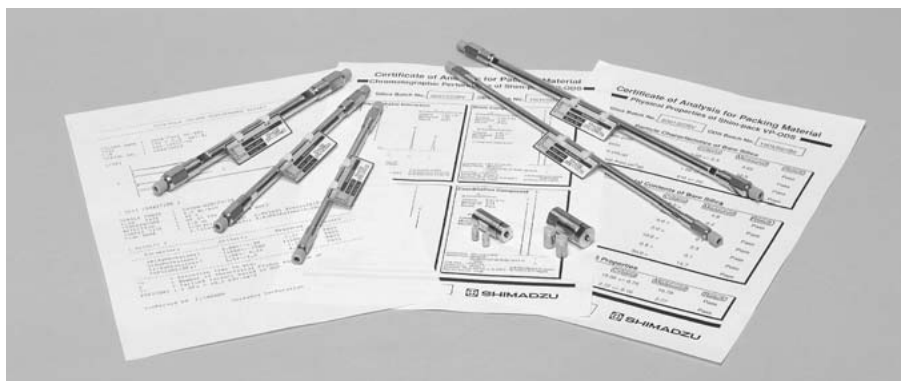


Fig. 1 Comparison of lot-to-lot repeatability (Silica-based material)



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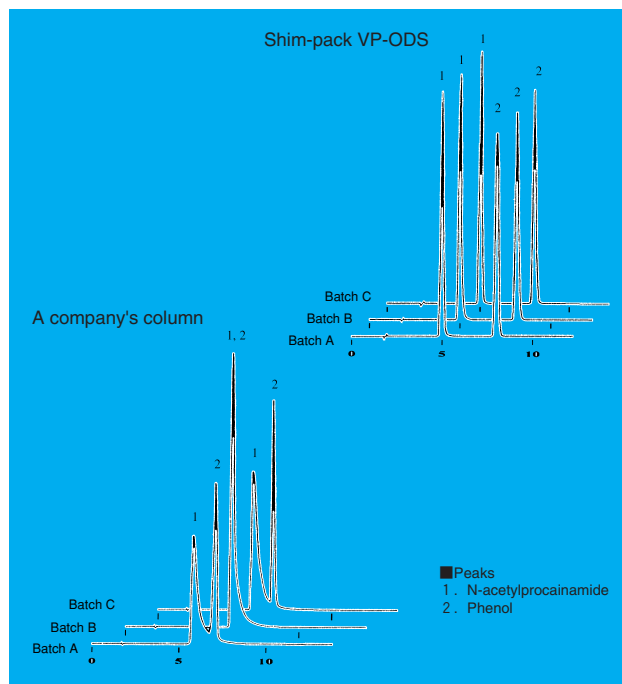


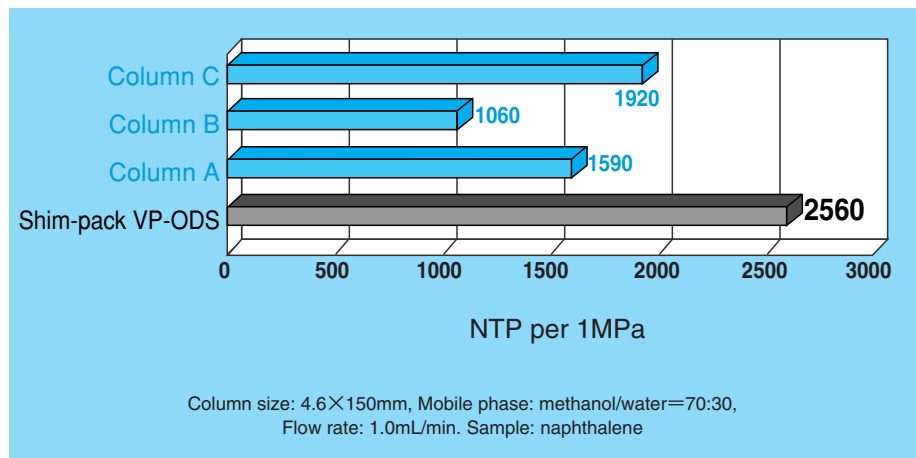
Fig. 1 Comparison of lot-to-lot repeatability (Silica-based material)

## 〈Packing characteristics of Shim-pack VP-ODS〉

Silica particles	Entirely porous, spherical silica particles with a high purity
Particle size	5 $\mu$ m
Pore size	12nm
Pore volume	1.25mL/g
Specific surface area	410m <sup>2</sup> /g
Trace metal content	30ppm max
Percentage of ODS	20%C
Carbon content	
ODS functional group	Mono-functional
End-capping	Used

## 〈Number of theoretical plates and column pressure〉

The higher NTP is and the lower the pressure is, the easier to handle the column is. Shim-pack VP-ODS shows superior performance shown by NTP per 1MPa.



## 〈Excellent peak shapes〉

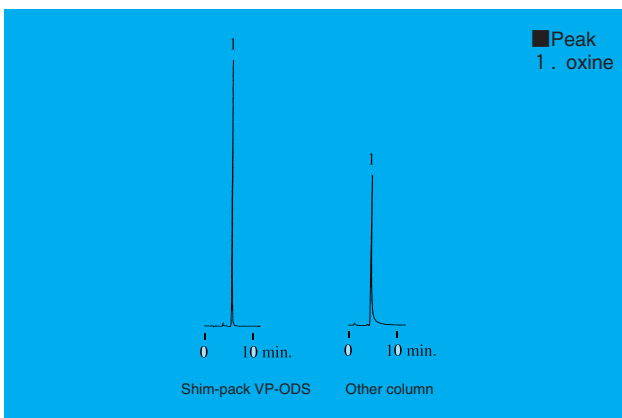


Fig. 2

Excellent peak shape of coordination compounds is achieved by the packing base material with less metal impurities.

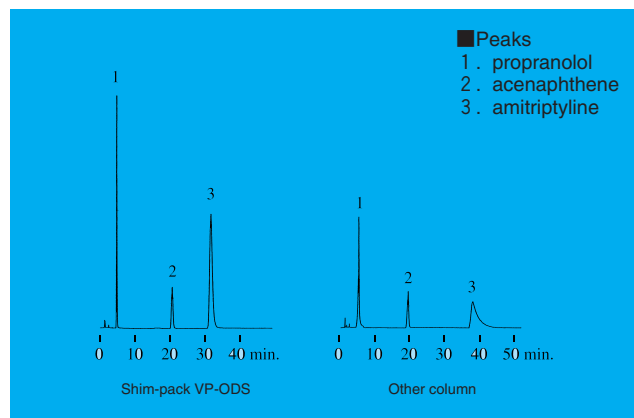


Fig. 3

Excellent peak shapes of basic compounds are achieved by thorough effective end-capping of the packings.

〈Certificate of Analysis (Quality certificate)〉

Shim-pack VP-ODS is shipped with three types of certificate of Analysis to support method validation for HPLC analysis.

1. Certificate of Analysis for Packing Material (Physical Properties)
2. Certificate of Analysis for Packing Material (Chromatographic Performance)
3. Shim-pack Column Performance Report

**Certificate of Analysis for Packing Material**  
Physical Properties of Shim-pack VP-ODS

Silica Batch No. S051222BV ODS Batch No. 19DVS01B0

**Particle Characteristics of Bare Silica**

	Criteria	Measured	Result
Particle Size (µm)	4.60 ± 0.3	4.65	Pass
Pore Size (nm)	12.0 ± 1.0	12.5	Pass
Pore Volume (mL/g)	1.25 ± 0.05	1.28	Pass
Specific Surface Area (m <sup>2</sup> /g)	410 ± 20	409	Pass

**Trace Metal Contents of Bare Silica**

	Criteria	Measured	Result
Na (ppm)	10.0 >	4.9	Pass
Cu (ppm)	10.0 >	6.2	Pass
Mg (ppm)	2.0 >	1.1	Pass
Al (ppm)	2.0 >	0.1	Pass
Fe (ppm)	10.0 >	2.3	Pass
Ti (ppm)	0.5 >	0.1	Pass
Total (ppm)	30.0 >	14.7	Pass

**ODS Properties**

	Criteria	Measured	Result
Total Carbon Content (%)	19.58 ± 0.75	19.79	Pass
ODS Coverage (µmol/m <sup>2</sup> )	2.72 ± 0.10	2.77	Pass

Verified by [ ] SHIMADZU

**Certificate of Analysis for Packing Material**  
Chromatographic Performance of Shim-pack VP-ODS

Silica Batch No. S051222BV ODS Batch No. 19DVS01B0

**Hydrophobic Interaction**

Column: 150 x 4.6 mm ID, 5 µm, HiChrom VP-ODS  
Mobile Phase: Acetonitrile  
Flow Rate: 1.0 mL/min  
Temperature: 40.0 °C  
Injection Volume: 10 µL  
Sample Volume: 10 µL

Criteria: 1.50-1.80  
Measured: 1.51  
Result: Pass

**Basic Compound**

Column: 150 x 4.6 mm ID, 5 µm, HiChrom VP-ODS  
Mobile Phase: Acetonitrile  
Flow Rate: 1.0 mL/min  
Temperature: 40.0 °C  
Injection Volume: 10 µL  
Sample Volume: 10 µL

Criteria: 1.50-1.80  
Measured: 1.54  
Result: Pass

**Acidic Compound**

Column: 150 x 4.6 mm ID, 5 µm, HiChrom VP-ODS  
Mobile Phase: Acetonitrile  
Flow Rate: 1.0 mL/min  
Temperature: 40.0 °C  
Injection Volume: 10 µL  
Sample Volume: 10 µL

Criteria: 1.45-1.60  
Measured: 1.50  
Result: Pass

**Coordination Compound 2**

Column: 150 x 4.6 mm ID, 5 µm, HiChrom VP-ODS  
Mobile Phase: Acetonitrile  
Flow Rate: 1.0 mL/min  
Temperature: 40.0 °C  
Injection Volume: 10 µL  
Sample Volume: 10 µL

Criteria: 3.35-3.71  
Measured: 3.50  
Result: Pass

Verified by [ ] SHIMADZU

**Shim-pack COLUMN PERFORMANCE REPORT**

COLUMN NAME : Shim-Pack VP-ODS  
SIZE : 250 X 4.6 mm I.D.  
P/N : 228-34927-72  
SERIAL NO. : SVK148828

START

5

18

15

1

2

3

4

TEST CONDITION

MOBILE PHASE : CH3OH/H2O=70/30  
FLOW RATE : 1.0 mL/min  
TEMPERATURE : 40 °C  
DETECTION : UV at 254 nm, 0.16 AUFS  
SAMPLE SIZE : 3.0 µL  
PEAK NAME : 1, Uracil(8.13u9), 2, Naphthyl Benzoate(1.65u9), 3, Toluene(6.45u9), 4, Naphthalene(1.29u9)

RESULTS

Parameters	Criteria	Measured	Result
1R(Naphthalene)	13.00±1.50	13.32	PASS
2R(Naphthalene)	2000±	2168	PASS
TIC(Naphthalene)	1.00±0.20	0.98	PASS
PRESSURE(MPa)	8.7±1.0	9.55	PASS

ns : Retention time in min.  
N : Number of theoretical Plates, based on JP  
TF : Tailfist factor, based on USP  
PRESSURE : 1 MPa = 10.2 kgf/cm<sup>2</sup>

Inspected by F.SAGAWA  
Shimadzu Corporation

Physical Properties of Shim-pack VP-ODS

Chromatographic Performance of Shim-pack VP-ODS

Shim-pack Column Performance Report

〈Application Data〉

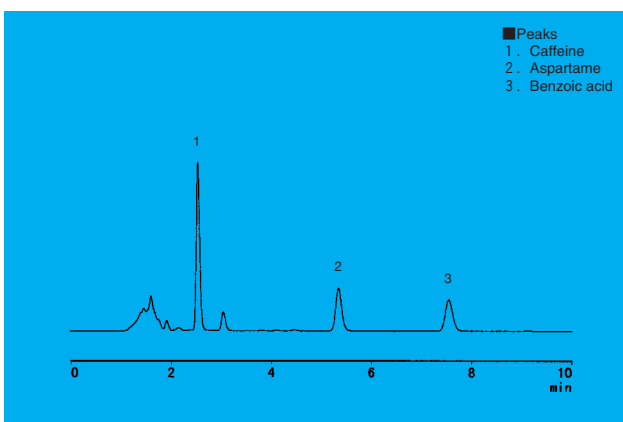


Fig. 4 Analysis of Aspartame in Soft Drink

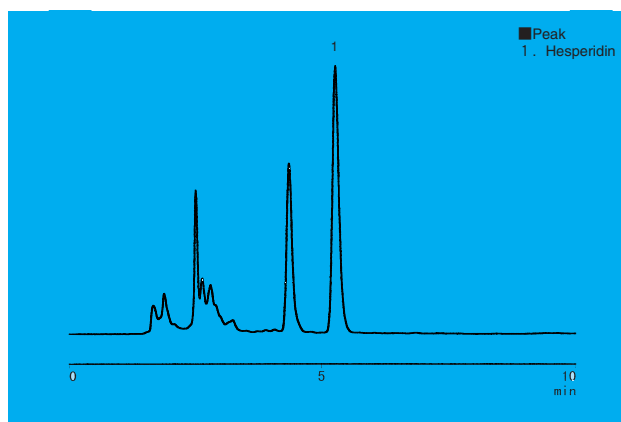


Fig. 5 Analysis of Hesperidine in Orange Juice

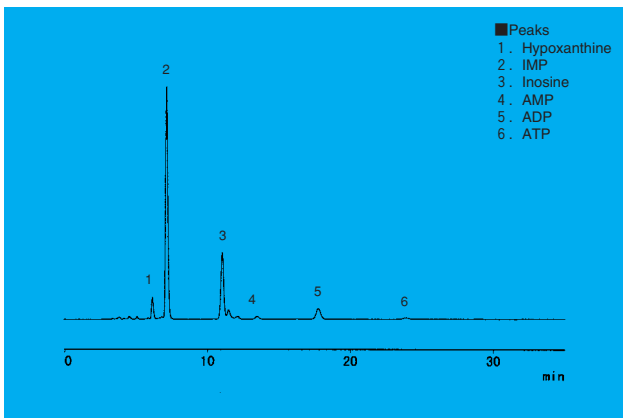


Fig. 6 Analysis of Nucleic Acid in Tuna Meat

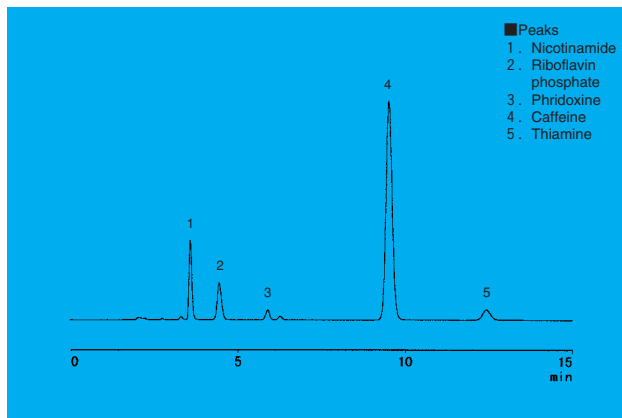


Fig. 7 Analysis of Vitamin B Group in Soft Drink

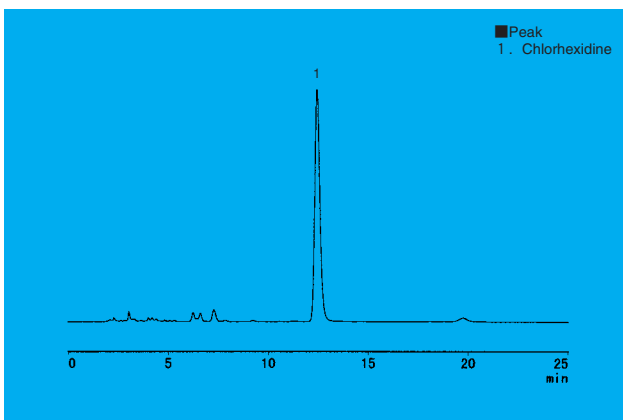


Fig. 8 Analysis of Chlorhexidine in Ointment

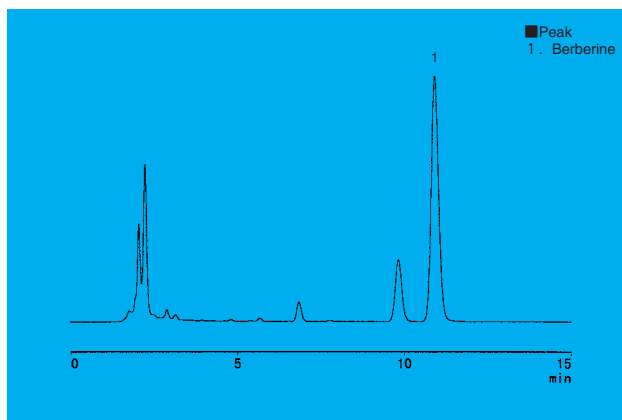


Fig. 9 Analysis of Berberine

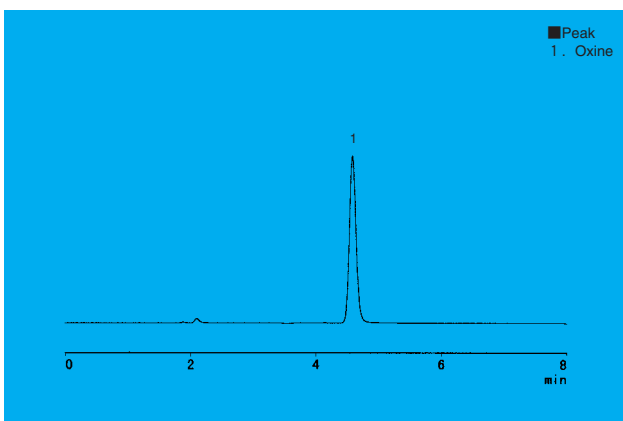


Fig. 10 Analysis of Oxine Standard

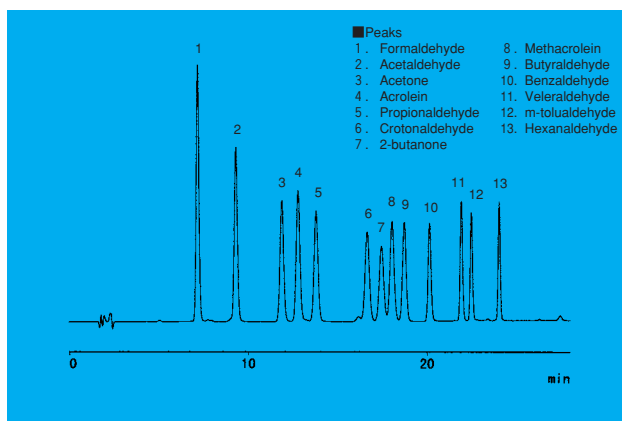


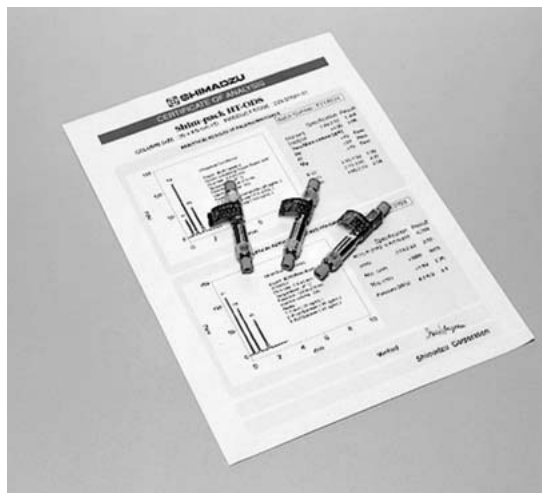
Fig. 11 Analysis of DHPH-Aldehydes, Ketones group Standard

## Shim-pack HT-ODS *High-speed HPLC columns*

### Best Partner For High Throughput Analysis

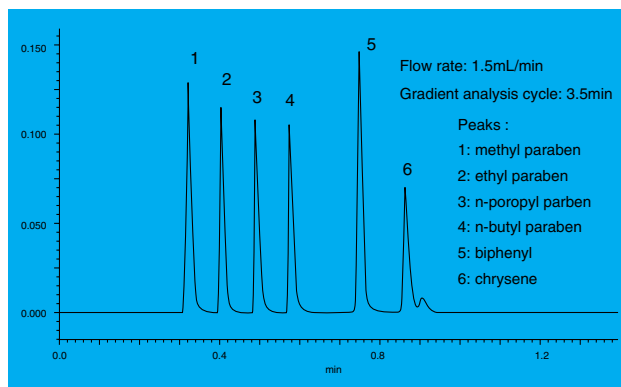
Recently, the demand for high throughput analysis has increased. For HPLC methods, it is essential to save analysis time and to obtain reliable results. To meet such requirements, Shimadzu has developed the High Throughput (HT) column packed with very fine non-porous, high purity silica particles. With this column, it is possible to realize ultra high speed analyses (Analysis times within 1 minute.) Moreover, quality certificates for the uniformity of the packing material are provided for each column so it's easy to comply with validation requirements.

We are sure this column makes your analyses faster and more reliable.



### High speed analysis

With 2 $\mu$ m and non-porous packing material, it is possible to drastically reduce the gradient analysis run times while obtaining the same peak resolution. It is possible to finish an analysis within 1 minute. Shim-pack HT-ODS provides the capability of ultra high speed and high resolution analysis.



### Validation compliance

We certify the reliability of the analysis data with a 3 column set made from different lots and attach quality certificates for the uniformity of the packing material.

### The highest performance with Shimadzu LC-2010, High-throughput HPLC

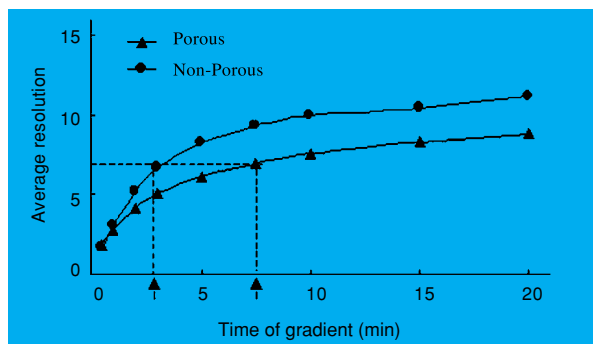
The combination of the Shim-pack HT-ODS and the Shimadzu LC-2010 is sure to help you to achieve additional high throughput in your research and analyses.



## High resolution by Non-porous silica

The right chart shows the time and the resolution in gradient analyses by a column packed with porous silica and one packed with non-porous silica. Non-porous silica shows higher resolution.

The time to obtain the same resolution with a non-porous column is less than the time required when using a porous silica column.

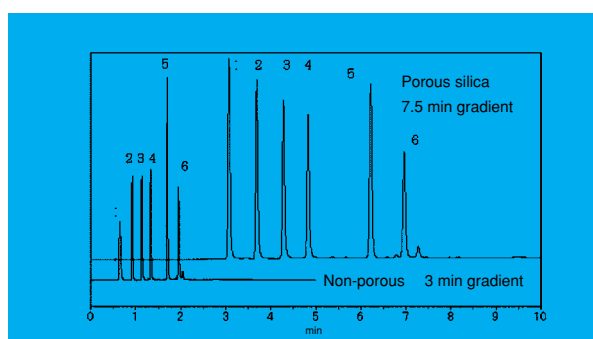


Time of gradient and resolution

## Benefit of non-porous silica column - High speed and reliable Analysis

The right chromatograms show gradient analyses with the same resolution by non-porous packed column and porous packed column. Non-porous column realizes high speed and reliable analysis.

Non-porous silica column can save your precious time and provide you more satisfactory performance.



Comparison of retention time between non-porous and porous columns.

## Shim-pack HT-ODS

Cat. No.	Description	I.D.×length (mm)	Remarks
228-37681-91	Shim-pack HT-ODS	4.6×30	2 μm non-porous silica ODS
228-37681-92	Shim-pack HT-ODS (3 column kit)	4.6×30	2 μm non-porous silica ODS A kit of columns with packings from three different production lots

## Shim-pack FC-ODS

High throughput and high resolution column packed with 3 μm porous high purity silica.

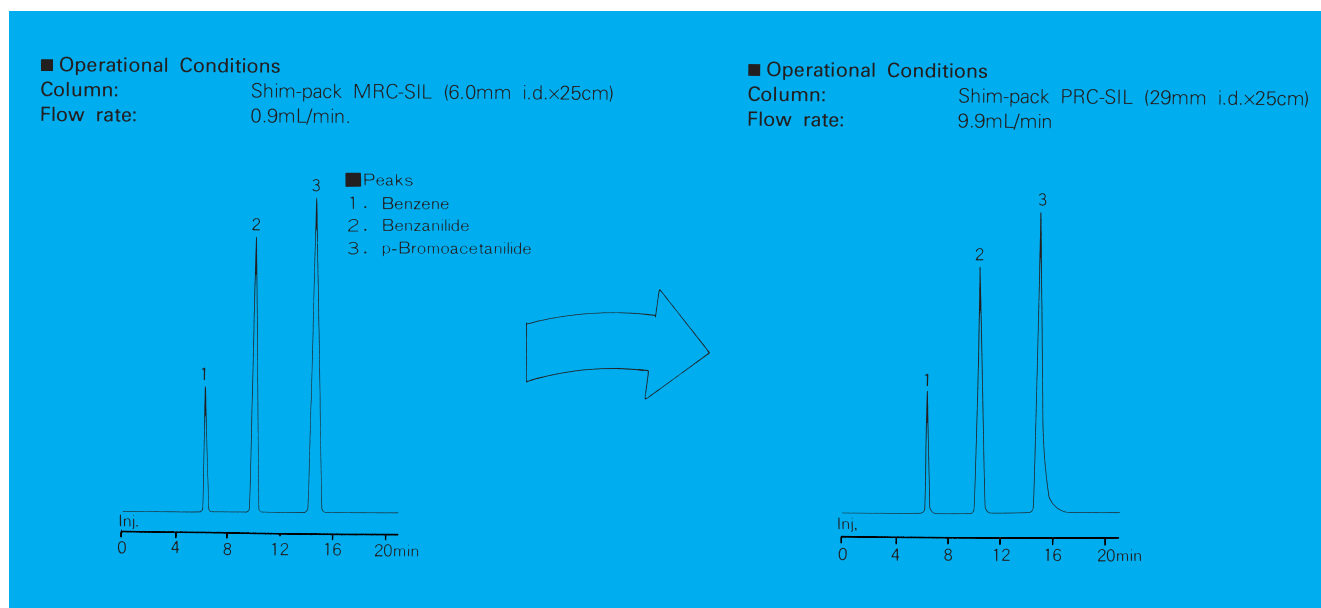
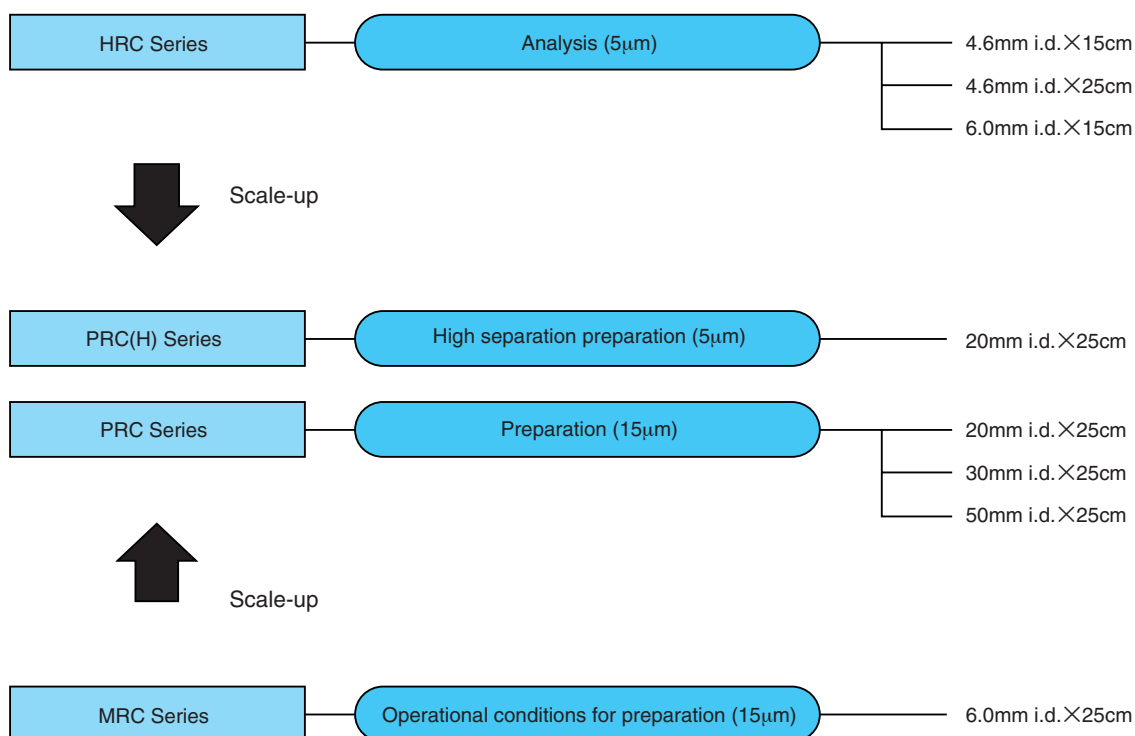
Applicable to wide application range from high throughput precisely separation analysis of complex matrix samples.

## Shim-pack FC-ODS

Cat. No.	Description	I.D.×length (mm)	Remarks
221-40511-91	Shim-pack FC-ODS	4.6×30	For fast analysis
221-40511-92	Shim-pack FC-ODS	4.6×75	For shorter analysis time
221-40511-93	Shim-pack FC-ODS	4.6×150	For high separation

## Shim-pack HRC, MRC, PRC

- High quality and outstanding performance ensured by stringent quality control.
- The same silica gel solid support is used in all the columns of this series.
- Adsorption even of basic compounds is completely eliminated by the Shimadzu's original secondary silylation method (except the -SIL series which is packed with silica particles without any surface treatment).
- The same packing material is used in the analytical and preparative columns, so that the operational parameters for analytical runs may be easily copied for preparative work.



## Shim-pack HRC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack HRC-SIL	Silica	5	Absorption	4.6mm i.d.×15cm	228-23460-91
				4.6mm i.d.×25cm	228-23460-92
				6.0mm i.d.×15cm	228-23460-93
Shim-pack HRC-ODS	Octadecyl group	5	Reversed phase	4.6mm i.d.×15cm	228-23463-91
				4.6mm i.d.×25cm	228-23463-92
				6.0mm i.d.×15cm	228-23463-93
Shim-pack HRC-C <sub>8</sub>	Octyl group	5	Reversed phase	4.6mm i.d.×15cm	228-24376-91
				4.6mm i.d.×25cm	228-24376-92
				6.0mm i.d.×15cm	228-24376-93
Shim-pack HRC-TMS	Trimethyl group	5	Reversed phase	4.6mm i.d.×15cm	228-24377-91
				4.6mm i.d.×25cm	228-24377-92
				6.0mm i.d.×15cm	228-24377-93
Shim-pack HRC-NH <sub>2</sub>	Aminopropyl group	5	Reversed, normal, ion exchange	4.6mm i.d.×15cm	228-24378-91
				4.6mm i.d.×25cm	228-24378-92
				6.0mm i.d.×15cm	228-24378-93
Shim-pack HRC-CN	Cyanopropyl group	5	Reversed phase, Normal phase	4.6mm i.d.×15cm	228-24379-91
				4.6mm i.d.×25cm	228-24379-92
				6.0mm i.d.×15cm	228-24379-93

## Guard Column for HRC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack GHRC-SIL	Silica	5	4.0mm i.d.×1cm	228-23462-91
Shim-pack GHRC-ODS	Octadecyl group	5	4.0mm i.d.×1cm	228-23465-91
Shim-pack GHRC-C <sub>8</sub>	Octyl group	5	4.0mm i.d.×1cm	228-24386-91
Shim-pack GHRC-TMS	Trimethyl group	5	4.0mm i.d.×1cm	228-24387-91
Shim-pack GHRC-NH <sub>2</sub>	Aminopropyl group	5	4.0mm i.d.×1cm	228-24388-91
Shim-pack GHRC-CN	Cyanopropyl group	5	4.0mm i.d.×1cm	228-24389-91



## Shim-pack PRC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack PRC-SIL	Silica	15	Adsorption	20mm i.d.×25cm	228-23461-93
Shim-pack PRC-SIL(K)				30mm i.d.×25cm	228-23461-94
Shim-pack PRC-SIL(L)		50mm i.d.×25cm		228-23461-95	
Shim-pack PRC-SIL(H)		20mm i.d.×25cm		228-23461-91	
Shim-pack PRC-ODS	Octadecyl group	15	Reversed phase	20mm i.d.×25cm	228-23464-93
Shim-pack PRC-ODS(K)				30mm i.d.×25cm	228-23464-94
Shim-pack PRC-ODS(L)		50mm i.d.×25cm		228-23464-95	
Shim-pack PRC-ODS(H)		20mm i.d.×25cm		228-23464-91	
Shim-pack PRC-C <sub>8</sub>	Octyl group	15	Reversed phase	20mm i.d.×25cm	228-24381-93
Shim-pack PRC-C <sub>8</sub> (H)		5		20mm i.d.×25cm	228-24381-91
Shim-pack PRC-TMS	Trimethyl group	15	Reversed phase	20mm i.d.×25cm	228-24382-93
Shim-pack PRC-TMS(H)		5		20mm i.d.×25cm	228-24382-91
Shim-pack PRC-NH <sub>2</sub>	Aminopropyl group	15	Reversed, normal, ion exchange	20mm i.d.×25cm	228-24383-93
Shim-pack PRC-NH <sub>2</sub> (H)		5		20mm i.d.×25cm	228-24383-91
Shim-pack PRC-CN	Cyanopropyl group	15	Reversed phase, normal phase	20mm i.d.×25cm	228-24384-93
Shim-pack PRC-CN(H)		5		20mm i.d.×25cm	228-24384-91

## Guard Column for PRC

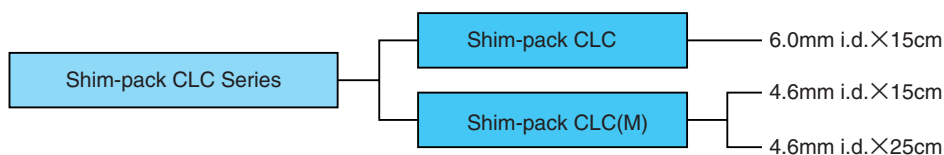
Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack GPRC-SIL	Silica	5	8mm i.d.×1.5cm	228-23462-92
Shim-pack GPRC-ODS	Octadecyl group	5	8mm i.d.×1.5cm	228-23465-92
Shim-pack GPRC-C <sub>8</sub>	Octyl group	5	8mm i.d.×1.5cm	228-24386-92
Shim-pack GPRC-TMS	Trimethyl group	5	8mm i.d.×1.5cm	228-24387-92
Shim-pack GPRC-NH <sub>2</sub>	Aminopropyl group	5	8mm i.d.×1.5cm	228-24388-92
Shim-pack GPRC-CN	Cyanopropyl group	5	8mm i.d.×1.5cm	228-24389-92
Shim-pack GPRC-SIL(K)	Silica	15	30mm i.d.×7.5cm	228-23462-93
Shim-pack GPRC-SIL(L)	Silica	15	50mm i.d.×5cm	228-23462-94
Shim-pack GPRC-ODS(K)	Octadecyl group	15	30mm i.d.×7.5cm	228-23465-93
Shim-pack GPRC-ODS(L)	Octadecyl group	15	50mm i.d.×5cm	228-23465-94

## Shim-pack MRC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack MRC-SIL	Silica	15	Adsorption	6.0mm i.d.×25cm	228-23461-92
Shim-pack MRC-ODS	Octadecyl group	15	Reversed phase	6.0mm i.d.×25cm	228-23464-92
Shim-pack MRC-C <sub>8</sub>	Octyl group	15	Reversed phase	6.0mm i.d.×25cm	228-24381-92
Shim-pack MRC-TMS	Trimethyl group	15	Reversed phase	6.0mm i.d.×25cm	228-24382-92
Shim-pack MRC-NH <sub>2</sub>	Aminopropyl group	15	Reversed, normal, ion, exchange	6.0mm i.d.×25cm	228-24383-92
Shim-pack MRC-CN	Cyanopropyl group	15	Reversed phase, normal phase	6.0mm i.d.×25cm	228-24384-92

## Shim-pack CLC

- Low cost, high-performance columns for wide field of applications.



### Shim-pack CLC (6.0mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack CLC-SIL	Silica	5	Adsorption	6.0mm i.d. x 15cm	228-00807-91
Shim-pack CLC-ODS	Octadecyl group	5	Reversed phase	6.0mm i.d. x 15cm	228-00808-91
Shim-pack CLC-C <sub>8</sub>	Octyl group	5	Reversed phase	6.0mm i.d. x 15cm	228-00809-91
Shim-pack CLC-TMS	Trimethyl group	5	Reversed phase	6.0mm i.d. x 15cm	228-00810-91
Shim-pack CLC-CN	Cyanopropyl group	5	Reversed-phase, normal phase	6.0mm i.d. x 15cm	228-00811-91
Shim-pack CLC-Phenyl	Phenyl group	5	Reversed phase	6.0mm i.d. x 15cm	228-00812-91
Shim-pack CLC-NH <sub>2</sub>	Aminopropyl group	5	Reversed, normal, ion, exchange	6.0mm i.d. x 15cm	228-16725-91

\* Also the Shim-pack VMA which is dedicated to the analysis of HVA and VMA is available.

### Shim-pack CLC(M) (4.6mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack CLC-SIL(M)	Silica	5	Adsorption	4.6mm i.d. x 15cm	228-17872-91
				4.6mm i.d. x 25cm	228-17872-92
Shim-pack CLC-ODS(M)	Octadecyl group	5	Reversed phase	4.6mm i.d. x 15cm	228-17873-91
				4.6mm i.d. x 25cm	228-17873-92
Shim-pack CLC-C <sub>8</sub> (M)	Octyl group	5	Reversed phase	4.6mm i.d. x 15cm	228-17874-91
				4.6mm i.d. x 25cm	228-17874-92
Shim-pack CLC-TMS(M)	Trimethyl group	5	Reversed phase	4.6mm i.d. x 15cm	228-17875-91
				4.6mm i.d. x 25cm	228-17875-92
Shim-pack CLC-CN(M)	Cyanopropyl group	5	Reversed-phase, normal phase	4.6mm i.d. x 15cm	228-17876-91
				4.6mm i.d. x 25cm	228-17876-92
Shim-pack CLC-Phenyl(M)	Phenyl group	5	Reversed phase	4.6mm i.d. x 15cm	228-17877-91
				4.6mm i.d. x 25cm	228-17877-92
Shim-pack CLC-NH <sub>2</sub> (M)	Aminopropyl group	5	Reversed, normal, ion, exchange	4.6mm i.d. x 15cm	228-17878-91
				4.6mm i.d. x 25cm	228-17878-92

\* Also the Shim-pack VMA which is dedicated to the analysis of HVA and VMA is available.

### Shim-pack G(4) Series (Guard Column for CLC series)

Column name	Dimensions	Cat. No.
Shim-pack G-SIL(4)	4.0mm i.d. x 1cm	228-18270-91
Shim-pack G-ODS	4.0mm i.d. x 1cm	228-18246-91
Shim-pack G-C <sub>8</sub> (4)	4.0mm i.d. x 1cm	228-18248-91
Shim-pack G-TMS(4)	4.0mm i.d. x 1cm	228-18262-91
Shim-pack G-CN(4)	4.0mm i.d. x 1cm	228-18266-91
Shim-pack G-Phenyl(4)	4.0mm i.d. x 1cm	228-18264-91
Shim-pack G-NH <sub>2</sub> (4)	4.0mm i.d. x 1cm	228-18268-91

## Shim-pack HRC, CLC

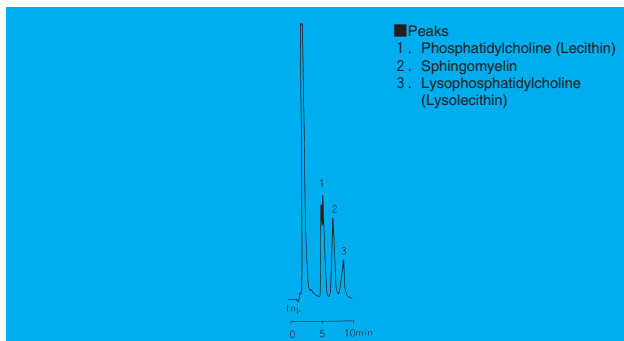


Fig. 12 Analysis of Phospholipids

## Operational Conditions

**Column:** Shim-pack CLC-SIL (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitrile/methanol/water (3/1/1)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 45°C  
**Detector:** UV spectrophotometric detector (205nm)

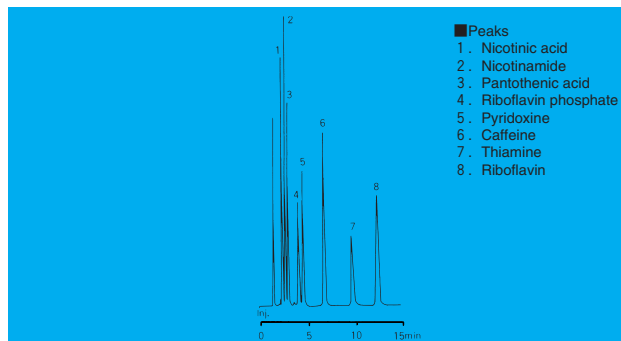


Fig. 13 Analysis of Vitamin B

## Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** [100mM phosphate buffer solution (pH 2.1) and 1.2mM sodium octane sulfonate] /acetonitrile (9/1)  
**Flow rate:** 1.2mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

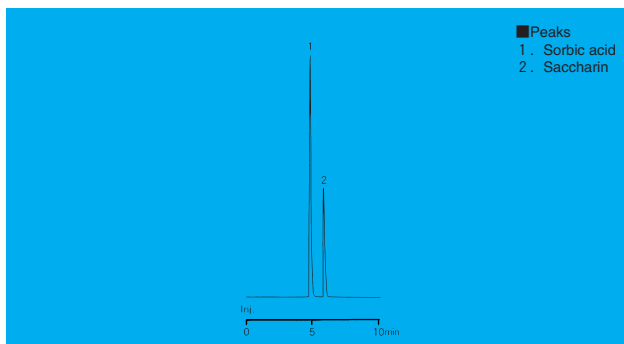


Fig. 14 Determination of Sorbic Acid and Saccharin

## Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Precolumn:** Shim-pack GRD-ODS  
**Mobile phase:** 10mM phosphoric acid buffer solution/acetonitrile (15/1)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (265nm)

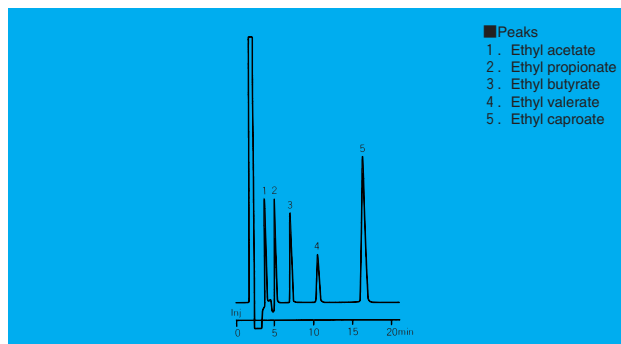


Fig. 15 Analysis of Fatty Acid Ethyl Esters

## Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitrile/water (1/1)  
**Flow rate:** 1.2mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

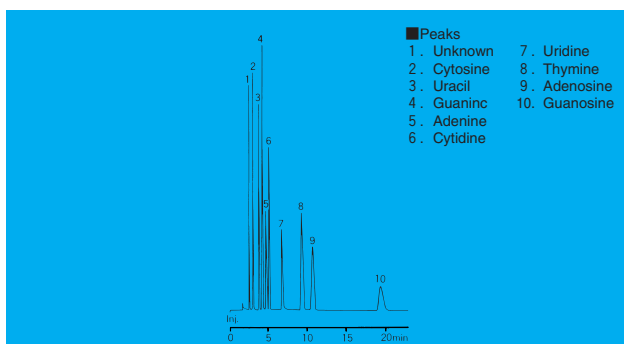


Fig. 16 Analysis of Nucleic Acid Related Compounds

## Operational Conditions

**Column:** Shim-pack HRC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1) and 0.2M sodium perchlorate  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (260nm)

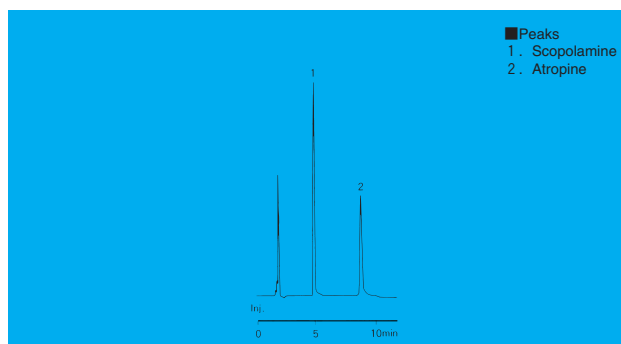


Fig. 17 Determination of Atropine and Scopolamine

## Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) /acetonitrile (5/1)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

## Shim-pack HRC, CLC

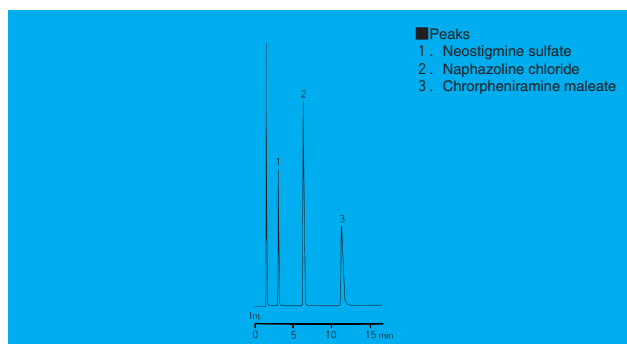


Fig. 18 Analysis of Eyewash

■ Operational Conditions  
**Column:** Shim-pack HRC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM Potassium dihydrogen phosphate /acetonitrile/phosphoric acid (70/30/0.05) with 0.1% sodium sulfonic octanoate added)  
**Flow rate:** 1.5mL/min.  
**Detector:** UV spectrophotometric detector (210nm)

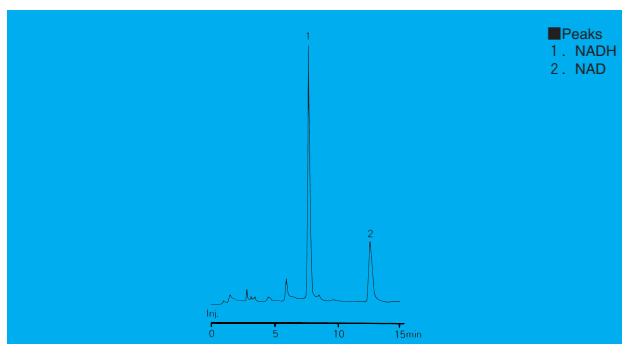


Fig. 19 Determination of NAD and NADH

■ Operational Conditions  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (260nm)

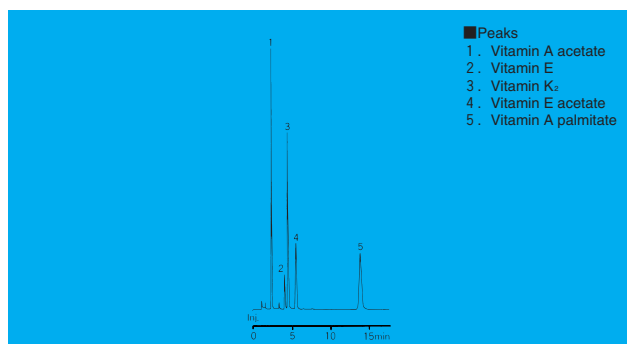


Fig. 20 Analysis of Oil Soluble Vitamines

■ Operational Conditions  
**Column:** Shim-pack HRC-ODS (4.6mm i.d.×15cm)  
**Mobile phase:** Methanol  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 55°C  
**Detector:** UV spectrophotometric detector (254nm)

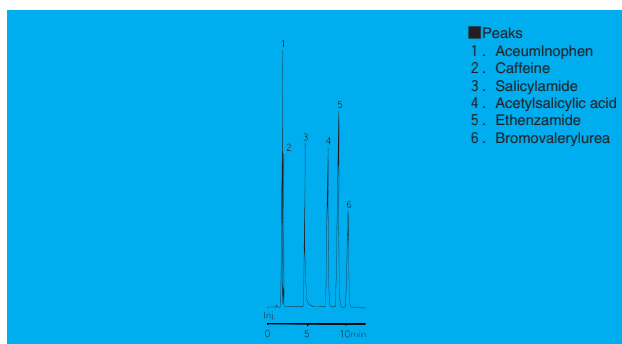


Fig. 21 Analysis of Analgesics/Antipyretics

■ Operational Conditions  
**Column:** Shim-pack HRC-ODS (4.6mm i.d.×15cm)  
**Mobile phase:** 10mM Ammonium dihydrogen phosphate (pH 2.5) /acetonitrile (4/1)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

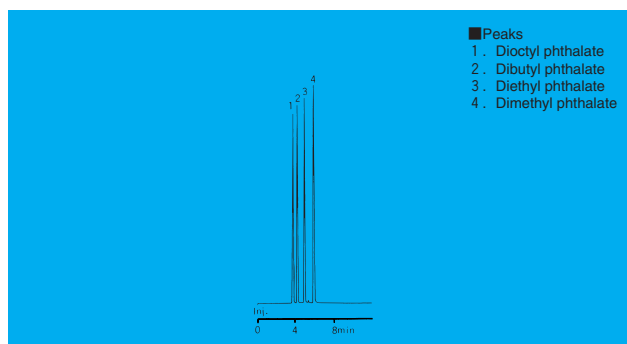


Fig. 22 Analysis of Phthalate Esters

■ Operational Conditions  
**Column:** Shim-pack HRC-SIL (4.6mm i.d.×25cm)  
**Mobile phase:** n-Hexane/ethanol (98/2)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

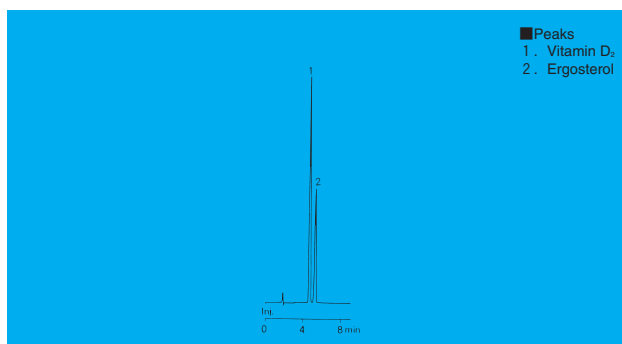


Fig. 23 Analysis of Vitamine D<sub>2</sub> and Ergosterol

■ Operational Conditions  
**Column:** Shim-pack HRC-SIL (4.6mm i.d.×25cm)  
**Mobile phase:** n-Hexane/ethanol (98/2)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

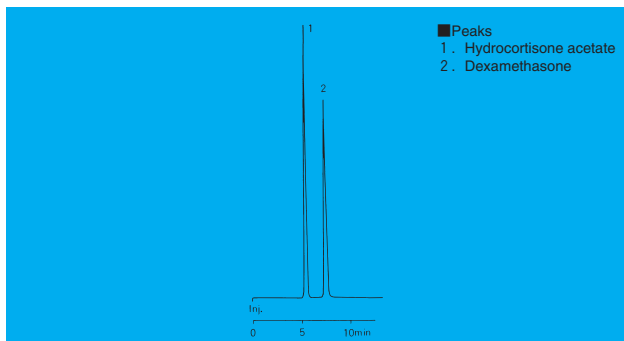


Fig. 24 Analysis of Steroid Hormones

■Operational Conditions

**Column:** Shim-pack CLC-CN (6.0mm i.d.×15cm)  
**Mobile phase:** Hexane/methanol (4/1)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

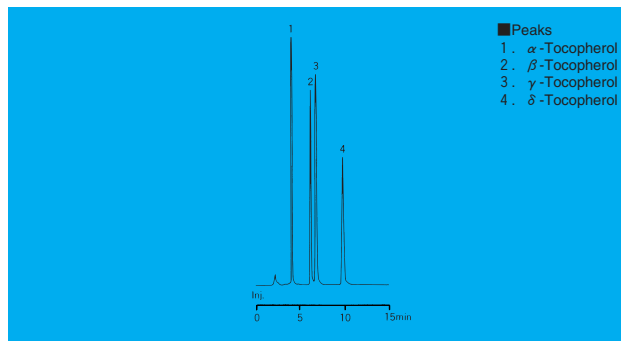


Fig. 25 Analysis of Tocopherols

■Operational Conditions

**Column:** Shim-pack CNC-NH<sub>2</sub> (6.0mm i.d.×15cm)  
**Mobile phase:** n-Hexane/isopropanol (25/1)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (297nm)

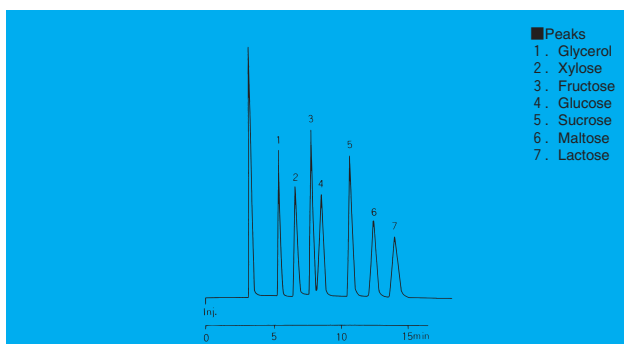


Fig. 26 Analysis of Sugars

■Operational Conditions

**Column:** Shim-pack CLC-NH<sub>2</sub> (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitrile/water (7/3)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** Refractive index detector

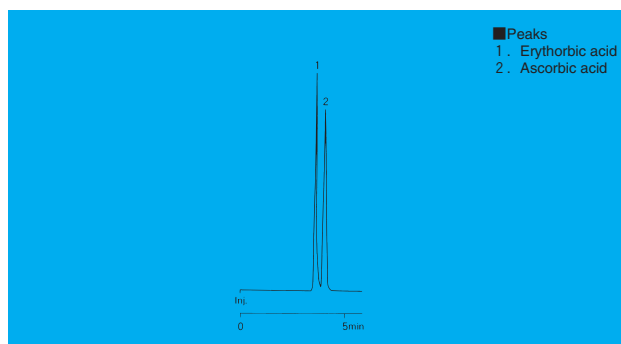


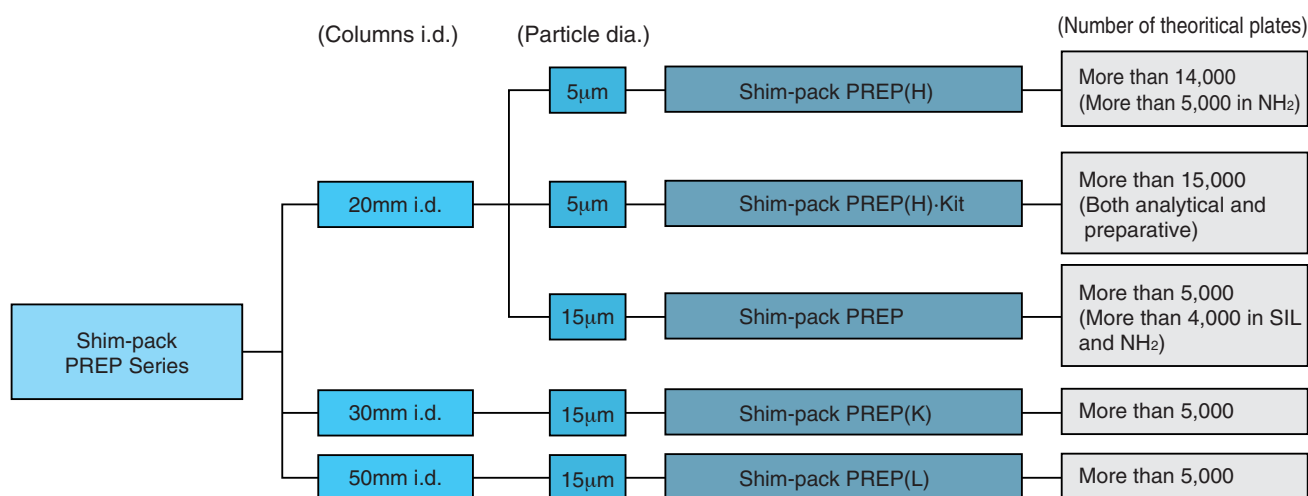
Fig. 27 Determination of Ascorbic Acid and Erythorbic Acid

■Operational Conditions

**Column:** Shim-pack CLC-NH<sub>2</sub> (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitrile/10mM phosphate buffer solution (pH 2.6) (3/1)  
**Column temperature:** 40°C  
**Flow rate:** 1.5mL/min  
**Detector:** UV spectrophotometric detector (245nm)

## Shim-pack PREP

- Shim-pack PREP Series are economical, high-performance columns for preparative LC.
- The columns are packed with fully porous spherical silica particles on which respective stationary phases are chemically bonded. (Except the PREP-SIL which is packed with silica particles without any surface treatment.)
- The residual silanol groups are end-capped by the unique silylation method. (Except the PREP-SIL)
- The PREP(H) series columns are packed with 5  $\mu\text{m}$  diameter particles (same as those used in the CLC series columns) to permit preparative LC with high resolution.
- The PREP(H) Kit is a set of an analytical and a preparative columns which are packed with packing material (5  $\mu\text{m}$  dia.) of the same lot.
- The PREP, PREP(K), and PREP(L) columns are universal LC columns, having an inner diameter of 20mm, 30mm, and 50mm, respectively.



### Shim-pack PREP(H) (5 $\mu\text{m}$ ) (20mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack PREP- $\text{C}_8$ (H)	Octyl group	5	Reversed	20mm i.d. $\times$ 25cm	228-17882-91
Shim-pack PREP-TMS(H)	Trimethyl group	5	Reversed	20mm i.d. $\times$ 25cm	228-17883-91
Shim-pack PREP-CN(H)	Cyanopropyl group	5	Normal, reversed	20mm i.d. $\times$ 25cm	228-17884-91
Shim-pack PREP-Phenyl(H)	Phenyl group	5	Reversed	20mm i.d. $\times$ 25cm	228-17885-91
Shim-pack PREP- $\text{NH}_2$ (H)	Aminopropyl group	5	Normal, reversed, ion exchange	20mm i.d. $\times$ 25cm	228-17886-91

### Shim-pack PREP(H) • Kit (5 $\mu\text{m}$ ) (20mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack PREP-SIL(H) • Kit	Silica	5	Adsorption	4.6mm i.d. $\times$ 25cm	228-17887-91
				20mm i.d. $\times$ 25cm	
Shim-pack PREP-ODS(H) • Kit	Octadecyl group	5	Reversed	4.6mm i.d. $\times$ 25cm	228-17888-91
				20mm i.d. $\times$ 25cm	

### ■ Shim-pack PREP (15 $\mu$ m) (20mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu$ m)	Separation mode	Dimensions	Cat. No.
Shim-pack PREP-SIL	Silica	15	Adsorption	20mm i.d. $\times$ 25cm	228-00814-91
Shim-pack PREP-ODS	Octadecyl group	15	Reversed	20mm i.d. $\times$ 25cm	228-00815-91
Shim-pack PREP-C <sub>8</sub>	Octyl group	15	Reversed	20mm i.d. $\times$ 25cm	228-00816-91
Shim-pack PREP-TMS	Trimethyl group	15	Reversed	20mm i.d. $\times$ 25cm	228-00817-91
Shim-pack PREP-CN	Cyanopropyl group	15	Normal, reversed	20mm i.d. $\times$ 25cm	228-00818-91
Shim-pack PREP-Phenyl	Phenyl group	15	Reversed	20mm i.d. $\times$ 25cm	228-00819-91
Shim-pack PREP-NH <sub>2</sub>	Aminopropyl group	15	Normal, reversed, ion, exchange	20mm i.d. $\times$ 25cm	228-17879-91

### ■ Shim-pack PREP(K) (15 $\mu$ m) (30mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu$ m)	Separation mode	Dimensions	Cat. No.
Shim-pack PREP-SIL(K)	Silica	15	Adsorption	30mm i.d. $\times$ 25cm	228-18273-91
Shim-pack PREP-ODS(K)	Octadecyl group	15	Reversed	30mm i.d. $\times$ 25cm	228-18319-91

### ■ Shim-pack PREP(L) (15 $\mu$ m) (50mm $\phi$ )

Column name	Stationary phase	Particle dia. ( $\mu$ m)	Separation mode	Dimensions	Cat. No.
Shim-pack PREP-SIL(L)	Silica	15	Adsorption	50mm i.d. $\times$ 25cm	228-18274-91
Shim-pack PREP-ODS(L)	Octadecyl group	15	Reversed	50mm i.d. $\times$ 25cm	228-18320-91

### ■ Shim-pack G(8), GK, and GL Guard columns

Column name	Use	Dimensions	Cat. No.
Shim-pack G-SIL(8)	Guard column for Shim-pack PREP and PREP(H)	8.0mm i.d. $\times$ 1.5cm	228-18270-92
Shim-pack G-ODS(8)		8.0mm i.d. $\times$ 1.5cm	228-18246-92
Shim-pack G-C <sub>8</sub> (8)		8.0mm i.d. $\times$ 1.5cm	228-18248-92
Shim-pack G-TMS(8)		8.0mm i.d. $\times$ 1.5cm	228-18262-92
Shim-pack G-CN(8)		8.0mm i.d. $\times$ 1.5cm	228-18266-92
Shim-pack G-Phenyl(8)		8.0mm i.d. $\times$ 1.5cm	228-18264-92
Shim-pack G-NH <sub>2</sub> (8)		8.0mm i.d. $\times$ 1.5cm	228-18268-92
Shim-pack GK-SIL	Guard column for Shim-pack PREP(K)	30mm i.d. $\times$ 7.5cm	228-18338-91
Shim-pack GK-ODS		30mm i.d. $\times$ 7.5cm	228-18321-91
Shim-pack GL-SIL	Guard column for Shim-pack PREP(L)	50mm i.d. $\times$ 5cm	228-18339-91
Shim-pack GL-ODS		50mm i.d. $\times$ 5cm	228-18322-91

## Shim-pack FLC,SBC, MBC

- The Shim-pack FLC series, SBC series, and MBC series ensure higher speeds, higher sensitivity and lower mobile phase consumption than the Shim-pack CLC series.

	(Column i.d.)	(Particle dia.)	
Shim-pack CLC	6mm i.d.,4.6mm i.d.	5 $\mu$ m	Universal use
Shim-pack FLC	4.6mm i.d.	3 $\mu$ m	Fast analysis
Shim-pack SBC	2.5mm i.d.	5 $\mu$ m	Mobile phase consumption is less than 1/3 compared with the Shim-pack CLC series. Applicable to ordinary HPLC systems.
Shim-pack MBC	1.0mm i.d.	10 $\mu$ m	Mobile phase consumption is less than 1/20 compared with the Shim-pack CLC series. Dedicated HPLC systems are required.

### Comparison of Shim-pack CLC and FLC

The Shim-pack FLC series columns use 3  $\mu$  m-diameter packing materials; the linear velocity of mobile phase can be increased without much lowering the column efficiency. This feature provide very fast analyses, which are quite useful for process control, for example.

Figure 28 shows a comparison data.

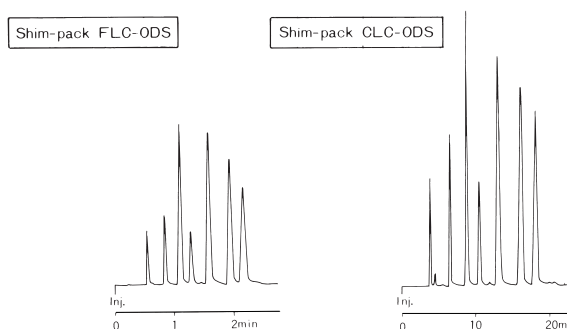


Fig. 28 Comparison Data of Shim-pack FLC and CLC

### Comparison of Shim-pack CLC, SBC, and MBC

These columns have different inner diameters. A smaller diameter column can provide a higher sensitivity, as shown in Fig. 29. Use of a small-diameter column, however, requires use of small-capacity sample injector and flow-thru cell; otherwise high sensitivity will not be provided. The Shim-pack SBC (2.5mm i.d.) is applicable to ordinary HPLC systems, while the Shim-pack MBC (1.0mm i.d.) must be used in a dedicated HPLC system.

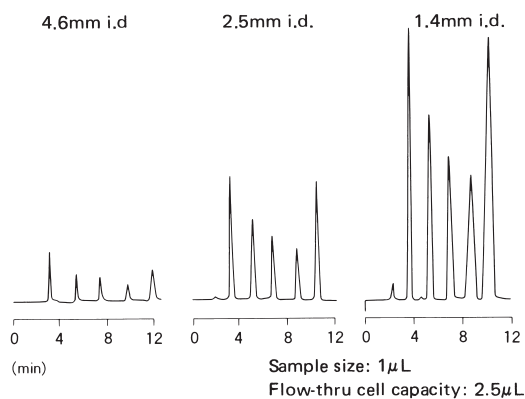


Fig. 29 Column Diameter and Peak Height



## Advantage of Pre-heater

In very fast LC using a Shim-pack FLC series column, a high mobile phase flow rate can result in broadened peaks due to a temperature gradient within the column.

A pre-heater connected between the sample injector and the column will solve this problem as demonstrated in Fig.30.

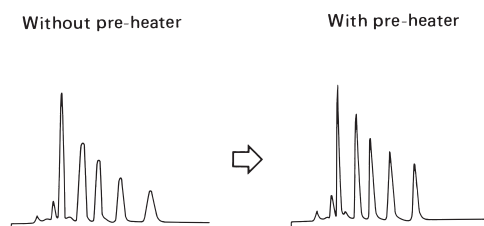


Fig. 30 Merit of Pre-heater

## Shim-pack FLC for fast LC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack FLC-ODS	Octadecyl group	3	Reversed	4.6mm i.d.×5cm	228-13375-91
Shim-pack FLC-SIL	Silica	3	Adsorption	4.6mm i.d.×5cm	228-13375-92
Shim-pack FLC-CN	Cyanopropyl group	3	Normal, reversed	4.6mm i.d.×5cm	228-13694-91
Shim-pack FLC-C <sub>8</sub>	Octyl group	3	reversed	4.6mm i.d.×5cm	228-13695-91
Shim-pack FLC-NH <sub>2</sub>	Aminopropyl group	3	Normal, reversed, ion exchange	4.6mm i.d.×5cm	228-13696-91

## Shim-pack SBC small-bore columns

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack SBC-ODS	Octadecyl group	5	Reversed	2.5mm i.d.×15cm	228-17268-91
Shim-pack SBC-C <sub>8</sub>	Octyl group	5	Reversed	2.5mm i.d.×15cm	228-17269-91
Shim-pack SBC-SIL	Silica	5	Adsorption	2.5mm i.d.×15cm	228-17270-91

## Shim-pack MBC micro-bore columns

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack MBC-ODS	Octadecyl group	10	Reversed	1.0mm i.d.×25cm	228-12812-02
				1.0mm i.d.×50cm	228-12812-05
Shim-pack MBC-SIL	Silica	10	Adsorption	1.0mm i.d.×50cm	228-12811-05
				1.0mm i.d.×100cm	228-12811-10
Shim-pack MBC-ACN	Aminopropyl group	10	Normal, reversed	1.0mm i.d.×25cm	228-12813-02
				1.0mm i.d.×50cm	228-12813-05
Shim-pack MBC-C <sub>8</sub>	Octyl group	10	Reversed	1.0mm i.d.×25cm	228-12814-02
				1.0mm i.d.×50cm	228-12814-05

\*Shim-pack MBC requires dedicated HPL system.

## Shim-pack FLC

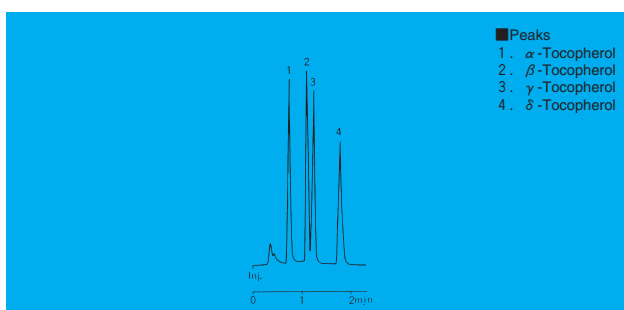


Fig. 31 Analysis of Tocopherols

### Operational Conditions

**Column:** Shim-pack FLC-SIL (4.6mm i.d.×5cm)  
**Mobile phase:** n-Hexane/dioxane/ethanol (98/2/0.2)  
**Flow rate:** 2.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (295nm)

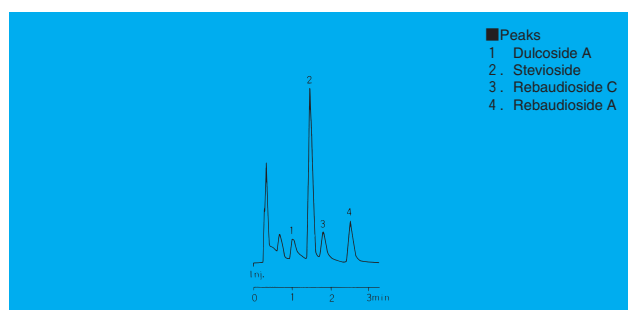


Fig. 32 Analysis of Sweet Components in Stevia Leaf

### Operational Conditions

**Column:** Shim-pack FLC-NH<sub>2</sub> (4.6mm i.d.×5cm)  
**Mobile phase:** Acetonitrile/water (8/2)  
**Flow rate:** 2.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (210nm)

## Shim-pack SBC

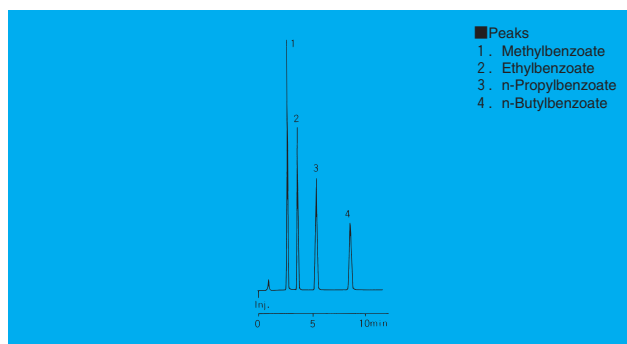


Fig. 33 Analysis of Benzoic Acid Esters

■ Operational Conditions  
**Column:** Shim-pack SBC-ODS (2.5mm i.d.×15cm)  
**Mobile phase:** Methanol/water (7/3)  
**Flow rate:** 0.4mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (250nm)

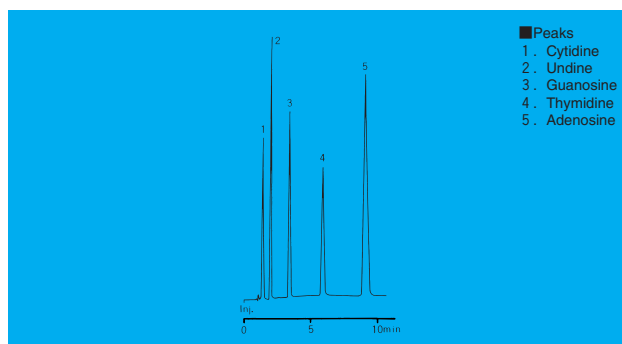


Fig. 34 Analysis of Nucleosides

■ Operational Conditions  
**Column:** Shim-pack SBC-C<sub>8</sub> (2.5mm i.d.×15cm)  
**Mobile phase:** 50mM potassium dihydrogenphosphate methanol (93/7)  
**Flow rate:** 0.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (260nm)

## Shim-pack SBC

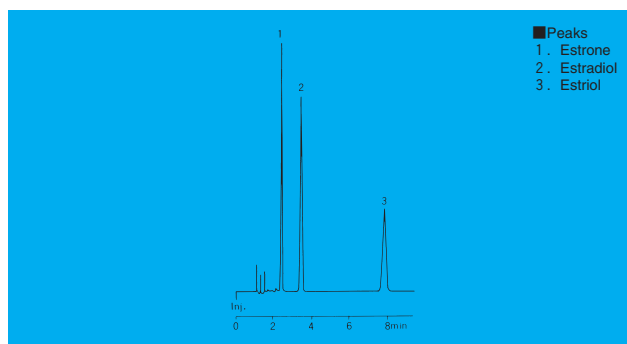


Fig. 35 Analysis of Estrogens

■ Operational Conditions  
**Column:** Shim-pack SBC-SIL (2.5mm i.d.×15cm)  
**Mobile phase:** Heptane/ethanol/methanol (90/80/2)  
**Flow rate:** 0.6mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (250nm)

## Shim-pack MBC

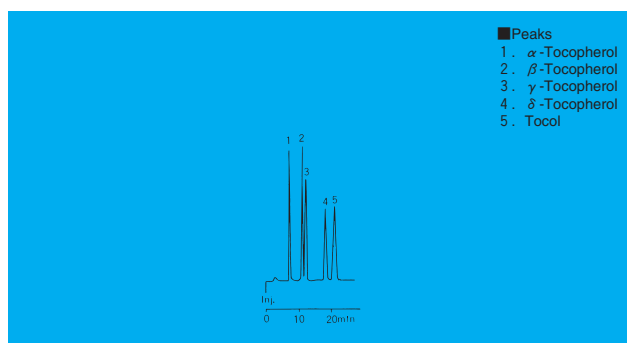


Fig. 36 Analysis of Tocopherols

■ Operational Conditions  
**Column:** Shim-pack MBC-SIL (1.0mm i.d.×50cm)  
**Mobile phase:** Hexane/dioxane/ethanol(98/2/0.2)  
**Flow rate:** 50 μL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (295nm)  
 (Micro flow-thru cell was used.)

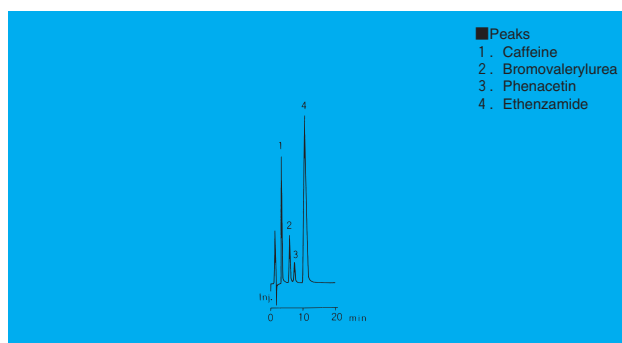
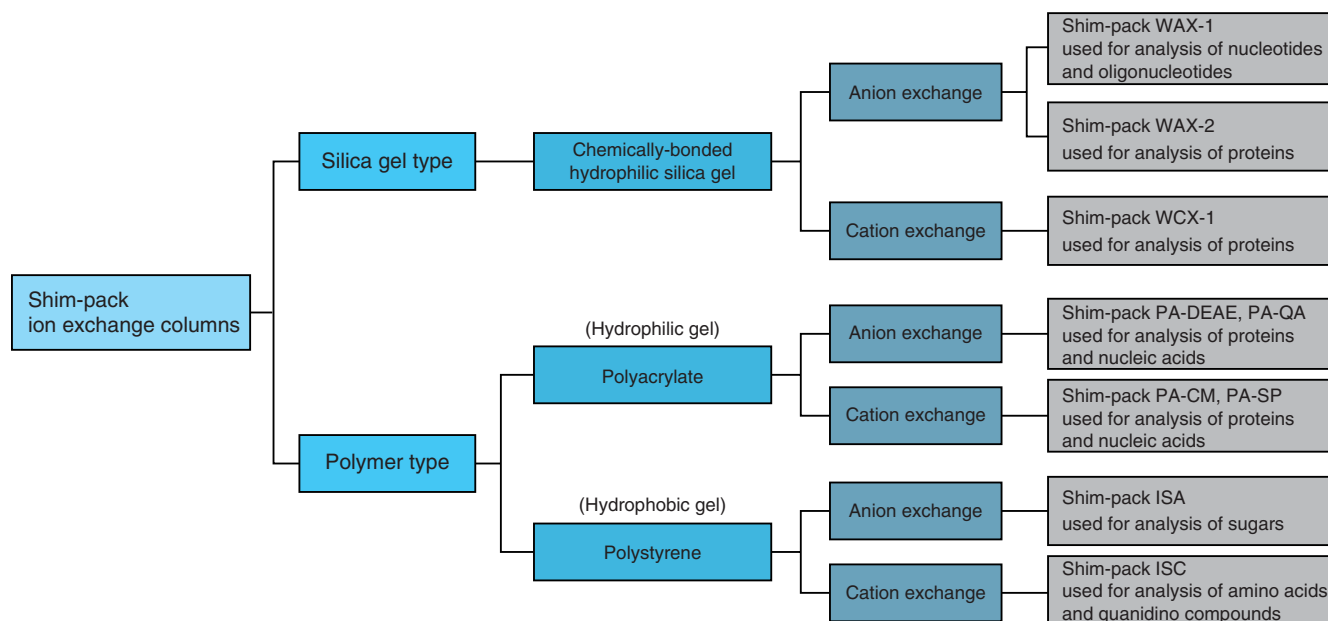


Fig. 37 Analysis of Febrifuge/Analgesics

■ Operational Conditions  
**Column:** Shim-pack MBC-ODS (1.0mm i.d.×25cm)  
**Mobile phase:** 10mM phosphoric acid buffer solution/acetonitrile (3/1)  
**Flow rate:** 100 μL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (210nm)  
 (Micro flow-thru cell was used.)

## Shim-pack Ion Exchange Columns

- The Shim-pack ion exchange columns are available in various types as shown below.



- Shim-pack ion exchange column series are classified into silica gel type and polymer type.
- The silica gel type columns ensure higher number of theoretical plates than the polymer type, because they use  $5\ \mu\text{m}$  or  $3\ \mu\text{m}$  diameter packing materials in contrast to  $10\ \mu\text{m}$  diameter packing materials used in the latter.
- The polymer type columns ensure better pH stability than the silica gel type and so they are applicable to wider pH range mobile phases.
- Polymer types are classified into polyacrylate type having hydrophilic gel and polystyrene type having hydrophobic gel. Therefore it is possible to use each depending on objecting samples.
- The Shim-pack WAX/WCX series columns are suitable for analysis of nucleotides, oligonucleotides, and proteins.
- The Shim-pack PA series columns use hydrophilic polymers as the solid support. They are especially suitable for the preparative LC of biological substances such as proteins and nucleic acids. The columns are available in two diameters, 8mm i.d. for determining the preparation conditions and 20mm i.d. for preparative work.
- The Shim-pack ISA/ISC series columns, which use polystyrene gel as the solid support, utilize electrostatic reaction and hydrophobic reaction. These columns are suitable for the analysis of sugars (ISA), amino acids (ISC-07), and guanidino compounds (ISC-05).

## Shim-pack WAX, WCX: Silica Gel Type Ion Exchange Column

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack WAX-1	Tertiary amino group	3	4.0mm i.d.×5cm	228-16225-91
Shim-pack WAX-2	Tertiary amino group	5	4.0mm i.d.×5cm	228-16365-91
Shim-pack WCX-1	Carboxyl group	5	4.0mm i.d.×5cm	228-16366-91

\* It is necessary to connect a precolumn (Cat. No. 228-16367-91) between the liquid pump and the sample injector.

## Shim-pack PA: Polyacrylate Type Ion Exchange Column

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack PA-DEAE	Diethyl amino ethyl group	10	8.0mm i.d.×10cm	228-20758-91
			20mm i.d.×10cm	228-20758-92
Shim-pack PA-QA	Tetramethyl ammonium group	10	8.0mm i.d.×10cm	228-20759-91
			20mm i.d.×10cm	228-20759-92
Shim-pack PA-CM	Carboxy methyl group	10	8.0mm i.d.×10cm	228-20760-91
			20mm i.d.×10cm	228-20760-92
Shim-pack PA-SP	Surforpropyl group	10	8.0mm i.d.×10cm	228-20761-91
			20mm i.d.×10cm	228-20761-92

## Shim-pack PAG: Guard Column for Shim-pack PA

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack PAG-DEAE	Diethyl amino ethyl group	10	8.0mm i.d.×1cm	228-20762-91
Shim-pack PAG-QA	Tetramethyl ammonium group	10	8.0mm i.d.×1cm	228-20763-91
Shim-pack PAG-CM	Carboxy methyl group	10	8.0mm i.d.×1cm	228-20764-91
Shim-pack PAG-SP	Surforpropyl group	10	8.0mm i.d.×1cm	228-20765-91

\* The Shim-pack PAG columns are also used for analysis. (See Fig. 46.)

## Shim-pack ISC, ISA: Polystyrene Type Ion Exchange Column

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack Amino-Na	Na type sulfone group	5	6.0mm i.d.×10cm	228-18837-91
Shim-pack Amino-Li	Li type sulfone group	5	6.0mm i.d.×10cm	228-18337-92
Shim-pack ISC-05/S0504	Na type sulfone group	5	4.6mm i.d.×3.8cm	228-09700-91
Shim-pack ISC-07/S1504	Na type sulfone group	7	4.0mm i.d.×15cm	228-09328-91
Shim-pack ISC-07/S1504Li	Li type sulfone group	7	4.0mm i.d.×15cm	228-00796-91
Shim-pack ISC-07/S2504	Quaternary ammonium group	7	4.0mm i.d.×25cm	228-09699-91

## Ammonia Trap Column

Column name	Dimensions	Cat. No.
Shim-pack ISC-30/S0504 (For trapping Na type ammonia)	4.0mm i.d.×5cm	228-14206-91
Shim-pack ISC-30/S0504Li (For trapping Li type ammonia)	4.0mm i.d.×5cm	228-00821-91

\* It is necessary to use one of the columns, in analysis of amino acids.

## Guard Column

\* The following are dedicated guard columns.

Column name	Dimensions	Cat. No.
ISA guard column	4.0mm i.d.×5cm	228-00823-91
ISC-07 guard column (Na type)	4.0mm i.d.×5cm	228-00802-91
ISC-07 guard column (Li type)	4.0mm i.d.×5cm	228-00797-91

## Shim-pack WCX

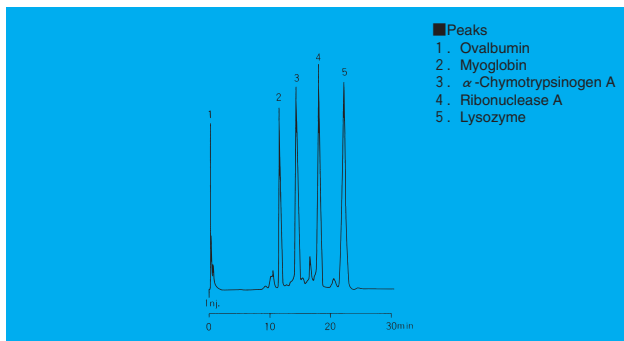


Fig. 38 Analysis of Protein Standard

**Operational Conditions**  
**Column:** Shim-pack WCX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** 20mM phosphate buffer solution (pH 6.0)/sodium sulfate, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector

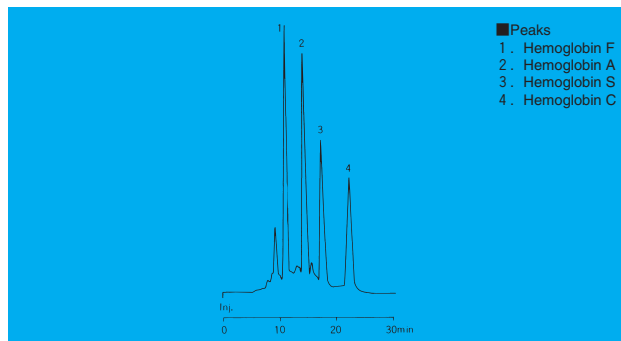


Fig. 39 Analysis of Hemoglobins

**Operational Conditions**  
**Column:** Shim-pack WCX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** 20mM phosphate buffer solution (pH 6.5)/sodium sulfate, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (415nm)

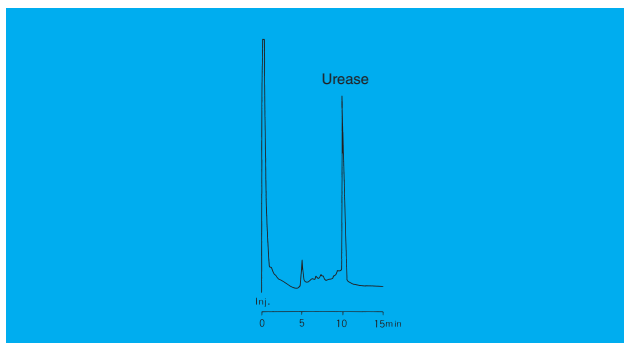


Fig. 40 Determination of Urease

**Operational Conditions**  
**Column:** Shim-pack WCX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** 20mM phosphate buffer solution (pH 6.0)/sodium sulfate, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

## Shim-pack WAX

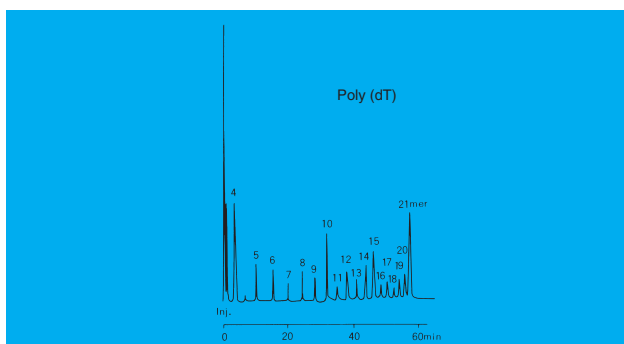


Fig. 41 Analysis of Synthesized DNAs

**Operational Conditions**  
**Column:** Shim-pack WAX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** Phosphate buffer solution (pH 6.8)/acetonitrile, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (260nm)

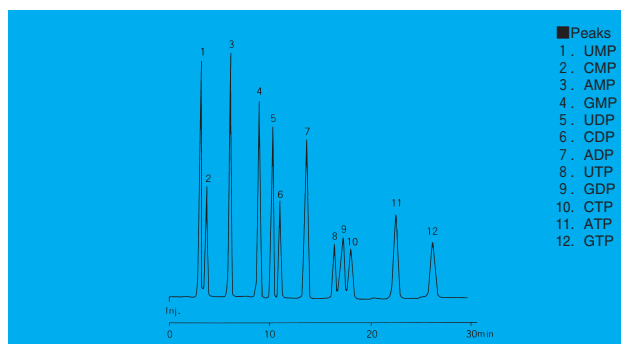


Fig. 42 Analysis of Mononucleotides

**Operational Conditions**  
**Column:** Shim-pack WAX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** 20mM phosphate buffer solution (pH 7)/480mM phosphoric acid buffer solution (pH 6.85), gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 45°C  
**Detector:** UV spectrophotometric detector (260nm)

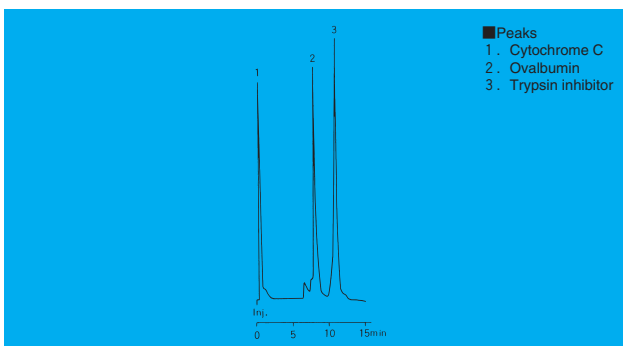


Fig. 43 Analysis of Protein Standard

**Operational Conditions**  
**Column:** Shim-pack WAX-2 (4.0mm i.d.×5cm)  
**Mobile phase:** Tris-sulfuric acid buffer solution (pH 7.5)/sodium sulfate, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

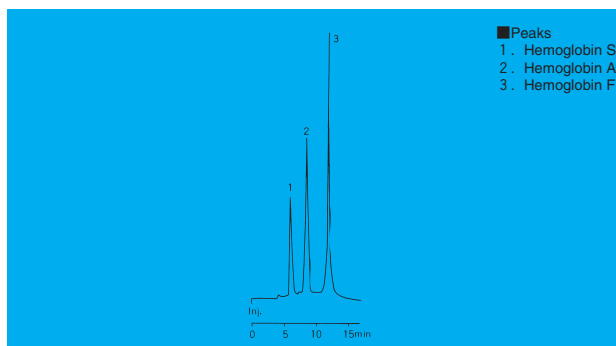


Fig. 44 Analysis of Hemoglobins

**Operational Conditions**  
**Column:** Shim-pack WAX-2 (4.0mm i.d.×5cm)  
**Mobile phase:** Tris-acetic acid buffer solution (pH 8) /sodium sulfate, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV-VIS spectrophotometric detector (415nm)

## Shim-pack PA

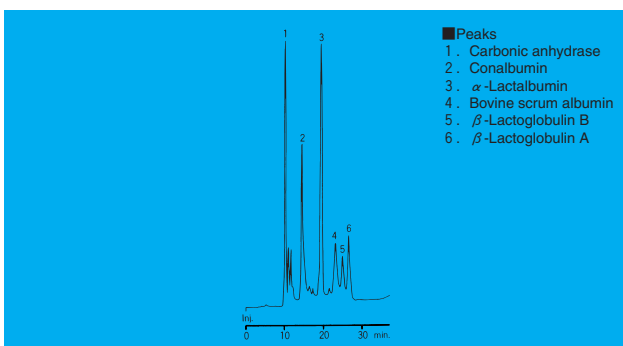


Fig. 45 Analysis of Protein Standard

**Operational Conditions**  
**Column:** Shim-pack PA-DEAE (8.0mm i.d.×1cm)  
**Mobile phase:** Tris-hydrochloric acid buffer solution (pH 8.0) /sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

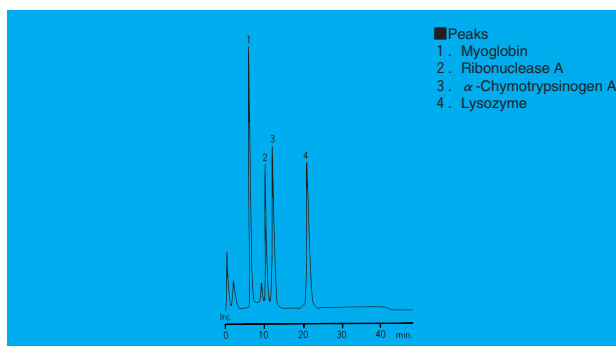


Fig. 46 Analysis of Protein Standard

**Operational Conditions**  
**Column:** Shim-pack PAG-CM (8.0mm i.d.×1cm)  
**Mobile phase:** Phosphate buffer solution (pH 6.6) /sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

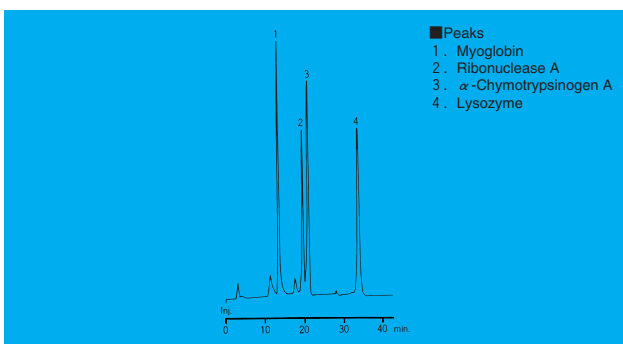


Fig. 47 Analysis of Protein Standard

**Operational Conditions**  
**Column:** Shim-pack PA-SP (8.0mm i.d.×1cm)  
**Mobile phase:** Phosphate buffer solution (pH 6.6) /sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

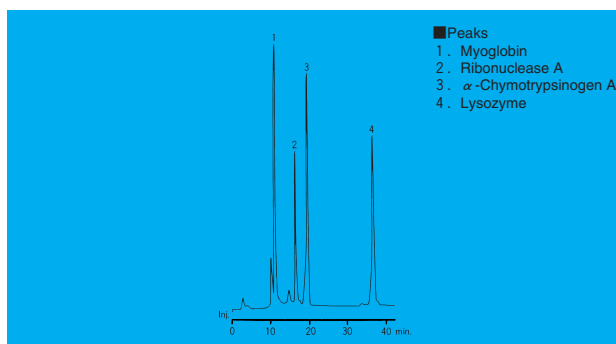


Fig. 48 Analysis of Protein Standard

**Operational Conditions**  
**Column:** Shim-pack PA-CM (8.0mm i.d.×1cm)  
**Mobile phase:** Phosphate buffer solution (pH 6.6) /sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

## Shim-pack ISA, ISC

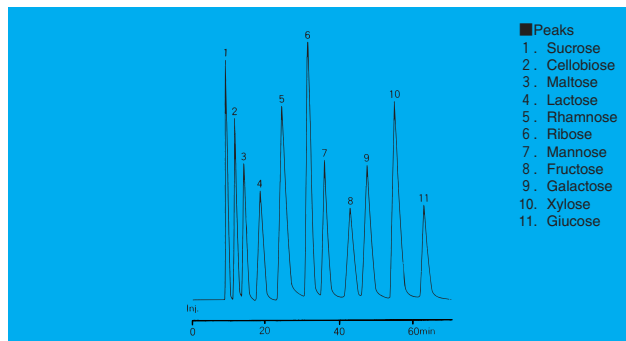


Fig. 49 Analysis of Saccharide Standard

## Operational Conditions

**Column:** Shim-pack ISA-07/S2504 (4.0mm i.d.×25cm)  
**Mobile phase:** Potassium borate buffer solution, gradient elution  
**Flow rate:** 0.6mL/min.  
**Column temperature:** 65°C  
**Detector:** Spectrofluorophotometric detector (EX. 348nm, Em. 430nm)  
**Method:** Post-column derivatization with arginine

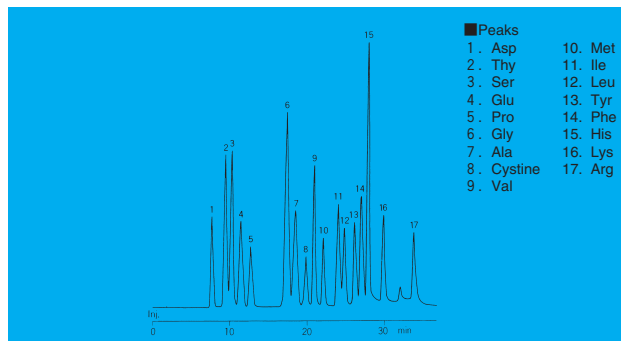


Fig. 50 Analysis of Amino Acid Standard

## Operational Conditions

**Column:** Shim-pack ISC-07/S1504 (4.0mm i.d.×15cm)  
**Mobile phase:** Sodium citrate buffer solution, gradient elution  
**Flow rate:** 0.3mL/min.  
**Column temperature:** 55°C  
**Detector:** Spectrofluorophotometric detector (EX. 348nm, Em. 450nm)  
**Method:** Post-column derivatization with ortho-phthalaldehyde

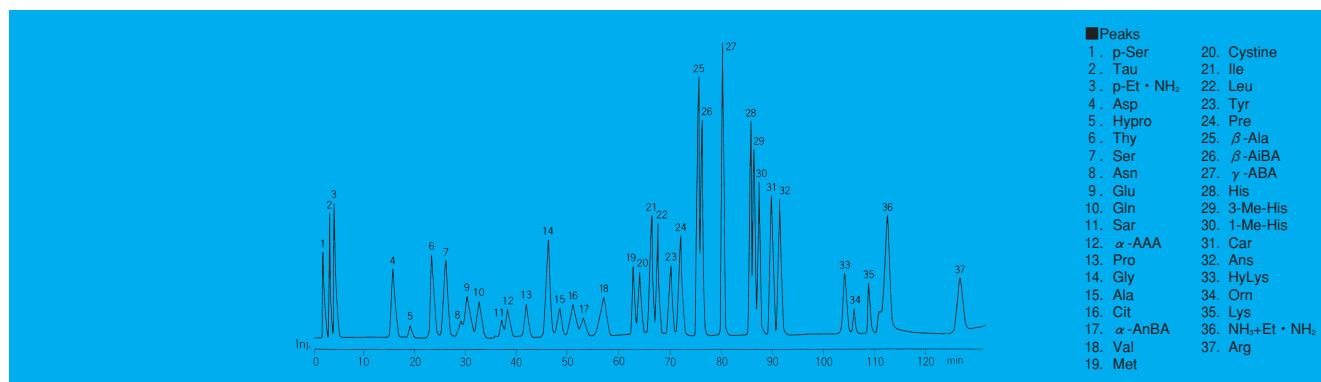


Fig. 51 Analysis of Biological Amino Acids

## Operational Conditions

**Column:** Shim-pack ISC-07/S1504Li (4.0mm i.d.×15cm)  
**Mobile phase:** Lithium citrate buffer solution, gradient elution  
**Flow rate:** 0.4mL/min.  
**Column temperature:** 38°C~58°C  
**Detector:** Spectrofluorophotometric detector (EX. 348nm, Em. 450nm)  
**Method:** Post-column derivatization with ortho-phthalaldehyde

## Shim-pack GPC

- Shim-pack GPC Series columns are used for the determination of analysis of organic solvent solution of tetrahydrofuran (800 Series), determination of chloroform (800C Series), and determination of dimethylformamide (800D series).
- The technique of GPC does not utilize such chemical reactions as partition, adsorption, and ion exchange, but a physical reaction, that is, separation based on molecular size of the sample components. Therefore, this method is suitable for the measurement of molecular weight distribution of high polymers and oligomers.
- Shim-pack GPC series columns are packed with polystyrene polymers of respective degree of cross-linking, so that you can choose a column that exactly meets your requirements, ranging from analysis of high polymers to that of oligomers.
- GPC-80M (80MC, 80MD) are mixed gel columns which get the straight of calibration curve in wide range of molecular weight.
- Shim-pack GPC-2000 Series are dedicated to preparative LC of tetrahydrofuran and chloroform. They ensure separation efficiency comparable to analytical columns as well as large preparative capacity.

### Shim-pack GPC (dedicated to determination of tetrahydrofuran)

Column name	Exclusion limit (polystyrene)	Dimensions	Cat. No.
Shim-pack GPC-801	$1.5 \times 10^3$	8.0mm i.d.×30cm	228-20803-91
Shim-pack GPC-802	$5 \times 10^3$	8.0mm i.d.×30cm	228-20804-91
Shim-pack GPC-8025	$2 \times 10^4$	8.0mm i.d.×30cm	228-20805-91
Shim-pack GPC-803	$7 \times 10^4$	8.0mm i.d.×30cm	228-20806-91
Shim-pack GPC-804	$4 \times 10^5$	8.0mm i.d.×30cm	228-20807-91
Shim-pack GPC-805	$4 \times 10^5$	8.0mm i.d.×30cm	228-20808-91
Shim-pack GPC-806	$4 \times 10^7$	8.0mm i.d.×30cm	228-20809-91
Shim-pack GPC-80M	$4 \times 10^7$ , Mixed gel	8.0mm i.d.×30cm	228-20810-91
Shim-pack GPC-807	$2 \times 10^8$	8.0mm i.d.×30cm	228-20811-91

### Guard Column (for the above)

Column name	Dimensions	Cat. No.
Shim-pack GPC-800P	4.6mm i.d.×1cm	228-20812-91

### Shim-pack GPC (dedicated to determination of chloroform)

Column name	Exclusion limit (polystyrene)	Dimensions	Cat. No.
Shim-pack GPC-801C	$1.5 \times 10^3$	8.0mm i.d.×30cm	228-20803-92
Shim-pack GPC-802C	$5 \times 10^3$	8.0mm i.d.×30cm	228-20804-92
Shim-pack GPC-8025C	$2 \times 10^4$	8.0mm i.d.×30cm	228-20805-92
Shim-pack GPC-803C	$7 \times 10^4$	8.0mm i.d.×30cm	228-20806-92
Shim-pack GPC-804C	$4 \times 10^5$	8.0mm i.d.×30cm	228-20807-92
Shim-pack GPC-805C	$4 \times 10^5$	8.0mm i.d.×30cm	228-20808-92
Shim-pack GPC-806C	$4 \times 10^7$	8.0mm i.d.×30cm	228-20809-92
Shim-pack GPC-80MC	$4 \times 10^7$ , Mixed gel	8.0mm i.d.×30cm	228-20810-92
Shim-pack GPC-807C	$2 \times 10^8$	8.0mm i.d.×30cm	228-20811-92

### Guard Column (for the above)

Column name	Dimensions	Cat. No.
Shim-pack GPC-800CP	4.6mm i.d.×1cm	228-20812-92



## Shim-pack GPC (dedicated to determination of dimethylformamide)

Column name	Exclusion limit (polystyrene)	Dimensions	Cat. No.
Shim-pack GPC-801D	$1.5 \times 10^3$	8.0mm i.d.×30cm	228-20803-93
Shim-pack GPC-802D	$5 \times 10^3$	8.0mm i.d.×30cm	228-20804-93
Shim-pack GPC-8025D	$2 \times 10^4$	8.0mm i.d.×30cm	228-20805-93
Shim-pack GPC-803D	$7 \times 10^4$	8.0mm i.d.×30cm	228-20806-93
Shim-pack GPC-804D	$4 \times 10^5$	8.0mm i.d.×30cm	228-20807-93
Shim-pack GPC-805D	$4 \times 10^6$	8.0mm i.d.×30cm	228-20808-93
Shim-pack GPC-806D	$4 \times 10^7$	8.0mm i.d.×30cm	228-20809-93
Shim-pack GPC-80MD	$4 \times 10^7$ , Mixed gel	8.0mm i.d.×30cm	228-20810-93
Shim-pack GPC-807D	$2 \times 10^8$	8.0mm i.d.×30cm	228-20811-93

## Guard Column (for the above)

Column name	Dimensions	Cat. No.
Shim-pack GPC-800DP	4.6mm i.d.×1cm	228-20812-93

## Shim-pack GPC-2000 Series

Column name	Exclusion limit (polystyrene)	Dimensions	Cat. No.
Shim-pack GPC-2001*	$1.5 \times 10^3$	20mm i.d.×30cm	228-23342-91
Shim-pack GPC-2002*	$5 \times 10^3$	20mm i.d.×30cm	228-23342-92
Shim-pack GPC-20025*	$2 \times 10^4$	20mm i.d.×30cm	228-23342-93
Shim-pack GPC-2003*	$7 \times 10^4$	20mm i.d.×30cm	228-23342-94
Shim-pack GPC-2001C**	$1.5 \times 10^3$	20mm i.d.×30cm	228-23343-91
Shim-pack GPC-2002C**	$5 \times 10^3$	20mm i.d.×30cm	228-23343-92
Shim-pack GPC-20025C**	$2 \times 10^4$	20mm i.d.×30cm	228-23343-93
Shim-pack GPC-2003C**	$7 \times 10^4$	20mm i.d.×30cm	228-23343-94

\*For preparative LC of tetrahydrofuran.

\*\*For preparative LC of chloroform.

## Guard Column (for the above)

Column name	Dimensions	Cat. No.
Shim-pack GPC-2000P (For preparative LC of tetrahydrofuran.)	8mm×5cm	228-20812-94
Shim-pack GPC-2000CP (For preparative LC of chloroform.)	8mm×5cm	228-20812-95

## Shim-pack GPC

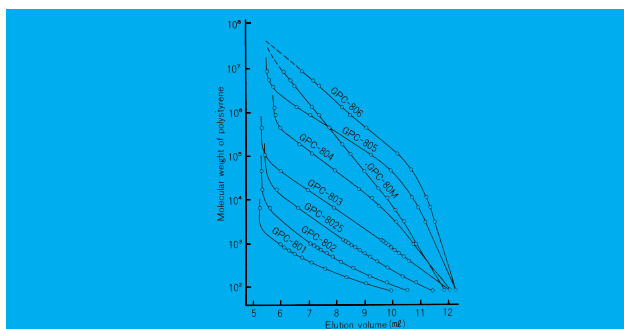


Fig. 52 Calibration Curves

### Operational Conditions

**Mobile phase:** Tetrahydrofuran  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

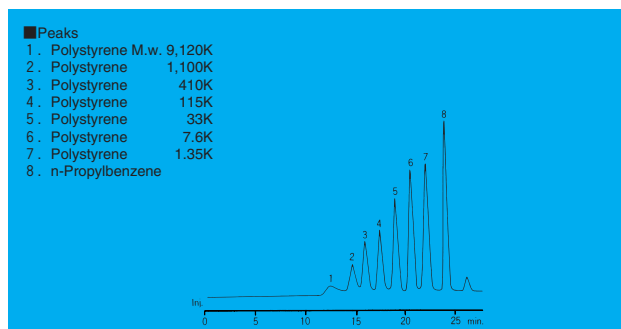


Fig. 53 Analysis of Polystyrene Standard

### Operational Conditions

**Column:** Shim-pack GPC-80M (2 columns in series)  
**Mobile phase:** Tetrahydrofuran  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

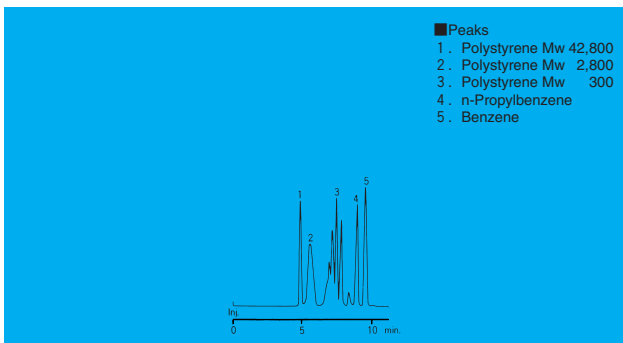


Fig. 54 Analysis of Polystyrene Standard

■Operational Conditions  
**Column:** Shim-pack GPC-802  
**Mobile phase:** Tetrahydrofuran  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (254nm)

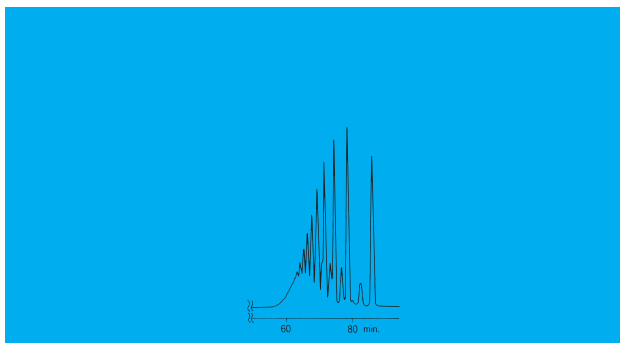


Fig. 55 Analysis of Epoxy Resin

■Operational Conditions  
**Column:** Shim-pack GPC-803 (4 pcs.) and Shim-pack GPC-8025 (4 pcs.)  
**Mobile phase:** Tetrahydrofuran  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 55°C  
**Detector:** UV spectrophotometric detector (254nm)

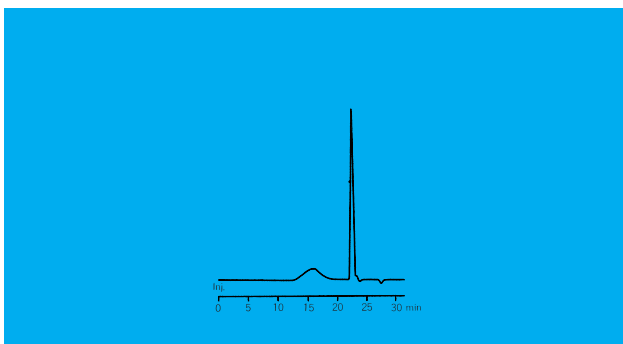


Fig. 56 Analysis of Polymethylbenzene

■Operational Conditions  
**Column:** Shim-pack GPC-80MC (2pcs.) (8mm i.d.×300mm)  
**Mobile phase:** Chloroform/Trichlorotrifluoroethane (8/2)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

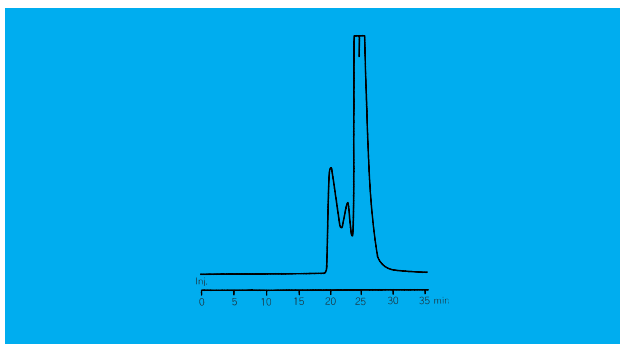


Fig. 57 Analysis of Polythioethersulfone

■Operational Conditions  
**Column:** Shim-pack GPC-80MC (8mm i.d.×300mm)  
**Mobile phase:** Chloroform  
**Flow rate:** 0.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (270nm)

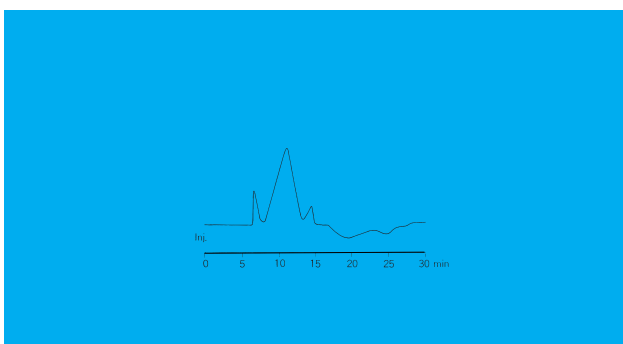


Fig. 58 Analysis of Vinyliden Poly fluoride

■Operational Conditions  
**Column:** Shim-pack GPC-80MD (8mm i.d.×300mm)  
**Mobile phase:** DMF (including 10mM LiCL)  
**Flow rate:** 0.8mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

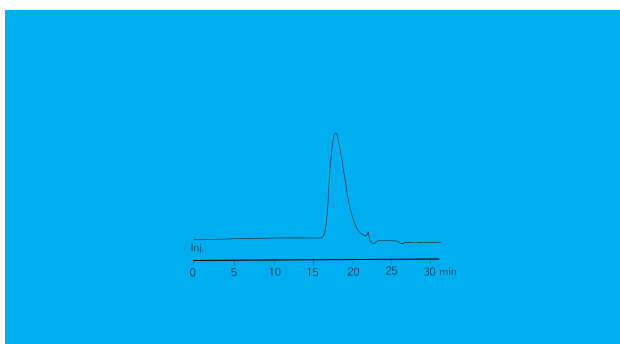
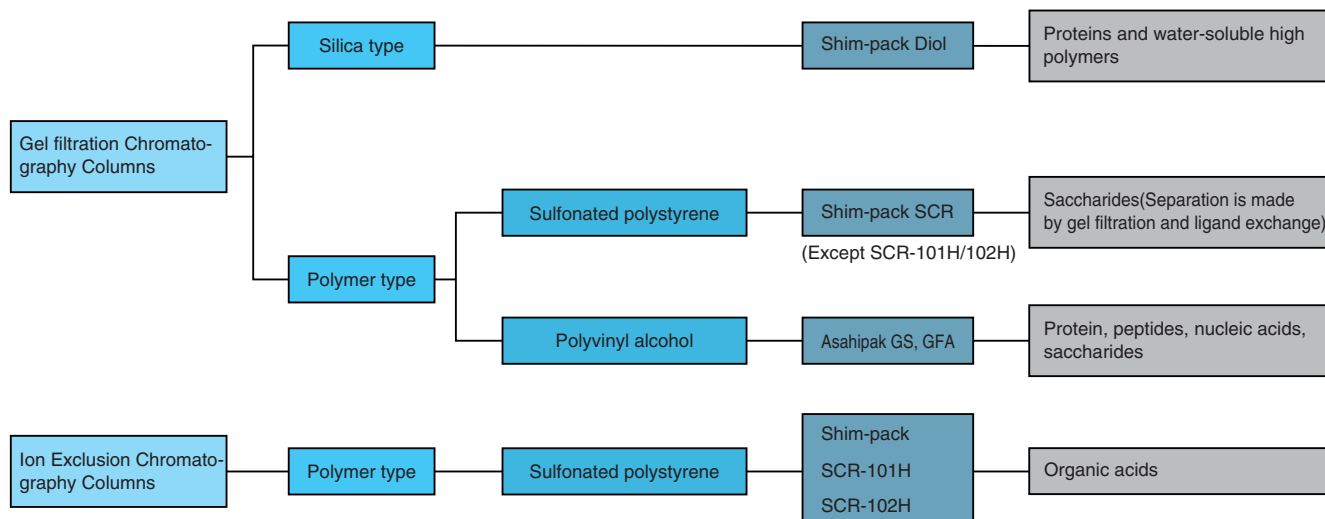


Fig. 59 Analysis of Polyetherimide

■Operational Conditions  
**Column:** Shim-pack GPC-80MD (2 pcs.) (8mm i.d.×300mm)  
**Mobile phase:** DMF (including 10mM LiCL)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

## Shim-pack Columns for Gel Filtration / Ion Exclusion Chromatography

- Gel filtration chromatography (GFC) is used to separate water-soluble high polymers such as polysaccharides, proteins, and nucleic acids, according to the molecular sizes.
- Shimadzu HPLC columns for gel filtration chromatography and ion exclusion chromatography may be classified as follows.



- Analysis of polysaccharides and oligosaccharides . . . . . Asahipak GS series are recommended.
- Analysis of monosaccharides . . . . . Shim-pack SCR series is recommended. Since the separation is made by a combination of gel filtration and ligand exchange, the selectivity differs with the type of cation.
- Analysis of proteins . . . . . Shim-pack Diol series, and Asahipak GS series are recommended. The silica type provides sharper peaks while the polymer type ensures better alkali resistancy.
- Analysis of organic acids . . . . . Ion exclusion chromatography using Shim-pack SCR-101H and SCR-102H is recommended. An acidic aqueous solution (e.g. aqueous solution of perchloric acid) is used as mobile phase and the H type sulfonated styrene polymer as stationary phase.

## Shim-pack SCR

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack SCR-101N	Na type sulfone group	10	7.9mm i.d.×30cm	228-07730-92
Shim-pack SCR-101C	Ca type sulfone group	10	7.9mm i.d.×30cm	228-17889-91
Shim-pack SCR-101P	Pb type sulfone group	10	7.9mm i.d.×30cm	228-17890-91
Shim-pack SCR-101H	H type sulfone group	10	7.9mm i.d.×30cm	228-07730-93
Shim-pack SCR-102H	H type sulfone group	7	8.0mm i.d.×30cm	228-17893-91

## Guard Column for Shim-pack SCR

Column name	Dimensions	Cat. No.
Guard column SCR(N) (Dedicated to SCR-101N)	4.0mm i.d.×5cm	228-09619-92
Guard column SCR(H) (Dedicated to SCR-101H)	4.0mm i.d.×5cm	228-09619-93
Guard column SCR(C) (Dedicated to SCR-101C)	4.0mm i.d.×5cm	228-17891-91
Guard column SCR(P) (Dedicated to SCR-101P)	4.0mm i.d.×5cm	228-17892-91
Guard column SCR-102H (Dedicated to SCR-102H)	6.0mm i.d.×5cm	228-17924-91

\* Use a guard column without fail.

## Shim-pack Diol

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack Diol-150	Diol group	5	7.9mm i.d.×25cm	228-14775-91
			7.9mm i.d.×50cm	228-14775-92
Shim-pack Diol-300	Diol group	5	7.9mm i.d.×25cm	228-14776-91
			7.9mm i.d.×50cm	228-14776-92

\* Install a precolumn (Cat. No. 228-16367-91) between the liquid pump and the sample injector.

## Shim-pack ION KS

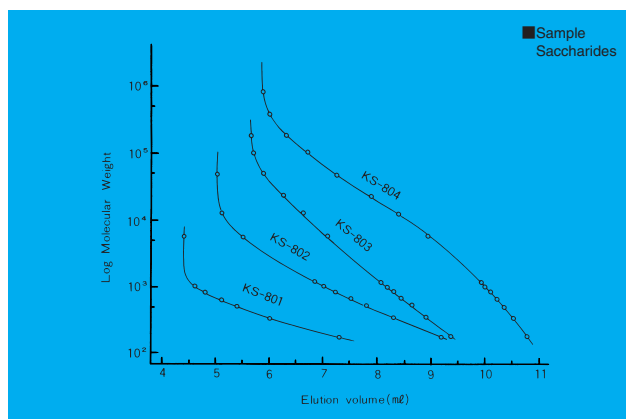


Fig. 60 Calibration Curves for ION KS Series Columns

### Operational Conditions

**Column:** Shim-pack ION KS (8.0mm i.d.×30cm)  
**Mobile phase:** Water  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 80°C  
**Detector:** Refractive index detector

## Shim-pack SCR

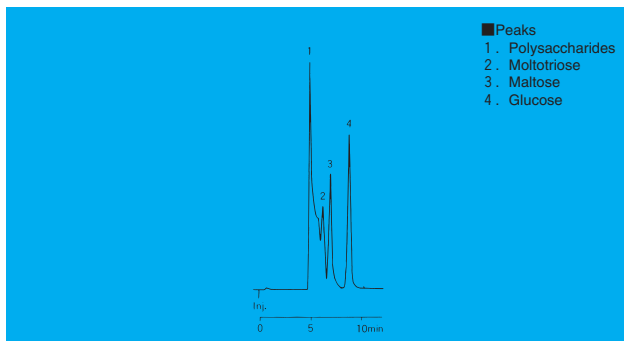


Fig. 61 Analysis of Dextrin

■ Operational Conditions  
**Column:** Shim-pack SCR-101N (7.9mm i.d.×30cm)  
**Mobile phase:** Water  
**Flow rate:** 0.8mL/min.  
**Column temperature:** 60°C  
**Detector:** Refractive index detector

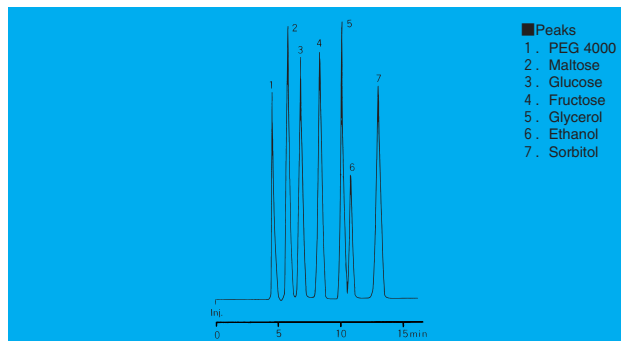


Fig. 62 Analysis of Saccharide Standard

■ Operational Conditions  
**Column:** Shim-pack SCR-101C (7.9mm i.d.×30cm)  
**Mobile phase:** Water  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 80°C  
**Detector:** Refractive index detector

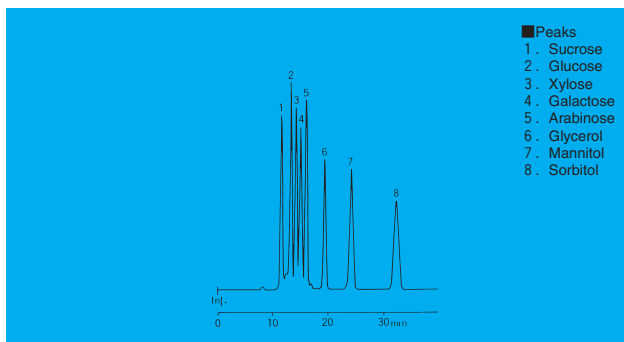


Fig. 63 Analysis of Saccharide Standard

■ Operational Conditions  
**Column:** Shim-pack SCR-101p (7.9mm i.d.×30cm)  
**Mobile phase:** Water  
**Flow rate:** 0.6mL/min.  
**Column temperature:** 80°C  
**Detector:** Refractive index detector

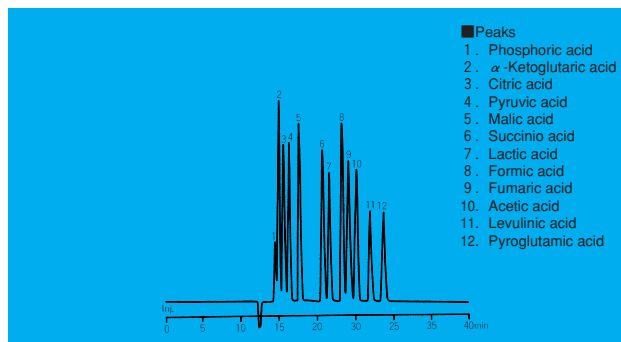


Fig. 64 Analysis of Organic Acids

■ Operational Conditions  
**Column:** Shim-pack SCR-102H (7.9mm i.d.×30cm) (2 pcs.)  
**Mobile phase:** 5mM p-Toluene sulfonic acids aqueous solution  
**Flow rate:** 0.8mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector (pH buffer organic acids analysis system)

## Shim-pack Diol

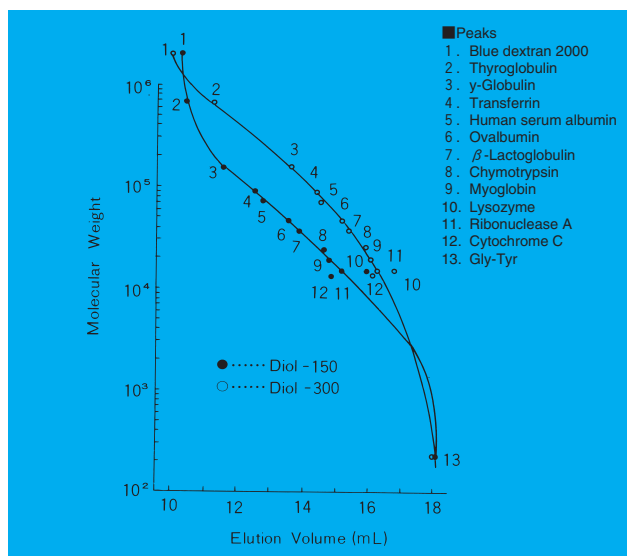


Fig. 65 Calibration Curve for Shim-pack Diol Columns

### Operational Conditions

**Column:** Shim-pack Diol (7.9mm i.d.×50cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 7) and 0.2M sodium sulfate  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

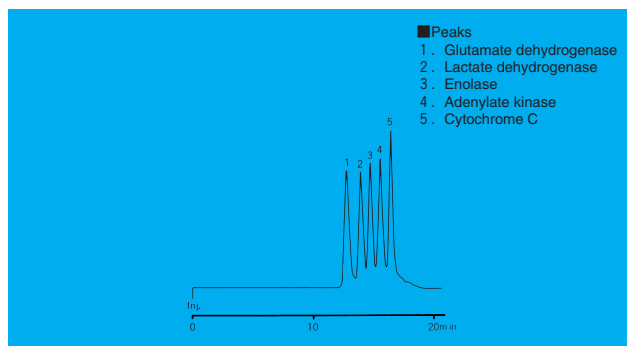


Fig. 66 Analysis of Protein Standard

### Operational Conditions

**Column:** Shim-pack Diol-300 (7.9mm i.d.×50cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 7) and 0.1M sodium chloride  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

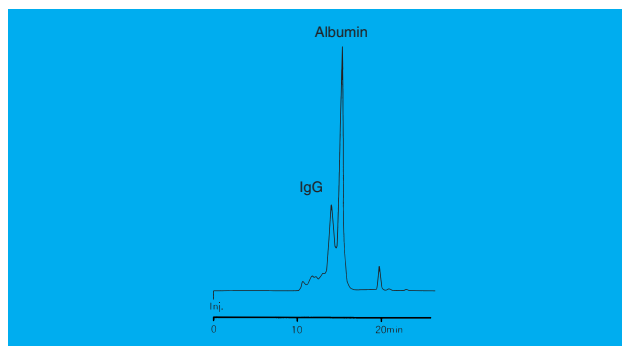


Fig. 67 Analysis of Human Serum

### Operational Conditions

**Column:** Shim-pack Diol-300 (7.9mm i.d.×50cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 7) and 0.2M sodium sulfate  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

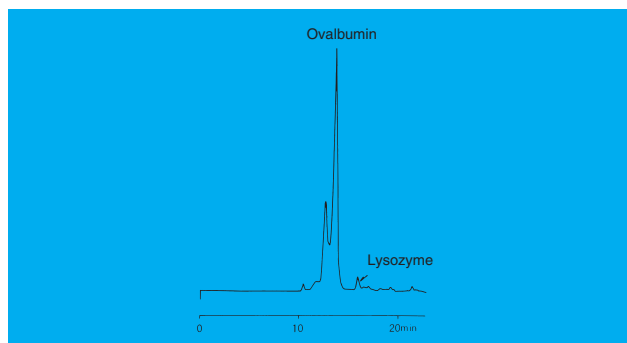


Fig. 68 Analysis of Albumin

### Operational Conditions

**Column:** Shim-pack Diol-150 (7.9mm i.d.×50cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 7) and 0.2M sodium sulfate  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

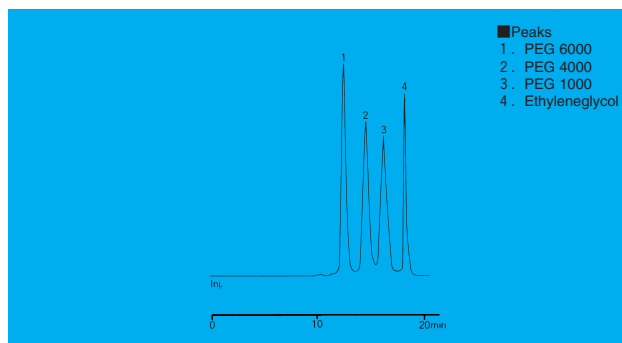


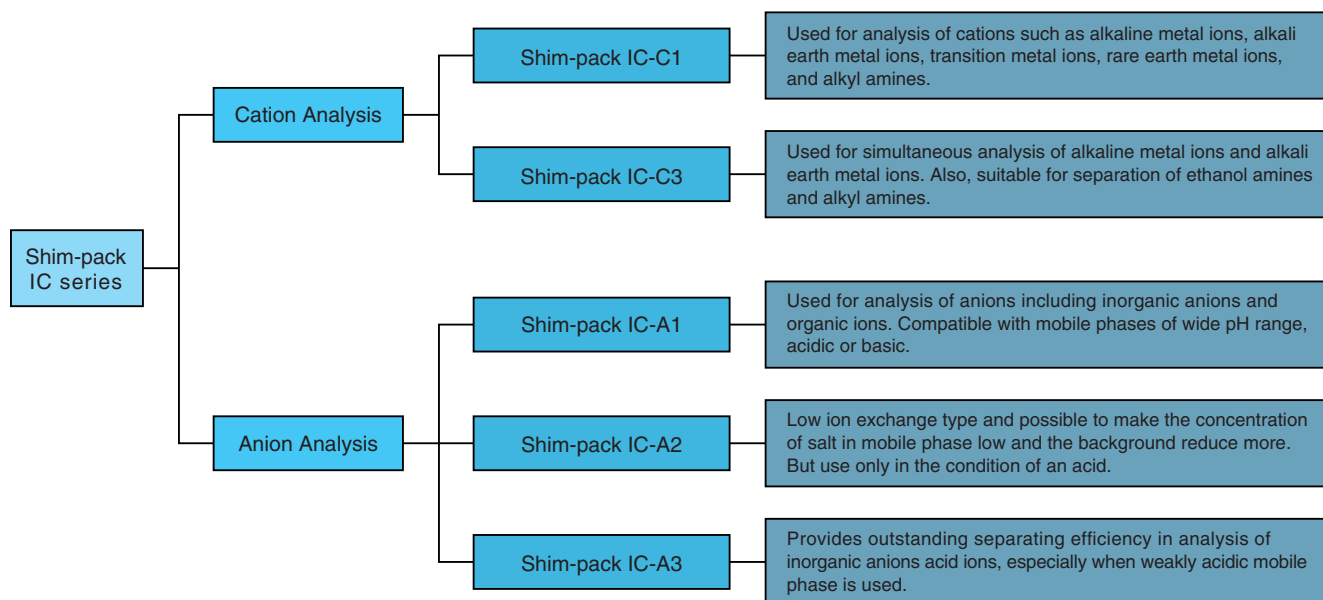
Fig. 69 Analysis of Polyethylene Glycol

### Operational Conditions

**Column:** Shim-pack Diol-150 (7.9mm i.d.×50cm)  
**Mobile phase:** Water  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** Refractive index detector

## Shim-pack IC

- Shim-pack IC series is ion exchange columns developed for ion chromatography.
- The Shim-pack IC series columns are classified as follows.



### Shim-pack IC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack IC-A1	Quaternary ammonium group	12.5	4.6mm i.d.×10cm	228-17733-91
Shim-pack IC-A1(S)	Quaternary ammonium group	10	2.0mm i.d.×15cm	228-33400-91
Shim-pack IC-A2	Quaternary ammonium group	10	4.6mm i.d.×5cm	228-17735-91
Shim-pack IC-A3	Quaternary ammonium group	5	4.6mm i.d.×15cm	228-31076-91
Shim-pack IC-A3(S)	Quaternary ammonium group	5	2.0mm i.d.×15cm	228-33366-91
Shim-pack IC-C1	Sulfone group	10	5.0mm i.d.×15cm	228-17737-91
Shim-pack IC-C1 PEEK	Sulfone group	10	4.6mm i.d.×10cm	228-33497-91
Shim-pack IC-C3	Carboxylic group	7	4.6mm i.d.×10cm	228-32329-91
Shim-pack IC-C3(S)	Carboxylic group	7	2.0mm i.d.×10cm	228-33367-91

\*Shim-pack IC(S) series are for semi-micro LC. The use of this with PIA-1000 is recommended.

### Shim-pack IC-G

Column name	Dimensions	Cat. No.
Shim-pack IC-GA1 (Dedicated to IC-A1)	4.6mm i.d.×1cm	228-17734-91
Shim-pack IC-GA2 (Dedicated to IC-A2)	4.0mm i.d.×1cm	228-17736-91
Shim-pack IC-GA3 (Dedicated to IC-A3)	4.6mm i.d.×1cm	228-31076-92
Shim-pack IC-GC1 (Dedicated to IC-C1)	4.0mm i.d.×1cm	228-17738-91
Shim-pack IC-GC1 PEEK (Dedicated to IC-C1 PEEK)	4.6mm i.d.×1cm	228-33497-92
Shim-pack IC-GC3 II (Dedicated to IC-C3)	4.6mm i.d.×7.5mm	228-41176-91

### Precolumn

Column name	Dimensions	Cat. No.
Shim-pack IC-PC1 (Dedicated to IC-C1)	8.0mm i.d.×5cm	228-17744-91

In analysis of alkali metal ions with the Shim-pack IC-C1, the use of the pre-column is recommended.

## Shim-pack IC

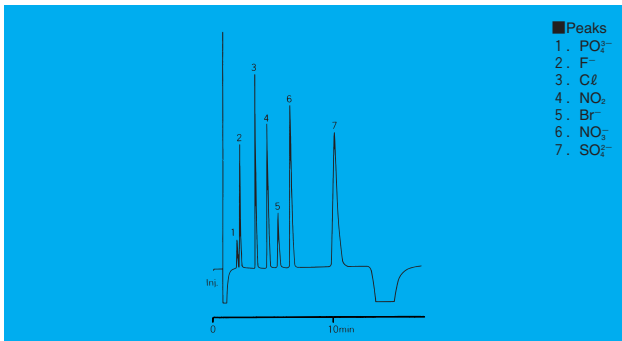


Fig. 70 Analysis of Inorganic Anions

**Operational Conditions**  
**Column:** Shim-pack IC-A3 (4.6mm i.d.×15cm)  
**Mobile phase:** 8.0mM p-Hydroxybenzoic acid and 3.2 mM Bis-Tris\*  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector  
 \*Bis(2-hydroxyethyl) iminotris (hydroxymethyl) methane

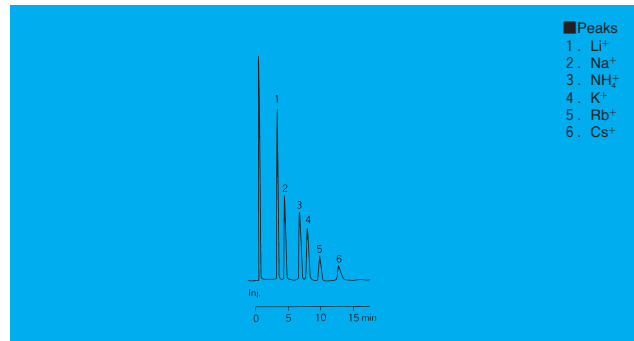


Fig. 71 Analysis of Alkali Metal Ions

**Operational Conditions**  
**Column:** Shim-pack IC-C1 (5.0mm i.d.×15cm)  
**Mobile phase:** 5mM nitric acid  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector

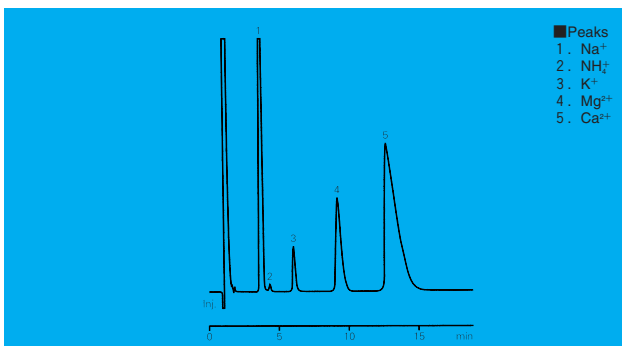


Fig. 72 Analysis of Cations in Water (includes ammonia)

**Operational Conditions**  
**Column:** Shim-pack IC-C3 (4.6mm i.d.×10cm)  
**Mobile phase:** 2.5mM oxalic acid  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector

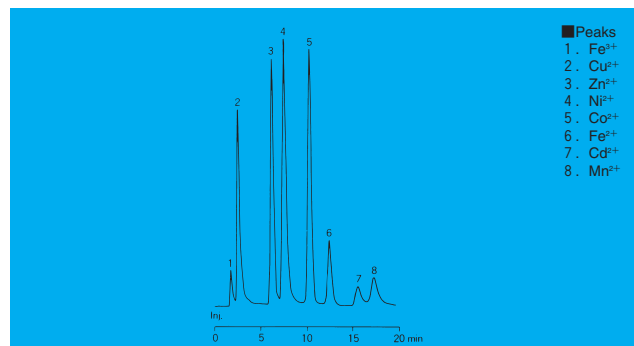


Fig. 73 Analysis of Transition Metal Ions

**Operational Conditions**  
**Column:** Shim-pack IC-C1 (5mm i.d.×15cm)  
**Mobile phase:** 0.3M lactic acid  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 40°C  
**Detector:** UV·VIS spectrophotometric detector  
**Method:** Post-column derivatization with 4-(2-pyridilazo resorcinol)



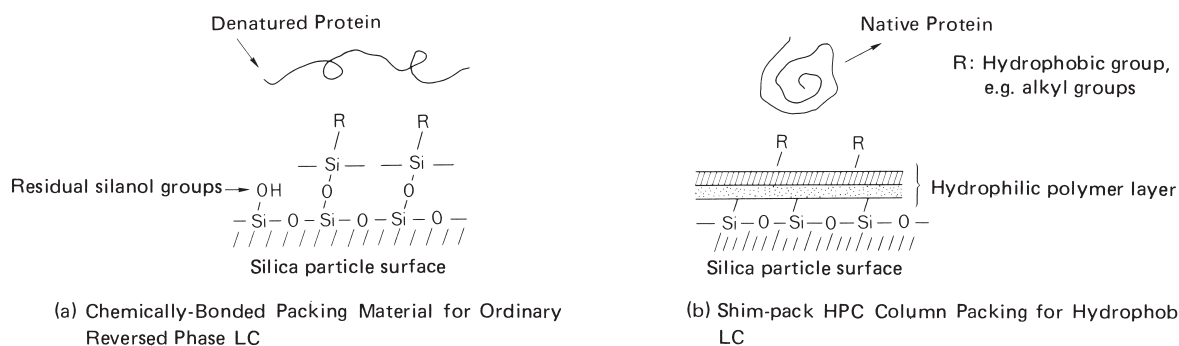
## Shim-pack HPC

- Shim-pack HPC columns have been specifically developed for the hydrophobic LC of biological high polymers such as proteins and nucleic acids.
- The columns are packed with fully-porous spherical particles, 5  $\mu\text{m}$  in diameter and 300  $\text{\AA}$  in pore diameter. The effect of the residual silanol groups is minimized by the unique chemical treatment technique.
- Ethyl groups or n-propyl groups are chemically bonded as stationary phase.

### Hydrophobic LC and Reversed Phase LC

Both of these techniques use hydrophobic column packings, but are different in the following points.

- The column packings for reversed phase LC are more hydrophobic than those for hydrophobic LC.
- In reversed phase LC, since organic solvent is generally used as mobile phase, the high-degree structures of proteins are often transmuted and the enzymes deactivated. In hydrophobic LC, on the other hand, since buffer solutions containing some salt are generally used, the high-degree structures of proteins are hardly transmuted.
- The retention of sample components is more dependent on the degree of hydrophobicity of the stationary phase in hydrophobic LC than in reversed phase LC. It is, therefore, more important to select the stationary phase, HPC-C2 or HPC-C3, in hydrophobic LC.



### Shim-pack HPC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack HPC-C2	Ethyl group	5	4.0mm i.d.×5cm	228-17792-91
Shim-pack HPC-C3	n-Propyl group	5	4.0mm i.d.×5cm	228-17793-91

\*It is necessary to install a pre-column between the liquid pump and the sample injector, to protect the Shim-pack HPC column.

## Shim-pack HPC

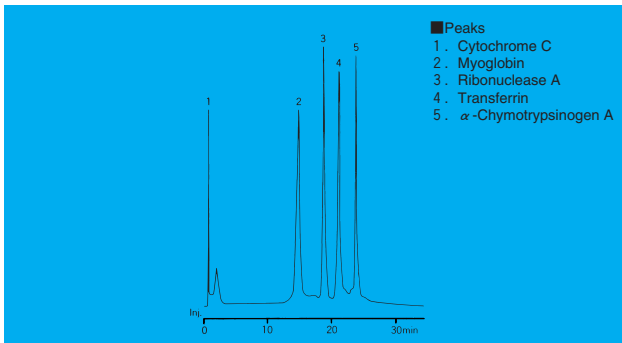


Fig. 74 Analysis of Protein Standard

### Operational Conditions

**Column:** Shim-pack HPC-C2 (4.0mm i.d.×5cm)  
**Mobile phase:** Ammonium sulfate/phosphate buffer solution (pH 7), gradient elution  
**Flow rate:** 0.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (220nm)

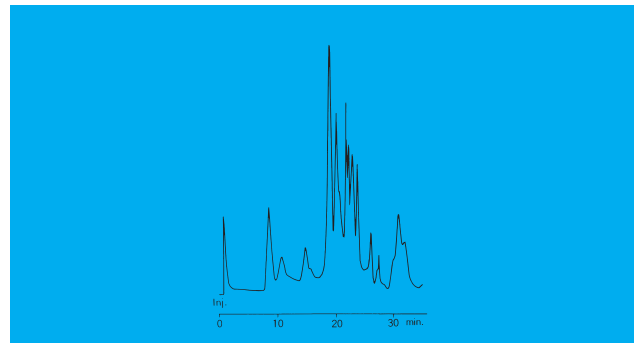


Fig. 75 Analysis of Snake Venom

### Operational Conditions

**Column:** Shim-pack HPC-C2 (4.0mm i.d.×5cm)  
**Mobile phase:** Ammonium sulfate/phosphate buffer solution (pH 7), gradient elution  
**Flow rate:** 0.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (220nm)

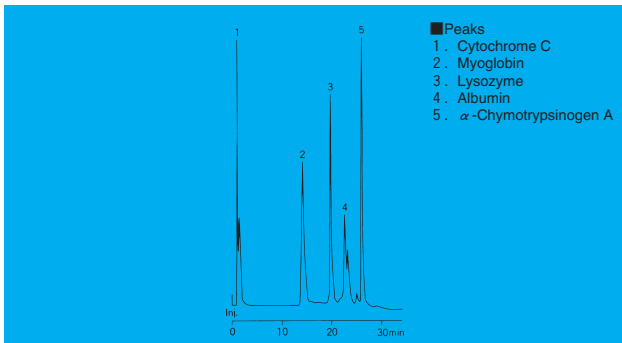


Fig. 76 Analysis of Glucosidase

### Operational Conditions

**Column:** Shim-pack HPC-C3 (4.0mm i.d.×5cm)  
**Mobile phase:** Ammonium sulfate/phosphate buffer solution (pH 7), gradient elution  
**Flow rate:** 0.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (220nm)

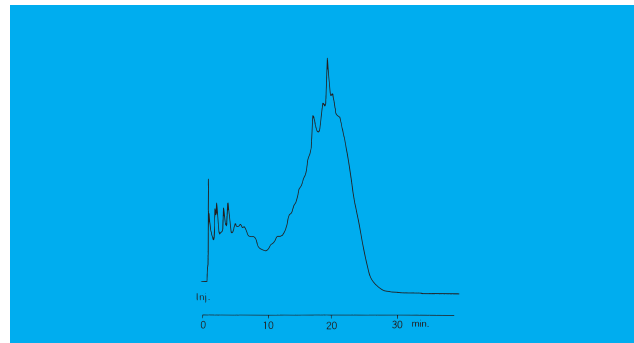


Fig. 77 Analysis of Ribonucleic Acids

### Operational Conditions

**Column:** Shim-pack HPC-C3 (4.0mm i.d.×5cm)  
**Mobile phase:** Ammonium sulfate/phosphate buffer solution (pH 7), gradient elution  
**Flow rate:** 0.5mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (260nm)

## Shim-pack SPC

- The Shim-pack SPC has specifically developed for automatic sample pretreatment and concentration by the column switching method.
- The Shim-pack SPC-RP3 column is packed with polymer particles and used for reversed phase LC.
- The Shim-pack SPC-AE1 column is packed with fully porous silica gel particles on which weakly basic anion exchange functions are chemically bonded.
- The Shimadzu Automatic Sample Pretreatment HPLC System is recommended.

### HPLC System with Automatic Sample Pretreatment Functions

This system is used for analysis of samples that contain many interfering compounds. In the first step, the sample is roughly separated in the pretreating column. Then, only the fraction that contains the target compounds is made to enter the analytical column, the switching being done with a valve. The sample pretreatment is carried out in a completely automated sequence.

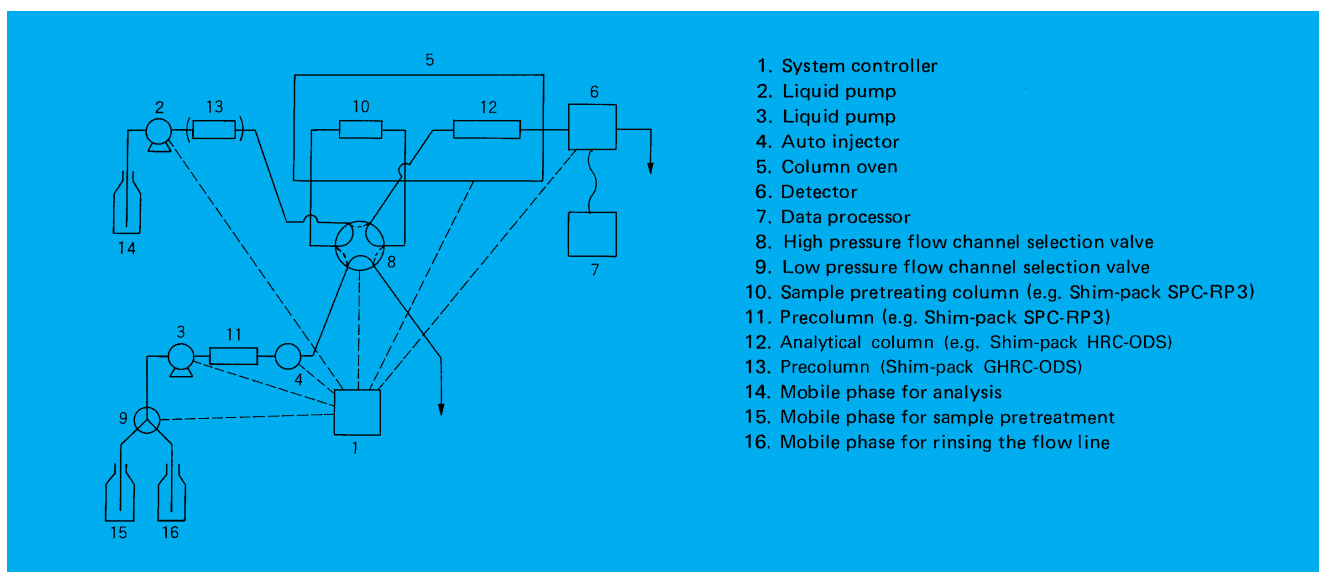


Fig. 78 Shimadzu Automatic Sample Pretreatment HPLC System

### Shim-pack SPC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack SPC-RP3	Polymer	9	Reversed phase	4.0mm i.d. $\times$ 3cm	228-33713-91
Shim-pack SPC-RP2	Polymer	10	Reversed phase	4.6mm i.d. $\times$ 1cm	228-18838-91
Shim-pack SPC-AE1	Tertiary amino group	10	Anion exchange	4.0mm i.d. $\times$ 1cm	228-17990-91

## Shim-pack GRD, Pre-column

- Undissolved materials in mobile phase is the cause of stuffing analytical column and consuming packing materials. Installing Shim-pack GRD-ODS, you can escape from these causes above. Connect this between the liquid pump and the sample injector bellow.
- Using GRD, Pre-column makes the life of analytical columns long.

### Shim-pack GRD-ODS

Effective particularly when reversed phase column is used and both mobile phase pH is higher than neutral and ion pair reagent involving nitrogen such as tetrabutyl ammonium is used.

### Pre-column Diol

The use of pre-column is recommended when using Shim-pack Diol, Shim-pack WAX, WCX, Shim-pack HPC series. Particularly when the concentration of the salt is high, it is available to eliminate un-dissolved materials in salt.

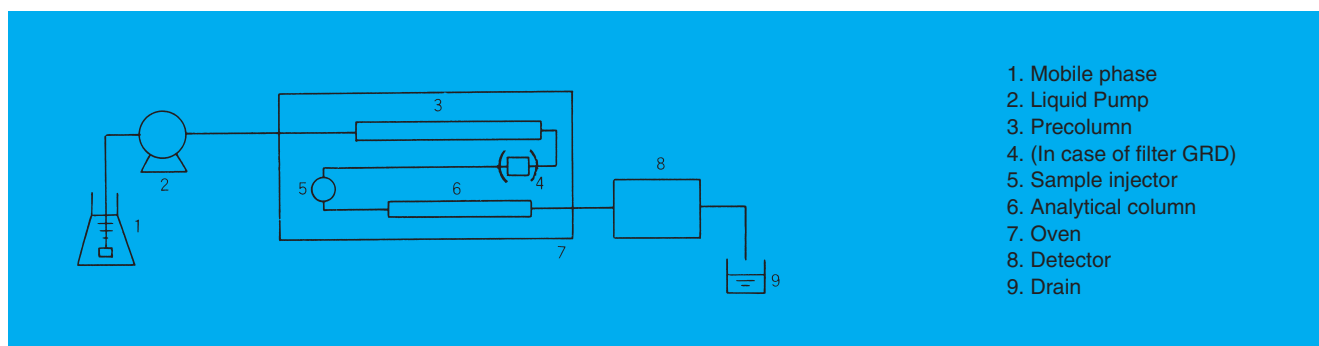


Fig. 79 Flow line having Pre-column

■ Shim-pack GRD, Pre-column (It is necessary to install a pre-column between the liquid pump and the sample injector.)

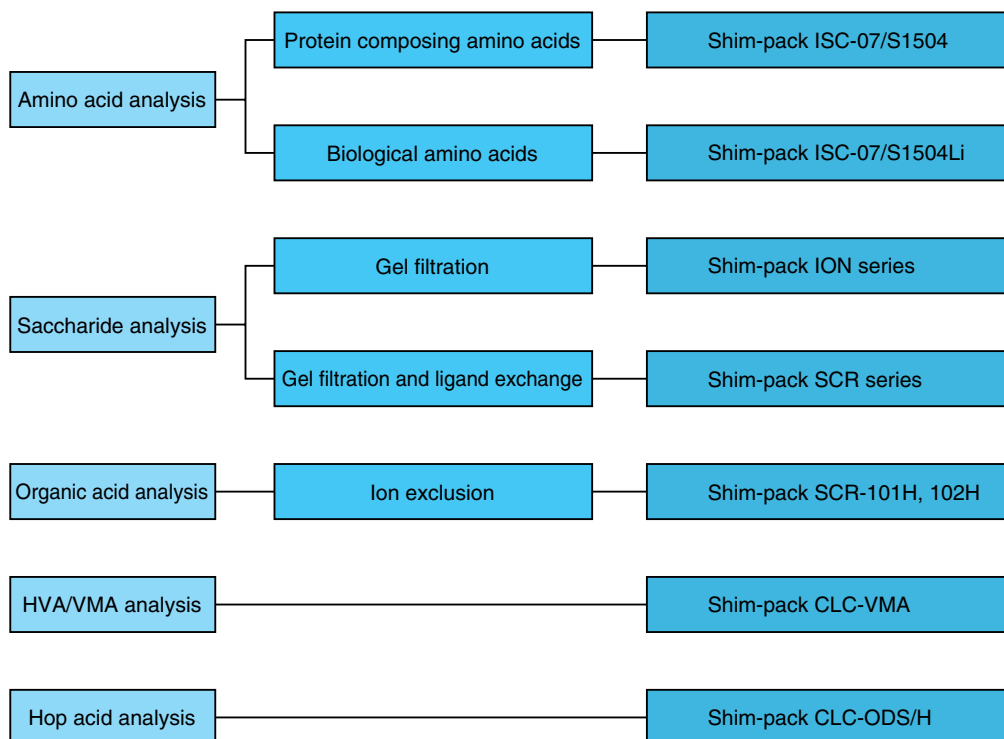
Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack GRD-ODS	Octadecyl group	30~50	4.0mm i.d.×25cm	228-16557-91
Pre-column Diol	Diol group	10	4.0mm i.d.×5cm	228-16367-91

### Packing materials of GRD-ODS

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Weight	Cat. No.
Gel GRD-ODS	Octadecyl group	30~50	10g	228-16831-91

## Dedicated Shim-pack Columns

● The Shim-pack Series columns may be classified as follows, according to the applicable type of samples.



Applications of the Shim-pack CLC-VMA column and the Shim-pack CLC-ODS/H column

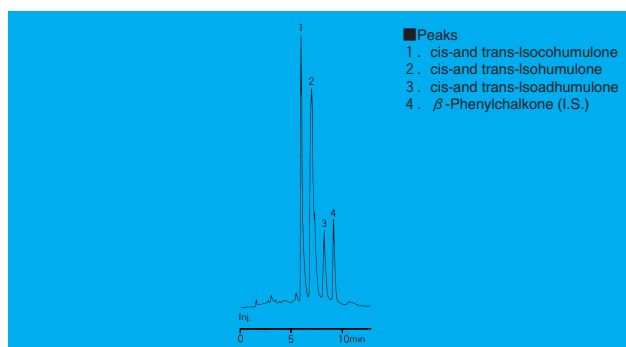


Fig. 80 Analysis of Hop Acids

■ Operational Conditions  
**Column:** Shim-pack CLC-ODS/H (4.6mm i.d.×25cm)  
**Mobile phase:** Methanol/water/phosphoric acid/10% tetra ethyl ammonium (77.5/22.5/1.1/3)  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (270nm)

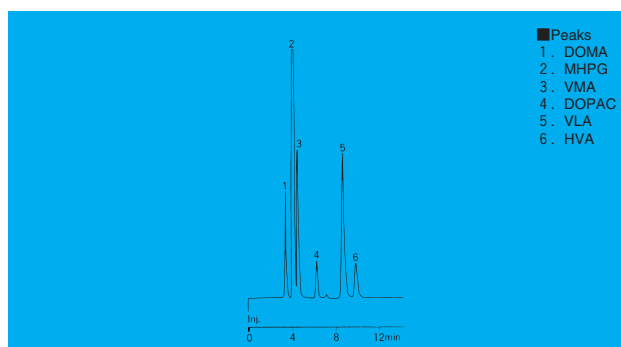


Fig. 81 Analysis of Catecholamine Metabolites

■ Operational Conditions  
**Column:** Shim-pack CLC-VMA (6.0mm i.d.×15cm)  
**Mobile phase:** 3mM tartaric acid/acetonitrile (7/3)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 40°C  
**Detector:** Fluorescence HPLC monitor (Ex. 280nm, Em. 320nm)

## Columns for Amino Acid Analysis

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack Amino-Na	Na type sulfone group	5	6.0mm i.d.×10cm	228-18837-91
Shim-pack Amino-Li	Li type sulfone group	5	6.0mm i.d.×10cm	228-18837-92
Shim-pack ISC-07/S1504Na	Na type sulfone group	7	4.0mm i.d.×15cm	228-09328-91
Shim-pack ISC-07/S1504Li	Li type sulfone group	7	4.0mm i.d.×15cm	228-00796-91

\* The following guard columns are available for amino acid analysis.

Column name	Dimensions	Cat. No.
ISC-07 guard column (Na type)	4.0mm i.d.×5cm	228-00802-91
ISC-07 guard column (Li type)	4.0mm i.d.×5cm	228-00797-91

\* The following column must be used in amino acid analysis.

Column name	Dimensions	Cat. No.
Shim-pack ISC-30/S0504Na (for trapping Na type ammonia)	4.0mm i.d.×5cm	228-14206-91
Shim-pack ISC-30/S0504Li (for trapping Li type ammonia)	4.0mm i.d.×5cm	228-00821-91

## Columns for Saccharide Analysis

Column name	Exclusion limit (dextran)	Dimensions	Cat. No.
Shim-pack ION KS-801	1×10 <sup>3</sup>	8.0mm i.d.×30cm	228-17894-91
Shim-pack ION KS-802	1×10 <sup>4</sup>	8.0mm i.d.×30cm	228-17895-91
Shim-pack ION KS-803	5×10 <sup>4</sup>	8.0mm i.d.×30cm	228-17896-91
Shim-pack ION KS-804	4×10 <sup>5</sup>	8.0mm i.d.×30cm	228-17897-91

\* The following dedicated guard column is available

Column name	Dimensions	Cat. No.
Shim-pack ION KS-800P (Common to ION KS series)	6.0mm i.d.×5cm	228-17898-91

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack SCR-101N	Na type sulfone group	10	7.9mm i.d.×30cm	228-07730-92
Shim-pack SCR-101C	Ca type sulfone group	10	7.9mm i.d.×30cm	228-17889-91
Shim-pack SCR-101P	Pb type sulfone group	10	7.9mm i.d.×30cm	228-17890-91

\* The following guard column must be used.

Column name	Dimensions	Cat. No.
SCR(N) guard column, dedicated to SCR-101N	4.0mm i.d.×5cm	228-09619-92
SCR(C) guard column, dedicated to SCR-101C	4.0mm i.d.×5cm	228-17891-91
SCR(P) guard column, dedicated to SCR-101P	4.0mm i.d.×5cm	228-17892-91

## Columns for Organic Acid Analysis

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack SCR-101H	H type sulfone group	10	7.9mm i.d.×30cm	228-07730-93
Shim-pack SCR-102H	H type sulfone group	7	8.0mm i.d.×30cm	228-17893-91

Column name	Dimensions	Cat. No.
SCR(H) guard column, dedicated to SCR-101H	4.0mm i.d.×5cm	228-09619-93
SCR-102H guard column, dedicated to SCR-102H	6.0mm i.d.×5cm	228-17924-91

## Columns for VMA/HVA Analysis

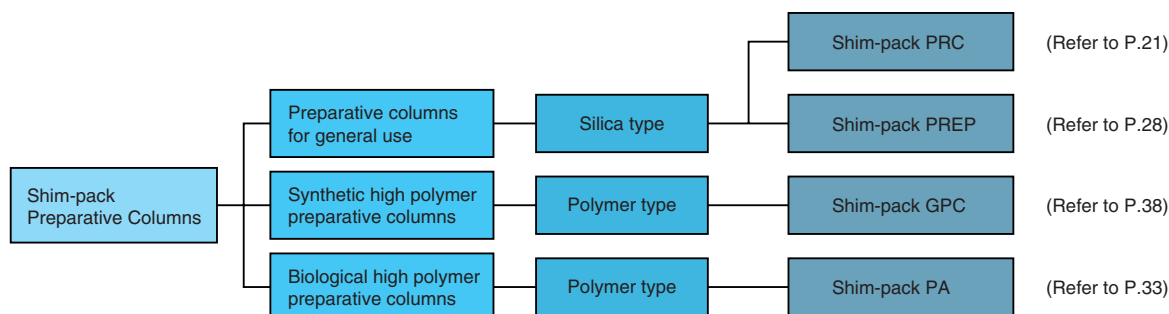
Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack CLC-VMA	Polar group	5	Reversed	6.0mm i.d.×15cm	228-17255-91

## Columns for Hop Acid Analysis

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack CLC-ODS/H	Octadecyl group	5	Reversed	4.6mm i.d.×25cm	228-00808-92

## Shim-pack Preparative Columns

- The Shim-pack preparative columns are available as follows.



Also STR (refer to P.70), Asahipak (refer to P.59) and CAPCELL PAK (refer to P.66) preparative columns are available.

### Standard values for maximum sample loads

When the sample load is increased beyond some maximum, extreme peak distortions can occur; the upper limit is the maximum acceptable sample load based on this peak distortion. The table below shows standard values of maximum sample loads for a semi-preparative column and preparative column each having a length of 250mm, in comparison with an ordinary analytical column. It should be noted that these standard values are applicable on condition that various requirements are satisfied: ion suppression for the desired component is achieved, the desired component is sufficiently soluble in the mobile phase, the type of sample injection solvent is considered, and other components do not elute near the target peak.

#### Standard Values of Maximum Sample Loads (column length is 250mm)

Column internal diameter (mm)	Sectional area (cm <sup>2</sup> )	Flow rate (ml min <sup>-1</sup> )	Maximum sample load (mg)
4.6	0.17	0.8	17
20.0	3.1	15	300
50.0	20	90	2000

#### Polymer type GPC Preparative Columns

##### Shim-pack GPC-2000 series

Preparative columns of Shim-pack GPC-800 series (Tetrahydrofuran)

Column name	Exclusion limit (polystyrene)	Dimensions	Cat. No.
Shim-pack GPC-2001	1.5×10 <sup>3</sup>	20mm i.d.×30cm	228-23342-91
Shim-pack GPC-2002	5×10 <sup>3</sup>	20mm i.d.×30cm	228-23342-92
Shim-pack GPC-20025	2×10 <sup>4</sup>	20mm i.d.×30cm	228-23342-93
Shim-pack GPC-2003	7×10 <sup>4</sup>	20mm i.d.×30cm	228-23342-94
Shim-pack GPC-2000P	(guard column)	8.0mm i.d.×5cm	228-20812-94

##### Shim-pack GPC-2000 series

Preparative columns of Shim-pack GPC-800C series (Chloroform)

Column name	Exclusion limit (polystyrene)	Dimensions	Cat. No.
Shim-pack GPC-2001C	1.5×10 <sup>3</sup>	20mm i.d.×30cm	228-23343-91
Shim-pack GPC-2002C	5×10 <sup>3</sup>	20mm i.d.×30cm	228-23343-92
Shim-pack GPC-20025C	2×10 <sup>4</sup>	20mm i.d.×30cm	228-23343-93
Shim-pack GPC-2003C	7×10 <sup>4</sup>	20mm i.d.×30cm	228-23343-94
Shim-pack GPC-2000CP	(guard column)	8.0mm i.d.×5cm	228-20812-95

## ■ Polymer Type Ion Exchange Preparative Column

### ■ Shim-pack PA

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack PA-DEAE	Diethylaminoethyl group	10	20mm i.d.×10cm	228-20758-92
Shim-pack PA-QA	Tetramethyl ammonium group	10	20mm i.d.×10cm	228-20759-92
Shim-pack PA-CM	Carboxymethyl group	10	20mm i.d.×10cm	228-20760-92
Shim-pack PA-SP	Sulfopropyl group	10	20mm i.d.×10cm	228-20761-92

### ■ Guard Column for Shim-pack PA

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack PAG-DEAE	Diethylaminoethyl group	10	8.0mm i.d.×1cm	228-20762-91
Shim-pack PAG-QA	Tetramethyl ammonium group	10	8.0mm i.d.×1cm	228-20763-91
Shim-pack PAG-CM	Carboxymethyl group	10	8.0mm i.d.×1cm	228-20764-91
Shim-pack PAG-SP	Sulfopropyl group	10	8.0mm i.d.×1cm	228-20765-91

## ■ Preparative column for general use

### ■ Shim-pack PRC

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	Cat. No.
Shim-pack PRC-SIL	Silica	15	Adsorption	20mm i.d.×25cm	228-23461-93
Shim-pack PRC-SIL(K)				30mm i.d.×25cm	228-23461-94
Shim-pack PRC-SIL(L)		50mm i.d.×25cm		228-23461-95	
Shim-pack PRC-SIL(H)		20mm i.d.×25cm		228-23461-91	
Shim-pack PRC-ODS	Octadecyl group	15	Reversed	20mm i.d.×25cm	228-23464-93
Shim-pack PRC-ODS(K)				30mm i.d.×25cm	228-23464-94
Shim-pack PRC-ODS(L)		50mm i.d.×25cm		228-23464-95	
Shim-pack PRC-ODS(H)		20mm i.d.×25cm		228-23464-91	
Shim-pack PRC-C <sub>8</sub>	Octyl group	15	Reversed	20mm i.d.×25cm	228-24381-93
Shim-pack PRC-C <sub>8</sub> (H)		5		20mm i.d.×25cm	228-24381-91
Shim-pack PRC-TMS	Trimethyl group	15	Reversed	20mm i.d.×25cm	228-24382-93
Shim-pack PRC-TMS(H)		5		20mm i.d.×25cm	228-24382-91
Shim-pack PRC-TMS	Aminopropyl group	15	Reversed, normal, ion exchange	20mm i.d.×25cm	228-24383-93
Shim-pack PRC-NH <sub>2</sub> (H)		5		20mm i.d.×25cm	228-24383-91
Shim-pack PRC-CN	Cyanopropyl group	15	Reversed-phase, normal phase	20mm i.d.×25cm	228-24384-93
Shim-pack PRC-CN(H)		5		20mm i.d.×25cm	228-24384-91

### ■ PRC guard column

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	Cat. No.
Shim-pack GPRC-SIL	Silica	5	8mm i.d.×1.5cm	228-23462-92
Shim-pack GPRC-ODS	Octadecyl group	5	8mm i.d.×1.5cm	228-23465-92
Shim-pack GPRC-C <sub>8</sub>	Octyl group	5	8mm i.d.×1.5cm	228-24386-92
Shim-pack GPRC-TMS	Trimethyl group	5	8mm i.d.×1.5cm	228-24387-92
Shim-pack GPRC-NH <sub>2</sub>	Aminopropyl group	5	8mm i.d.×1.5cm	228-24388-92
Shim-pack GPRC-CN	Cyanopropyl group	5	8mm i.d.×1.5cm	228-24389-92
Shim-pack GPRC-SIL(K)	Silica	15	30mm i.d.×7.5cm	228-23462-93
Shim-pack GPRC-SIL(L)	Silica	15	50mm i.d.×5cm	228-23462-94
Shim-pack GPRC-ODS(K)	Octadecyl group	15	30mm i.d.×7.5cm	228-23465-93
Shim-pack GPRC-ODS(L)	Octadecyl group	15	50mm i.d.×5cm	228-23465-94



## Preparative purification of Paeoniflorin in paeony root

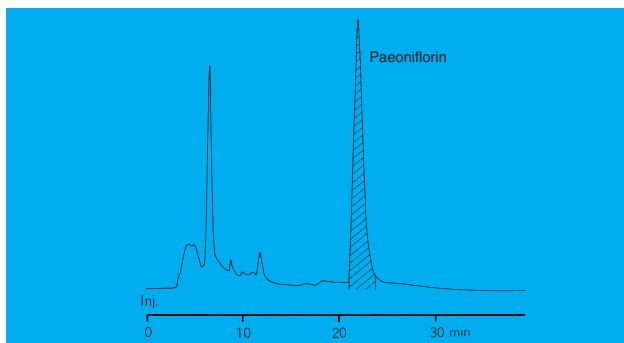


Fig. 82

### Large scale preparative conditions

**Sample:** Paeonyroot extract, 200mL, pumped-injection  
**Column:** Shim-pack PREP-ODS (L) (50mm i.d.×25cm)  
**Mobile phase:** Acetonitriles/water (1/6)  
**Flow rate:** 100mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (230nm, 0.5mL cell)

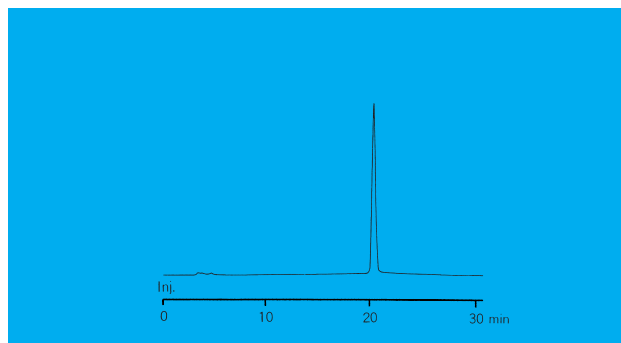


Fig. 83

### Fraction purity test condition

**Sample:** Fraction 5 μL  
**Column:** Shim-pack CLC-ODS (M) (4.6mm i.d.×25cm)  
**Mobile phase:** Acetonitriles/water (1/6)  
**Flow rate:** 0.8mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (230nm, 10mmL. cell)

## Preparative purification of Paeonol in Moutan Bark

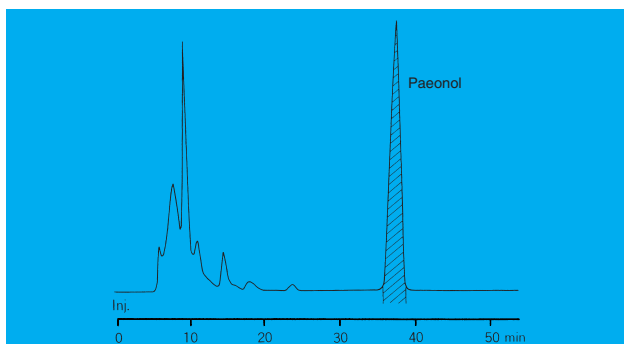


Fig. 84

### Large scale preparative conditions

**Sample:** Moutan bark extract, 200mL, pumped-injection  
**Column:** Shim-pack PREP-ODS (L) (50mm i.d.×25cm, 15 μm)  
**Mobile phase:** 0.1%(v/v)Acetate water/Acetonitriles (3/2)  
**Flow rate:** 60mL/min.  
**Detector:** UV spectrophotometric detector (254nm, 0.2mL cell)

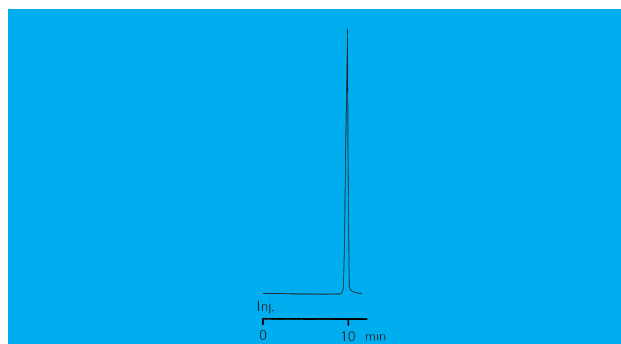


Fig. 85

### Fraction purity test condition

**Sample:** Fraction 10 μL  
**Column:** STR ODS-H (4mm i.d.×15cm, 5 μm)  
**Mobile phase:** 0.1%(v/v)Acetate water/Acetonitriles (3/2)  
**Flow rate:** 0.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (254nm)

## Preparative purification of Peptides

When separating peptides in reversed phase chromatography, changing mobile phase into acidic makes it possible to restrain the sticking of peptides to the surface of the silica gel. In case of preparative purification, the point is that evaporating a fraction easily to make the mobile phase acidic using trifluoroacetic acid and acetate

### Preparative Scale Purification of Peptides

Column	Reversed phase TMS(C <sub>1</sub> ), C <sub>4</sub> , C <sub>8</sub> , ODS(C <sub>18</sub> ) (size-exclusion or ion exchange column, as required)
Mobile phase	Acetonitrile/water (or 2-propanol), in the presence of 0.1 to 1%(v/v) trifluoroacetic acid, formic acid or acetic acid.
Detector	210-230nm
Fraction concentration	Evaporator (organic solvent) → freeze-drying (water) → storage at -20°C

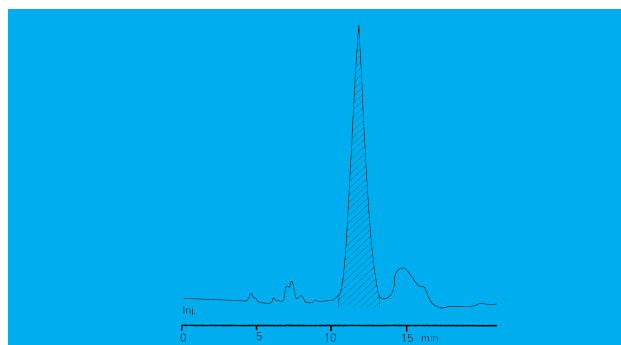
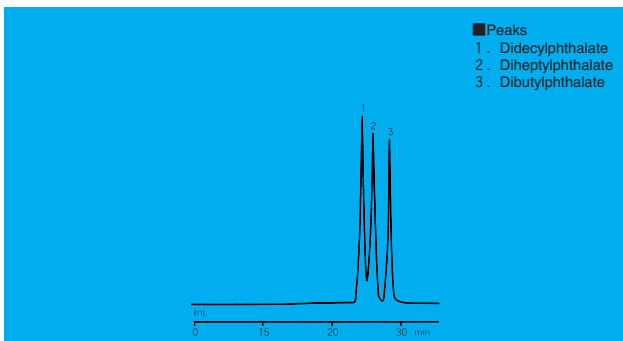


Fig. 86 Preparative purification of synthetic Peptides

**Sample:** Synthetic peptide (10 amino acids) 100mg/10mL aqueous solution  
**Column:** Shim-pack PREP-ODS(H)kit (20mm i.d.×250mm L.)  
**Mobile phase:** A: 0.1%(v/v) aqueous trifluoroacetic acid  
 B: 0.1%(v/v) solution of trifluoroacetic acid in acetonitrile  
 Gradient B 8 → 18% (20 min)  
**Flow rate:** 15mL/min.  
**Column temperature:** Ambient  
**Detector:** 220nm

## ■ Preparative in using GPC column



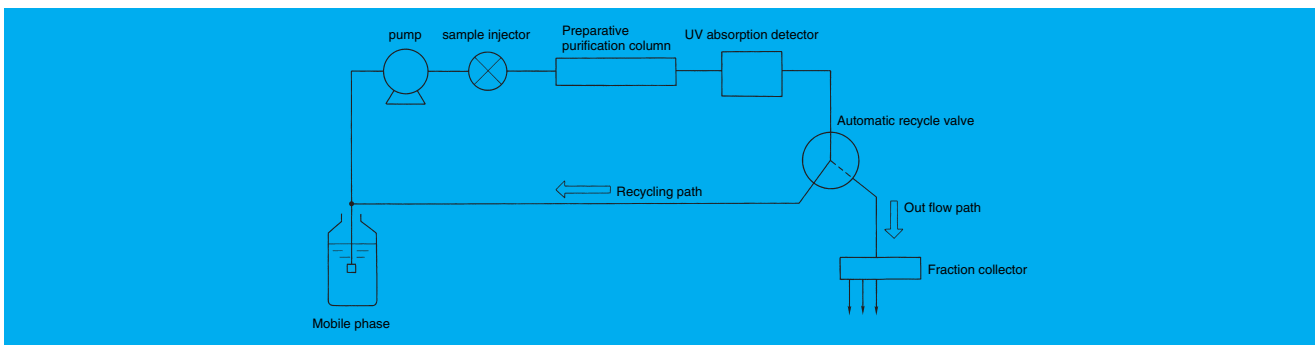
**Fig. 87 Separation of Esters phthalate in using GPC column**

**Column:** Shim-pack GPC 2001C (20mm i.d.×30cm)×2  
**Mobile phase:** Chloroform  
**Flow rate:** 3mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (260nm, 0.5mL. cell)

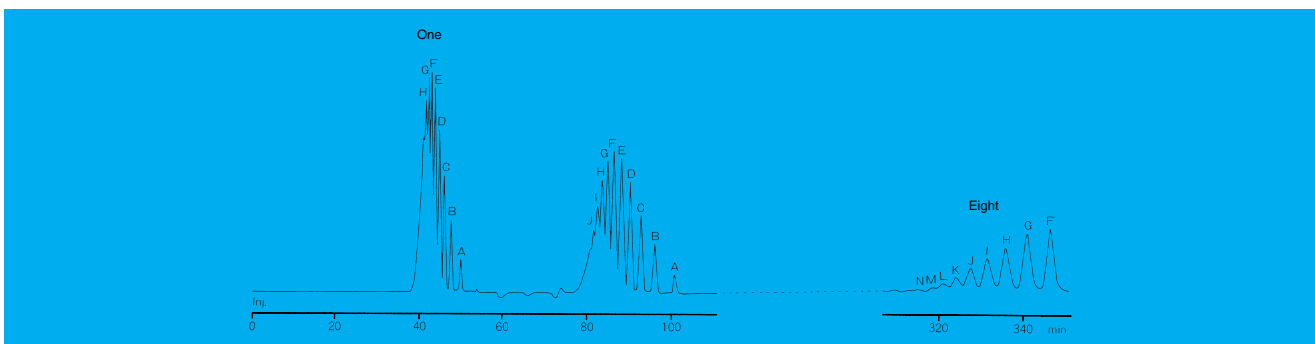
Various HPLC conditions must be accepted to increase the sample load. One of these conditions, mobile phase that has high sample solubilities should be used. In this point, GPC using chloroform as mobile phase has an advantage rather than reversed phase chromatography. With reference to the separation of low molecular, GPC is inferior to reversed phase chromatography. However, recycling method might be used to improve separations when the difference of the molecular size can be noticed and peak top is separated.

## ■ Recycling in preparative purification

Since preparative purification columns become more expensive as their size increases, it may be inevitable to use a column of such length that high resolution is not ensured. In these cases, an enhanced separating effect can be obtained as if the column length were increased by re-introducing the eluate band containing the desired component eluted from the column back into the column inlet. Figure 88 shows a flow chart for the recycling procedure. Switching the recycle valve permits choice of the out flow path leading to the fraction collector and the recycling path leading to the pump inlet. In the recycling mode, no additional mobile phase is consumed at all.



**Fig. 88 Flow Chart of Recycling System**



**Fig. 89 Recycle separation chromatogram of Styrene oligama**

**Sample:** Polystyrene molecular weight marker (molecular weight 760) 1%(W/V) chloroform solution 0.5mL injection  
**Column:** Shim-pack GPC-20025C+2002C+2001C  
**Mobile phase:** Chloroform  
**Column temperature:** Ambient  
**Detector:** Refractive index detector (128×12<sup>nd</sup>RIUFS)

## Shim-pack BIO(T)

- The Shim-pack BIO(T) series columns, using titanium column tubing, have been developed for biocompatible HPLC system, e.g. the Shimadzu LC-7A System.
- The Shim-pack BIO(T) series columns are available in three types: reversed phase, ion exchange, and hydrophobic.

### Merits of Titanium Column

In the HPLC of biological samples, corrosive liquids such as sodium chloride solution and hydrochloric acid are often used as the mobile phase.

Titanium is far more resistant against corrosion by sodium chloride solution and hydrochloric acid, especially against that by halogen ions, than the 316 stainless steel which is generally used in HPLC systems.

The Shim-pack BIO(T) series is recommended for such cases.

### 〈Corrosion Resistancy Comparison of Titanium and Stainless Steel〉

Solvent	Concentration (%)	Temperature (°C)	316 stainless steel	Pure titanium
Hydrochloric acid	1	25	◎	◎
	10	25	×	○
Sulfuric acid	1	25	◎	◎
	10	25	○	○
Nitric acid	25	boiling	◎	◎
	65	boiling	○	◎
Acetic acid	10	boiling	◎	◎
	60	boiling	○	◎
Formic acid	10	25	○	◎
Sodium chloride	25	25	○	◎
Ammonium chloride	40	25	○	◎
Zinc chloride	20	25	○	◎
Ferrous chloride	30	25	×	◎
Sodium hypochlorite	5	25	△	◎

◎ : Less than 0.125mm/year  
△ : 0.5 ~ 1.25mm/year

○ : 0.125 ~ 0.5mm/year  
× : More than 1.25mm/year

### ■ Shim-pack BIO(T) Series

Column name	Separation mode	Particle dia. (μm)	Pore diameter (Å)	Dimensions	Cat. No.
Shim-pack CLC-ODS(T)	Reversed	5	100	4.6mm i.d.×15cm	228-18062-91
Shim-pack WAX-1T	Anion exchange	3	100	4.6mm i.d.×5cm	228-18257-91
Shim-pack WAX-2T	Anion exchange	5	300	4.6mm i.d.×5cm	228-18258-91
Shim-pack WCX-1T	Cation exchange	5	300	4.6mm i.d.×5cm	228-18259-91
Shim-pack HPC-C2T	Hydrophobic	5	300	4.6mm i.d.×5cm	228-18260-91
Shim-pack HPC-C3T	Hydrophobic	5	300	4.6mm i.d.×5cm	228-18261-91

## Shim-pack BIO(T)

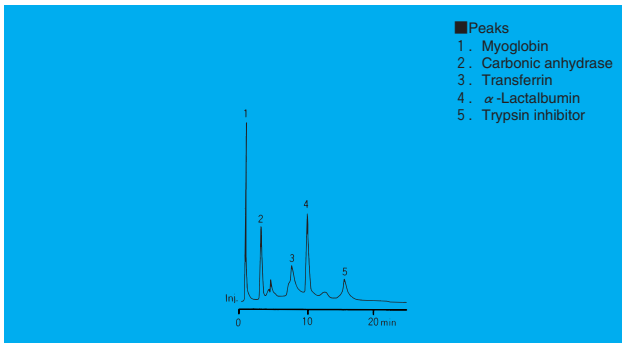


Fig. 90 Analysis of Protein Standards

■Operational Conditions  
**Column:** Shim-pack WAX-2T (4.6mm i.d.×5cm)  
**Mobile phase:** Tris-hydrochloric acid buffer solution (pH 8.0) /sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

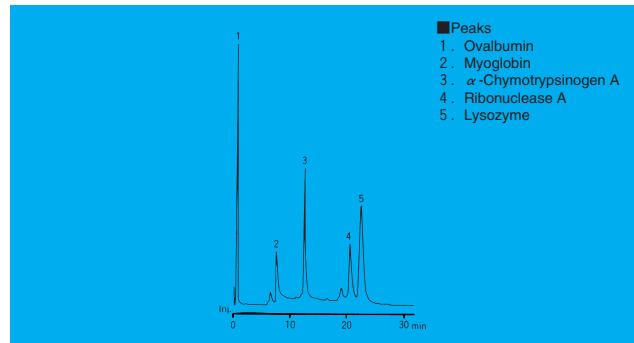


Fig. 91 Analysis of Protein Standards

■Operational Conditions  
**Column:** Shim-pack WCX-1T (4.6mm i.d.×5cm)  
**Mobile phase:** Phosphate buffer solution (pH 6.0)/sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (280nm)

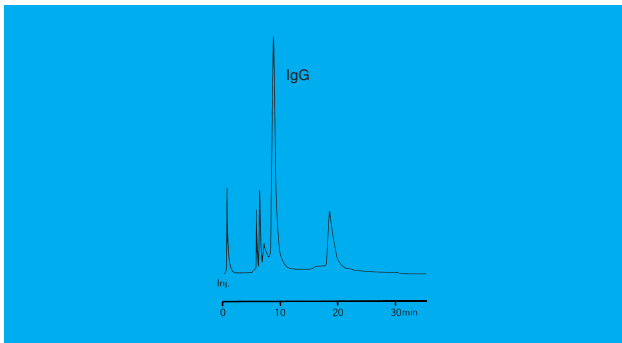


Fig. 92 Determination of IgG Monoclonal Antibody in Mouse Ascites Fluid

■Operational Conditions  
**Column:** Shim-pack WAX-2T (4.6mm i.d.×5cm)  
**Mobile phase:** Tris-hydrochloric acid buffer solution (pH 8.0) /sodium chloride, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (280nm)

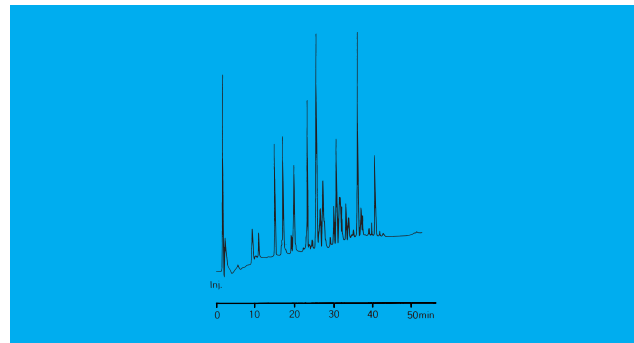


Fig. 93 Analysis of Decomposition Products of Trypsin in Cytoch C

■Operational Conditions  
**Column:** Shim-pack CLC-ODS(T) (4.6mm i.d.×5cm)  
**Mobile phase:** 0.1% aqueous solution of TFA/0.1% acetonitrile solution of TFA, gradient elution  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (215nm)

## Asahipak® ODP

- The Asahipak ODP column is packed with synthetic hard polymer particles on which octadecyl groups are chemically bonded and is used for reversed phase LC.
- Having no silanol groups, this column is especially effective for the HPLC of basic compounds, e.g. some drugs.
- The column is stable in a pH range of 2~13. It can be washed with an alkaline solution.
- The gel has a minimum swelling and contraction, and so the column can be used in a wide mobile phase condition, from 100% water to 100% water-soluble organic solvent.
- High repeatability in gradient LC.

### Asahipak® ODP

Column name		Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak ODP-50	4D	C <sub>18</sub> group	5	4.6mm i.d.×15cm	406-020
	4E			4.6mm i.d.×25cm	406-021
	6D			6.0mm i.d.×15cm	406-017
	6E			6.0mm i.d.×25cm	406-018

### Guard Column (Analytical)

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak ODP-50G	C <sub>18</sub> group	5	4.6mm i.d.×1cm	406-022
			6.0mm i.d.×1cm	406-019

### Guard Column (Analytical)

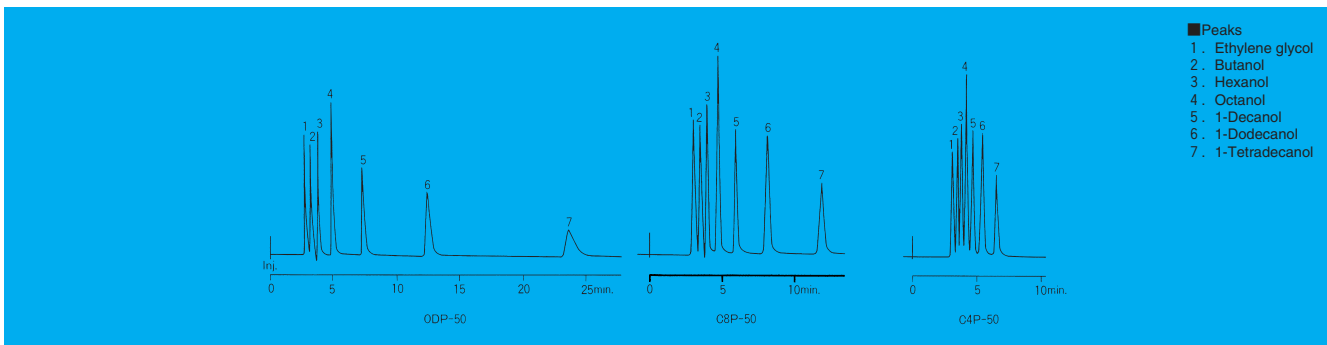
Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak C8P-50	C <sub>8</sub> group	5	4.6mm i.d.×15cm	406-023
			4.6mm i.d.×25cm	406-024
Asahipak C8P-50G			4.6mm i.d.× 1cm	406-025

### Asahipak® C4P

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak C4P-50	C <sub>4</sub> group	5	4.6mm i.d.×15cm	406-026
			4.6mm i.d.×25cm	406-027

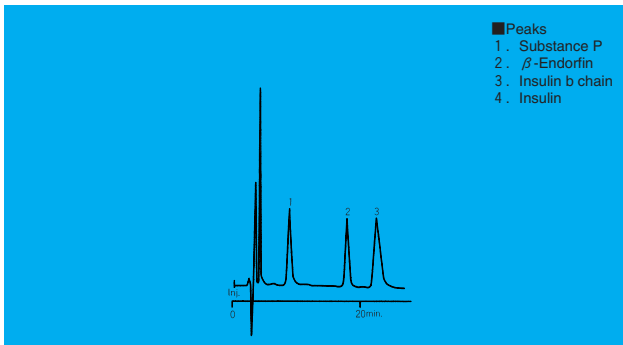
### Guard Column (Analytical)

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak C4P-50G	C <sub>4</sub> group	5	4.6mm i.d.×1cm	406-028



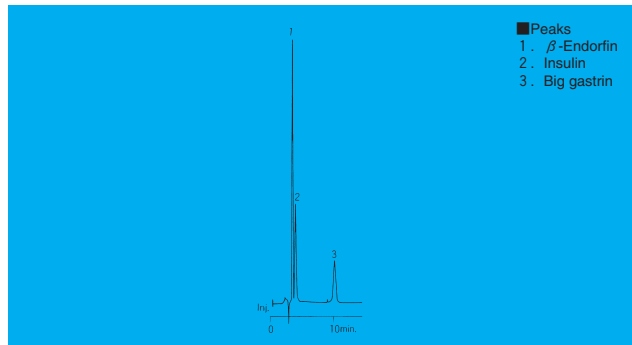
**Fig. 94 Separation of Alkylalcohols Group**

**Operational Conditions**  
**Column:** Asahipak ODP-50, C8P-50, C4P-50 (4.6mm i.d.×15cm)  
**Mobile phase:** Methanol 80/water 20  
**Flow rate:** 0.6mL/min.  
**Column temperature:** 30°C  
**Detector:** Refractive index detector



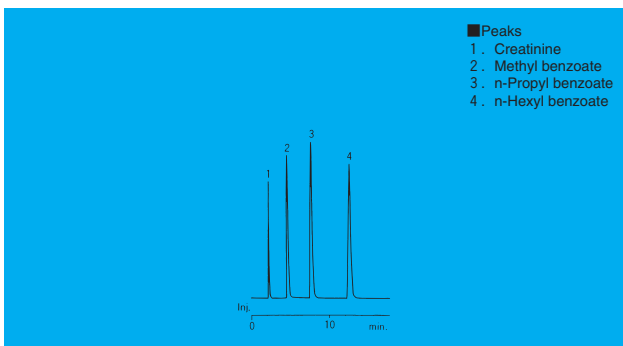
**Fig. 95 Separation of Macromolecule Peptide**

**Operational Conditions**  
**Column:** Asahipak C8P-50, (4.6mm i.d.×15cm)  
**Mobile phase:** 0.05%TFA (pH=2.3) 73/Acetonitriles 27  
**Flow rate:** 0.6mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (220nm)



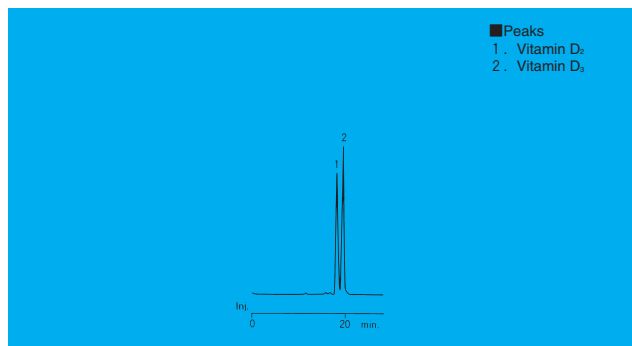
**Fig. 96 Separation of Macromolecule Peptide**

**Operational Conditions**  
**Column:** Asahipak C4P-50, (4.6mm i.d.×15cm)  
**Mobile phase:** 0.05%TFA (pH=2.3) 72/Acetonitriles 28  
**Flow rate:** 0.6mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (220nm)



**Fig. 97 Analysis of Standard Samples**

**Operational Conditions**  
**Column:** Asahipack ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitrile/water (7/3)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (254nm)



**Fig. 98 Separation of Vitamin D<sub>2</sub> and D<sub>3</sub>**

**Operational Conditions**  
**Column:** Asahipak ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitrile/water (95/5)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (260nm)

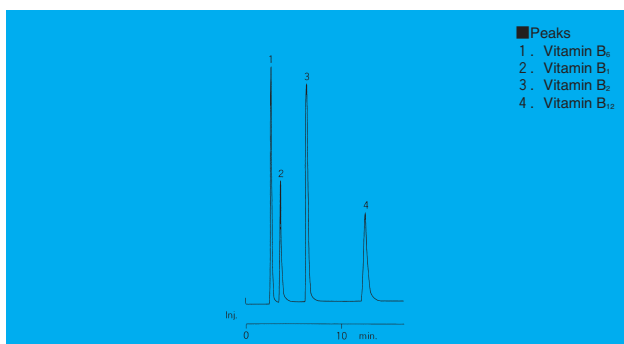


Fig. 99 Separation of Vitamins B Group

■ Operational Conditions

**Column:** Asahipak ODP-50, (6.0mm i.d.×15cm)  
**Mobile phase:** Acetonitriles 1/50mM phosphoric acid sodium  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (254nm)

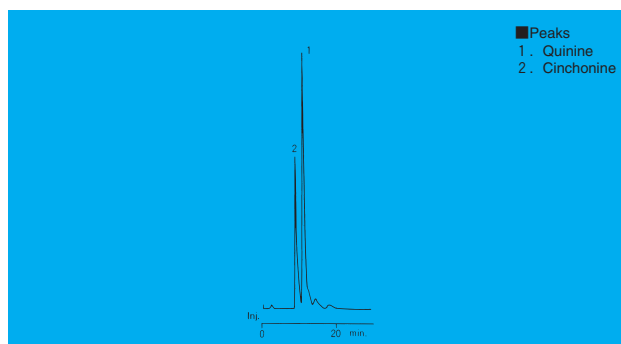


Fig. 100 Separation of Quinine and Cinchonine

■ Operational Conditions

**Column:** Asahipak ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** Methanol/acetonitrile/10mM sodium phosphate (2/1/1) (pH 9)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (250nm)

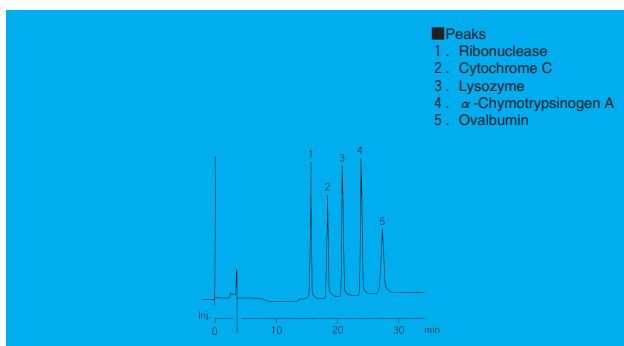


Fig. 101 Separation of Protein Standard

■ Operational Conditions

**Column:** Asahipak ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** Solvent A: Water and 0.05% TFA  
 Solvent B: Acetonitrile and 0.05% TFA  
 A/B=Linear gradient from 90/10 to 40/60, 30minutes  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (280nm)

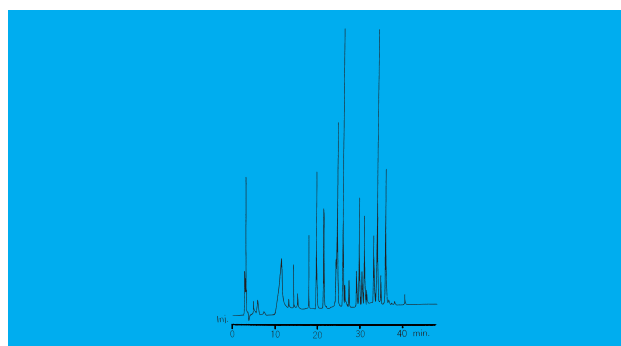


Fig. 102 Analysis of Decomposition Products of Trypsin in Cytochrome C

■ Operational Conditions

**Column:** Asahipak ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** Solvent A: Water and 0.1% TFA  
 Solvent B: Acetonitrile and 0.1% TFA  
 Linear gradient from solvent A to solvent B in 100minutes.  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 25°C  
**Detector:** UV spectrophotometric detector (215nm)

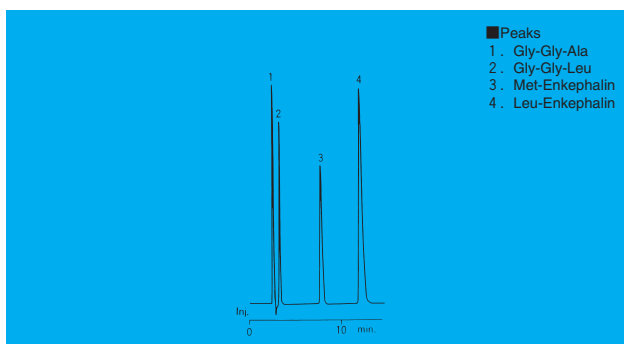


Fig. 103 Analysis of Peptide Mixture

■ Operational Conditions

**Column:** Asahipak ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** 0.05% TFA/acetonitrile (4/1)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (210nm)

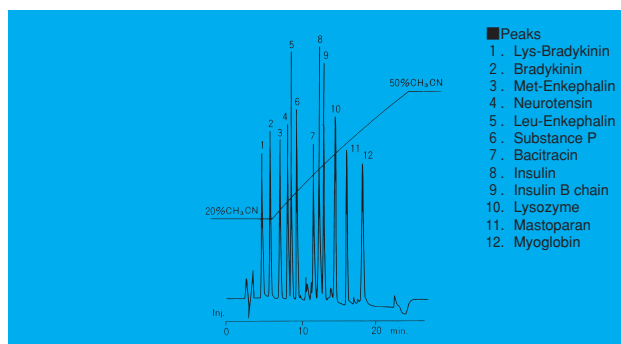


Fig. 104 Analysis of Peptide Mixture

■ Operational Conditions

**Column:** Asahipak ODP-50 (6.0mm i.d.×15cm)  
**Mobile phase:** Solvent A: 0.05% TFA/acetonitrile (4/1)  
 Solvent B: 0.05% TFA/acetonitrile (1/1)  
 Linear gradient from solvent A to solvent B in 20 minutes  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** UV spectrophotometric detector (220nm)

## Asahipak® GS

- The Asahipak GS series columns are packed with hard gel particles made of vinyl alcohol polymer.
- The allowable pH range of mobile phase is 2~12 for the GS-320 and GS-520, and 2~9 for the other columns.
- Various organic solvents can be used.
- Aqueous solutions of salts and buffer solutions can be used as mobile phase. (It is recommended that the concentration is below 0.5M.) Mobile phases that contain urea, guanidine HCl, or SDS (sodium dodecyl sulfate) are also applicable.

### High-Resolution Type (7.6mm i.d.×30cm)

Type	Column name	Average particle diameter (μm)	Exclusion limit (molecular weight) (Pullulan)	Number of theoretical plates	GLC Cat. No.
Applicable both to aqueous solutions and organic solvent	GF-310HQ	5	40,000	>16,000	406-323
	GF-510HQ	5	300,000	>16,000	406-324
	GF-710HQ	9	1,000,000	>10,000	406-325
	GF-7MHQ	7	1,000,000	>11,000	406-326
Applicable to aqueous solutions	GS-220HQ	6	3,000	>16,000	406-318
	GS-320HQ	6	40,000	>16,000	406-319
	GS-520HQ	7	300,000	>15,000	406-320
	GS-620HQ	7	2,000,000	>15,000	406-321

### Guard Columns

Column name	Dimensions	Grade	GLC Cat. No.
GF-1G 7B	7.6mm i.d.×10mm	310, 310HQ, 510, 510HQ 710, 710HQ, 700, 710HQ	406-327
GS-2G 7B	7.6mm i.d.×10mm	220, 220HQ, 320, 320HQ 520, 520HQ, 620, 620HQ	406-322



### ■ Applicable Organic Solvents

Solvent conditions		Applicability			
Solvent	Concentration	310, 510 710, 7M	320 520	220 620	GS-710
Aqueous solution	0~0.5M	●	●	●	●
Methanol	0~100%	●	●	Below 20%	●
Ethanol	0~100%	●	●	Below 20%	●
Acetonitrile	0~50%	●	●	●	Not applicable
	51~100%	●			
THF	0~100%	●			Not applicable
DMF	0~100%	●			
Acetone	0~100%	●			
Chloroform	0~100%	●			
DMSO	0~50%	●			
	51~100%	×			

### ■ Applications

310	Hydrophobic polymer, oligomers, monomers, surfactants, steroids, hydrophilic polymers
510	Proteins, nucleic acids, sugars, gelatins, hydrophobic/hydrophilic polymers
220	Oligosaccharides, peptides, hydrophilic, oligomers, monomers
320	Peptides, nucleic acid constituents, sugars, vitamins, blood metabolites
520	Plasma proteins, albumins, globulin, sugars, hydrophilic polymers, nucleic acids
620	Highly polymerated proteins, nucleic acids, gelatins
710	Polysaccharides, dextran, hydrophilic polymers

### ■ Calibration Curves

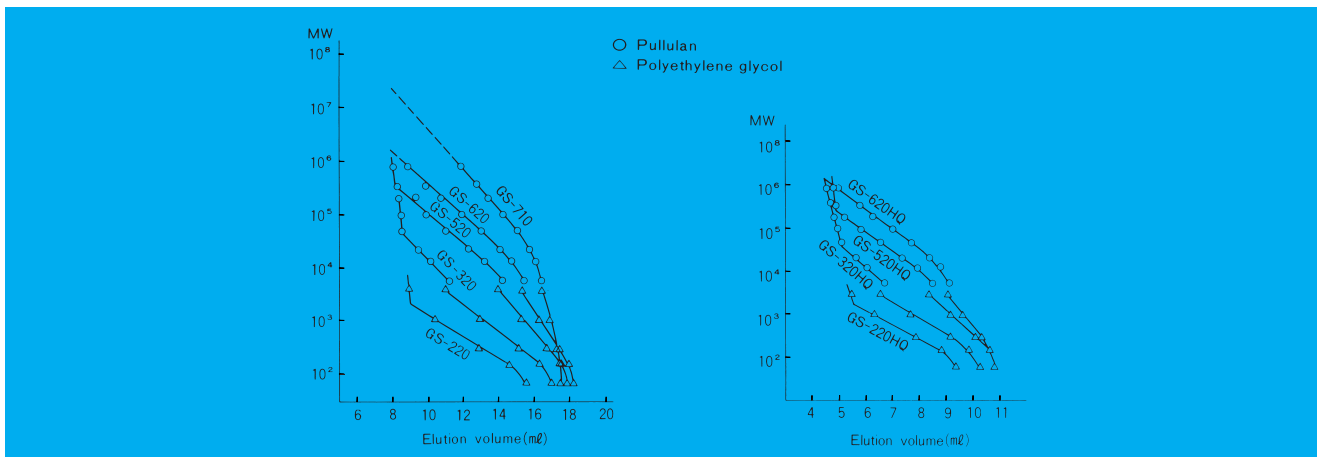


Fig. 105 Columns for Aqueous Mobile Phases

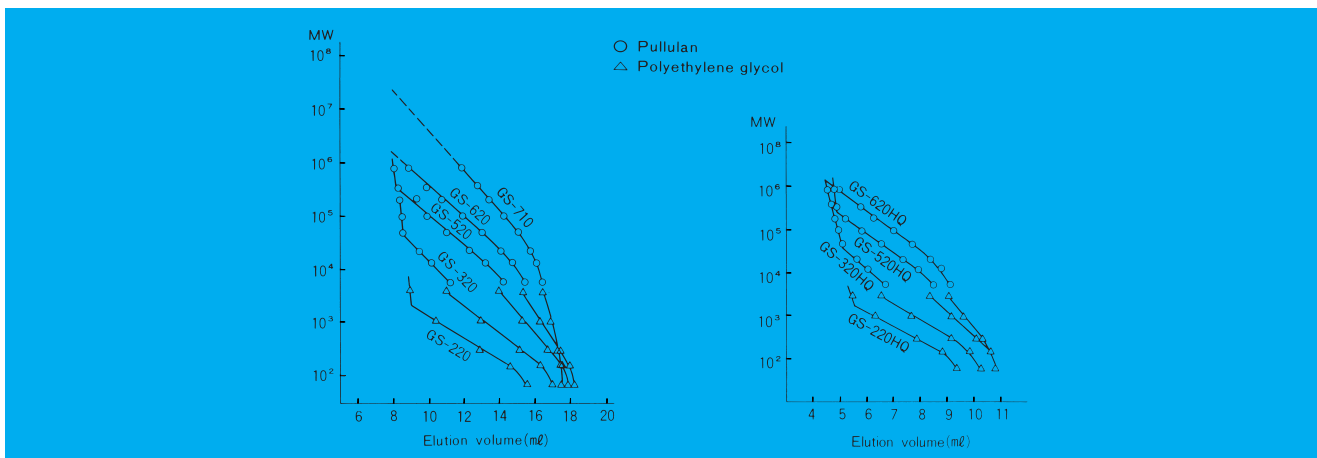


Fig. 106 Columns for Aqueous/Organic Solvent Mobile Phases

## Asahipak® ES

- The Asahipak ES series columns, developed for high-speed ion exchange chromatography, are packed with vinyl alcohol polymer particles into which ion exchange groups have been introduced.
- Since the gel particles have many ion exchange groups and alcoholic hydroxyl groups, hence have little hydrophobic adsorption, the columns are suitable for the separation of ionic biological compounds.
- The columns ensure high yield for proteins, enzymes, and nucleic acids.
- The allowable pH range of mobile phase is 2~12.
- The columns are available in two types, anion exchange type and cation exchange type.

### Analytical Column (7.6mm i.d.×10cm)

Column name	Average particle diameter (μm)	Ion exchange group	Ion exchange capacity (meq/g)	Number of theoretical plates	GLC Cat. No.
ES-502C	9±0.5	Carboxyl group	0.55±0.02	>3,000	406-109
ES-502N	9±0.5	Diethylamino ethyl group	0.55±0.02	>3,000	406-108

The use of preparative guard column GS-20G (228-18754-02) is recommended.

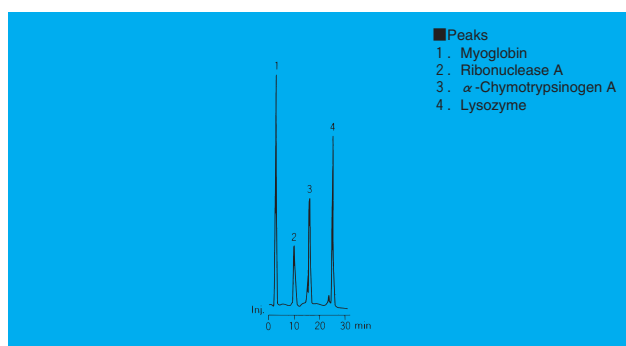


Fig. 107 Analysis of Protein Standard

#### Operational Conditions

**Column:** Asahipak ES-502C (7.6mm i.d.×10cm)  
**Mobile phase:** Solvent A: Sodium phosphate (pH 7.0)  
 Solvent B: 50mM sodium phosphate and 500mM sodium chloride (pH 7.0)  
 Linear gradient from solvent A to solvent B in 20 minutes  
**Flow rate:** 1.0mL/min.  
**Detector:** UV spectrophotometric detector (280nm)

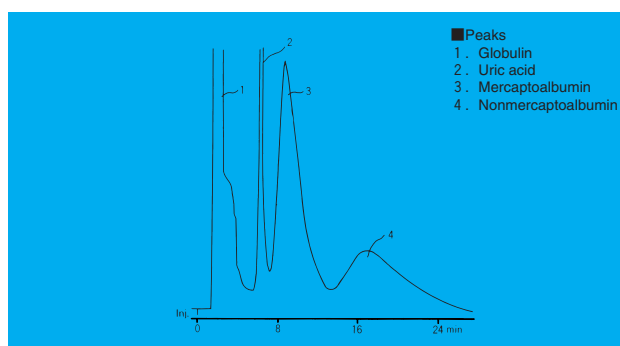


Fig. 108 Determination of Mercaptoalbumin and Nonmercaptoalbumin in Control Serum

#### Operational Conditions

**Column:** Asahipak ES-502N (7.6mm i.d.×10cm)  
**Mobile phase:** 50mM N-methyl piperazine, 400mM sodium sulfate, and 0.3% ethanol (pH 4.8)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 35°C  
**Detector:** UV spectrophotometric detector (280nm)

## Asahipak® NH2P

- The Asahipak NH2P series columns are for sugar analysis, packed with vinylalcoholcopolymer with polyamines chemically bonded.
- High number of theoretical plates and symmetrical peak can be acquired between monosaccharides and oligosaccharides.
- Available under the condition of the alkali (less than pH=13) and you can wash the column easily as well as getting high separation.
- The column is stable in a pH range of 2~13.

### Asahipak NH2P

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak NH2P-50 4E	NH <sub>2</sub> group	5	4.6mm i.d.×25cm	406-030

### Guard Column (Analytical)

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Dimensions	GLC Cat. No.
Asahipak NH2P-50G 4A	NH <sub>2</sub> group	5	4.6mm i.d.×1cm	406-031

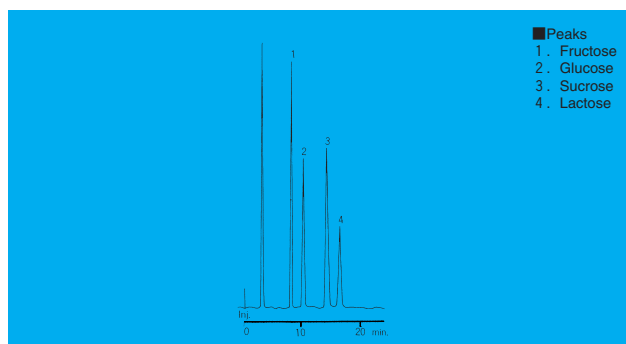


Fig. 109 Separation of Monosaccharides and Disaccharides

#### Operational Conditions

**Column:** Asahipak NH2P-50, (4.6mm i.d.×25cm)  
**Mobile phase:** Acetonitriles 75/Water 25  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** Refractive index detector

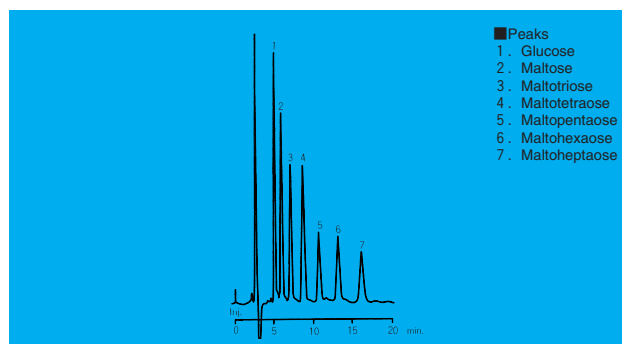


Fig. 110 Separation of Oligosaccharides

#### Operational Conditions

**Column:** Asahipak NH2P-50, (4.6mm i.d.×25cm)  
**Mobile phase:** Acetonitriles 60/Water 40  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** Refractive index detector

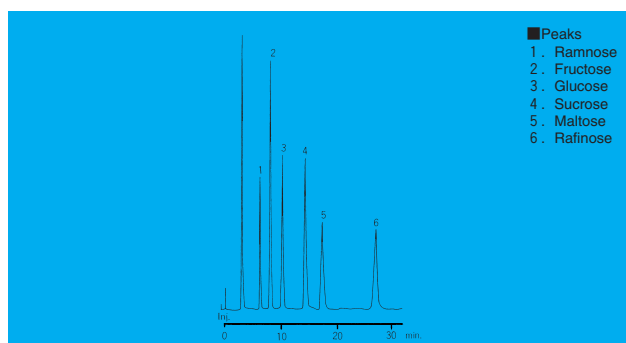


Fig. 111 Separation of Monosaccharides, Disaccharides and Trisaccharides

#### Operational Conditions

**Column:** Asahipak NH2P-50, (4.6mm i.d.×25cm)  
**Mobile phase:** Acetonitriles 75/Water 25  
**Flow rate:** 1.0mL/min.  
**Column temperature:** 30°C  
**Detector:** Refractive index detector

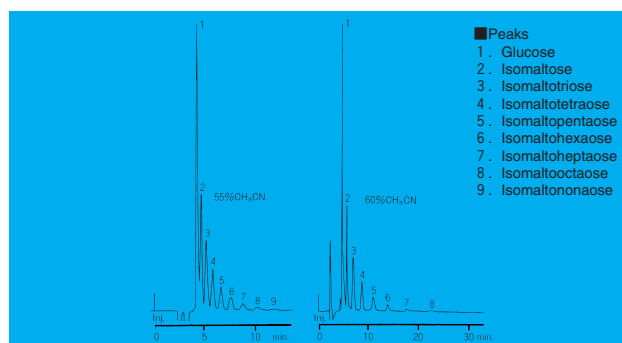


Fig. 112 Separation of Dextran hydrolysis substances

#### Operational Conditions

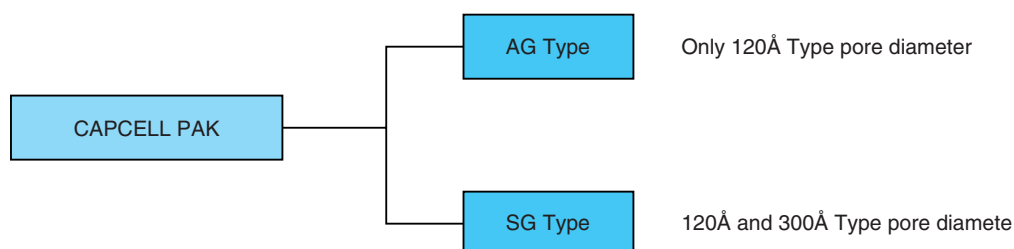
**Column:** Asahipak NH2P-50, (4.6mm i.d.×25cm)  
**Mobile phase:** Acetonitriles/Water  
**Flow rate:** 1.0mL/min.  
**Detector:** Refractive index detector

## CAPCELL PAK

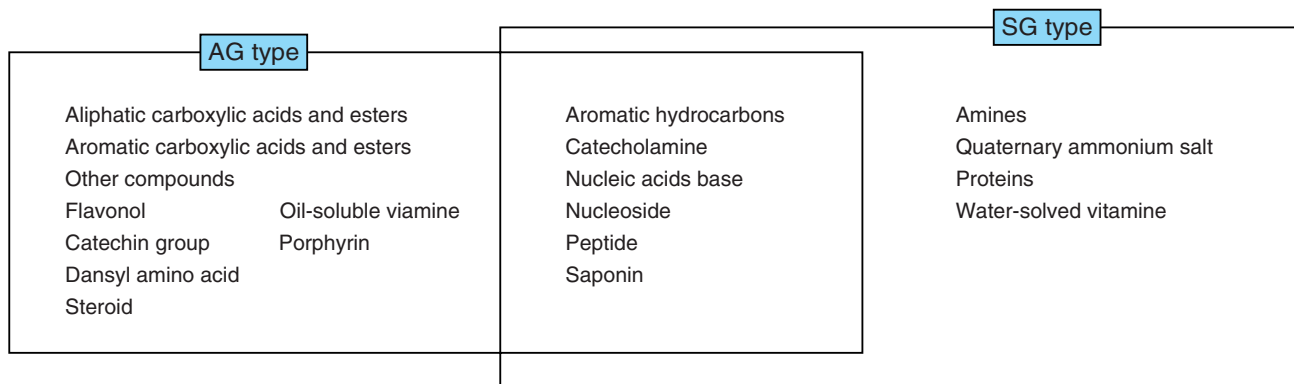
### (1) Features of CAPCELL PAK

As covered with silicones polymer on the surface of silica, superior in chemical stability rather than silica type materials.

### (2) CAPCELL PAK is classified as follows.



### (3) Selection of CAPCELL PAK columns



As for the amount of micro-metal involved in silica, SG type has fewer than AG type.

### ■ CAPCELL PAK AGType (C18 Type)

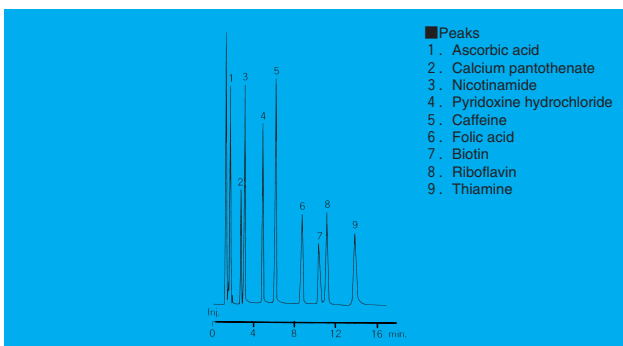
Stationary phase	Pore diameter (Å)	Particle dia. (μm)	Column i.d.	Column length (mm)	GLC Cat. No.	
C18	120	5	4.6	35	12501	
				150	12503	
				250	12504	
			6.0	35	12506	
				150	12508	
				50	15501	
			20	250	15504	
				30	250	16501

**■ CAPCELL PAK AGType (C8 Type)**

Stationary phase	Pore diameter (Å)	Particle dia. ( $\mu\text{m}$ )	Column i.d.	Column length (mm)	GLC Cat. No.
C8	120	5	4.6	35	21501
				150	21503
				250	21504
			6.0	35	21506
				150	21508
			20	50	25501
250	25503				

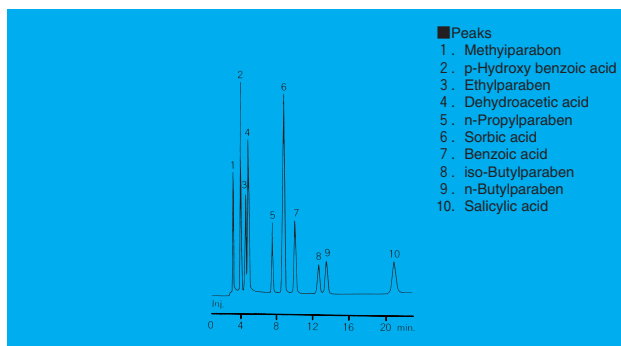
**■ CAPCELL PAK SGType (C18 Type)**

Stationary phase	Pore diameter (Å)	Particle dia. ( $\mu\text{m}$ )	Column i.d.	Column length (mm)	GLC Cat. No.
C18	120	5	4.6	35	12510
				150	12512
				250	12513
			6.0	35	13510
				150	13512
			20	50	16510
				250	16513
			30	250	16501
			C18	300	5
150	12522				
250	12523				
6.0	35	13520			
	150	13522			
20	50	16520			
	250	16523			



**Fig. 113 Separation of Aqueous Vitamine Group**

**Operational Conditions**  
**Column:** C<sub>18</sub> SG120 (4.6mm i.d.×25cm)  
**Mobile phase:** Water (pH2.1, H<sub>3</sub>PO<sub>4</sub>)/CH<sub>3</sub>CN =9/1, 1.5mM heptanesulfonic acid  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

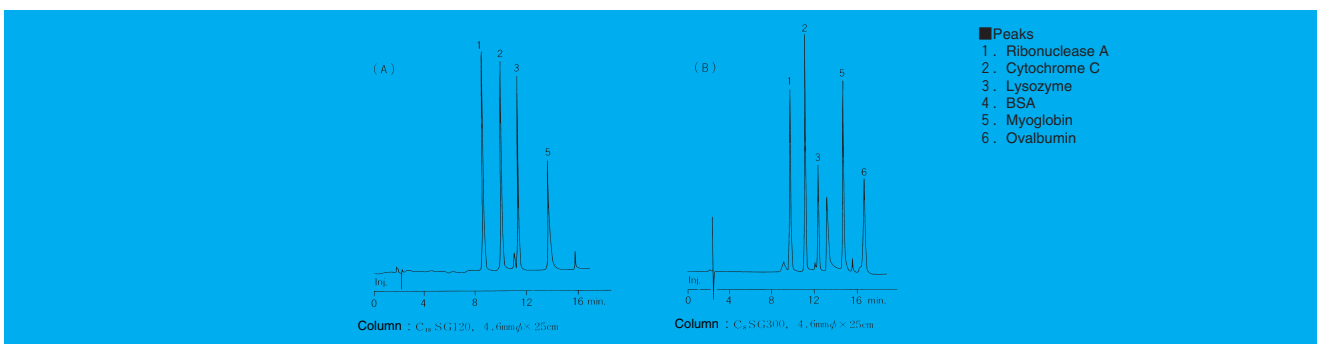


**Fig. 114 Separation of an Antiseptic Component**

**Operational Conditions**  
**Column:** C<sub>18</sub> SG120 (4.6mm i.d.×25cm)  
**Mobile phase:** 0.05M NaH<sub>2</sub>PO<sub>4</sub>(pH4.5)/MeOH/CH<sub>3</sub>CN=50/35/15, 4mM cetyltrimethylammonium chloride  
**Flow rate:** 1mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (235nm)  
**Injection volume:** 10 μL (5 μg/mL)

### CAPCELL PAK SG Type (C8 Type)

Stationary phase	Pore diameter (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	GLC Cat. No.
C8	120	5	4.6	35	21511
				150	21513
				250	21514
			6.0	35	21516
				150	21518
				250	25511
C8	300	5	20	50	25513
				250	25513
			4.6	35	21521
				150	21523
			6.0	250	21524
				35	21526
150	21528				
250	25521				
20	50	25521			
	250	25523			



**Fig. 115 Separation of Protein with SG type columns**

**Operational Conditions**  
**Mobile phase:** (A) 0.1% TFA/H<sub>2</sub>O (B) 0.1% TFA/CH<sub>3</sub>CN (B) 15% → 60% (15min) Gradient  
**Flow rate:** 1.5mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (214nm)

## Zorbax

- The Zorbax columns are high-performance columns developed by DuPont, the pioneer in HPLC.
- The columns are packed with fully-porous, spherical silica particles on which the respective stationary phases are chemically bonded. (Except Zorbax SIL)

### Zorbax

Column name	Stationary phase	Particle dia. ( $\mu\text{m}$ )	Separation mode	Dimensions	GLC Cat. No.
Zorbax SIL	Silica	5	Adsorption	4.6mm i.d. $\times$ 15cm	883952-701
				4.6mm i.d. $\times$ 25cm	880952-701
Zorbax ODS	Octadecyl group	5	Reversed	4.6mm i.d. $\times$ 15cm	883952-702
				4.6mm i.d. $\times$ 25cm	880952-702
Zorbax C <sub>8</sub>	Octyl group	5	Reversed	4.6mm i.d. $\times$ 15cm	883952-706
				4.6mm i.d. $\times$ 25cm	880952-706
Zorbax TMS	Trimethyl group	5	Reversed	4.6mm i.d. $\times$ 15cm	883952-710
				4.6mm i.d. $\times$ 25cm	880952-710
Zorbax NH <sub>2</sub>	Aminopropyl group	5	Reversed, normal, anion, exchange	4.6mm i.d. $\times$ 15cm	883952-708
				4.6mm i.d. $\times$ 25cm	880952-708
Zorbax CN	Cyanopropyl group	5	Reversed, normal	4.6mm i.d. $\times$ 15cm	883952-705
				4.6mm i.d. $\times$ 25cm	880952-705
Zorbax SAX	Quaternary ammonium group	5	Anion exchange	4.6mm i.d. $\times$ 15cm	883952-703
				4.6mm i.d. $\times$ 25cm	880952-703
Zorbax SCX-300	Sulfonic group	7	Cation exchange	4.6mm i.d. $\times$ 15cm	883952-704
				4.6mm i.d. $\times$ 25cm	880952-704

### Guard Column Pre-column Kit

In analyses with a Zorbax column, it is recommended to use a guard column or a pre-column so that a longer service life is expected of the Zorbax column.

The kit consists of an empty column, packing material, and a filter.

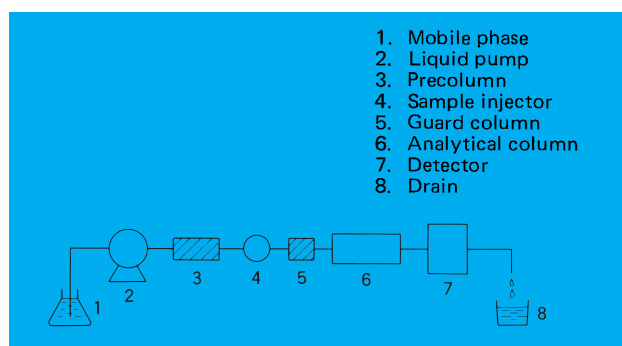


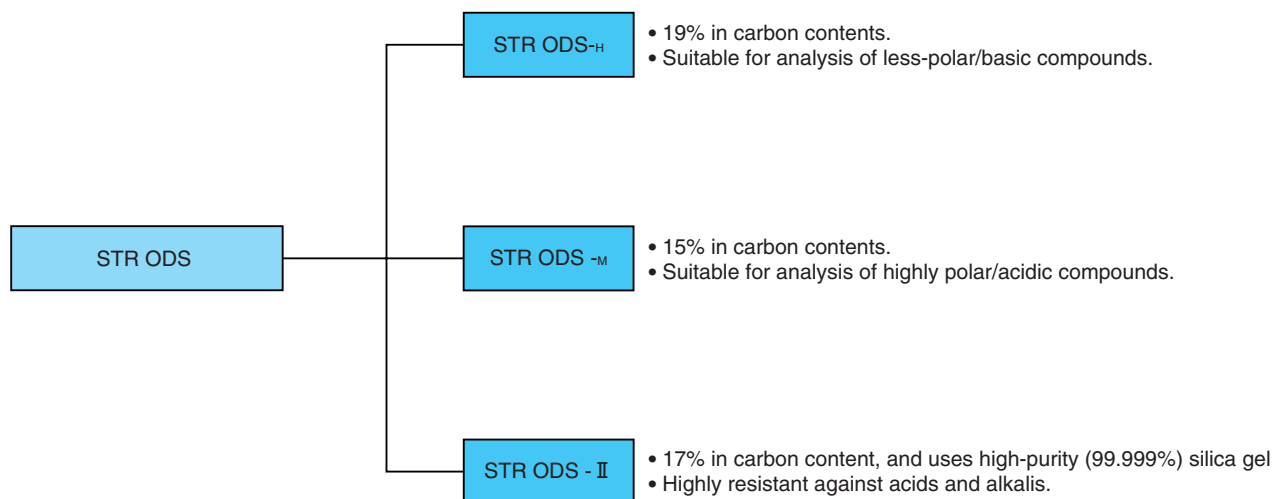
Fig. 116 Flow Line Having Guard Column and Pre-column

## STR ODS Series

(STR : Shimadzu Techno Research)

- STR ODS Series columns using octadecyl group as the stationary phase feature low cost and wide field of applications.

■ The STR ODS Series is available in the following three types:



### ■ The STR ODS-II Columns

#### ■ STR ODS Column for Analytical HPLC

Column name	Pore dia. (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	GLC Cat. No.
STR ODS-II	120	5	4.0	150	404-019
				250	404-018
			4.6	150	404-021
				250	404-020
STR ODS-II PEEK	120	5	4.6	150	404-027
				250	404-026

#### ■ STR ODS Column for Preparative HPLC

Column name	Pore dia. (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	GLC Cat. No.
STR ODS-II	120	5	20.0	250	404-030

#### ■ Guard Column for Analytical HPLC

Column name	Pore dia. (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	GLC Cat. No.
STR ODS-II	120	5	4.0	10	404-023
			4.6	10	404-024
			6.0	10	404-025
STR ODS-II PEEK	120	5	4.6	10	404-028

#### ■ Guard Column for Preparative HPLC

Column name	Pore dia. (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	GLC Cat. No.
STR ODS-II	120	5	20.0	50	404-031



## STR ODS Columns

### 1. Analytical HPLC

Column name	Pore dia. (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	Cat. No.
ODS-H	100	5	4	50	228-21336-02
				150	228-21336-01
				250	228-21336-03
			4.6	150	228-21336-04
				250	228-21336-05
ODS-M	100	5	4	150	228-21830-01
				250	228-21830-02
			4.6	150	228-21830-03
				250	228-21830-04

### 2. Preparative HPLC

Column name	Pore dia. (Å)	Particle dia. (μm)	Column i.d. (mm)	Column length (mm)	Cat. No.
PREP ODS-H	120	5	20	250	228-21831-01
				50	228-21831-02
			4.6	250	228-21831-03
PREP ODS-M	120	5	20	250	228-21832-01
				50	228-21832-02
			4.6	250	228-21832-03

### STR ODS-H

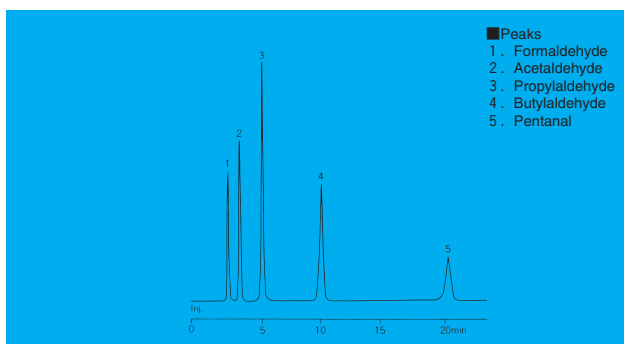


Fig. 117 Analysis of Aldehydes

#### Operational Conditions

**Column:** ODS-H (4.0 mm i.d. × 15 cm)  
**Mobile phase:** Methanol/water (40/60)  
**Column temperature:** 40°C  
**Flow rate:** 0.8 mL/min.  
**Detector:** Fluorophotometric detector (Ex: 360 nm, Em: 440nm)

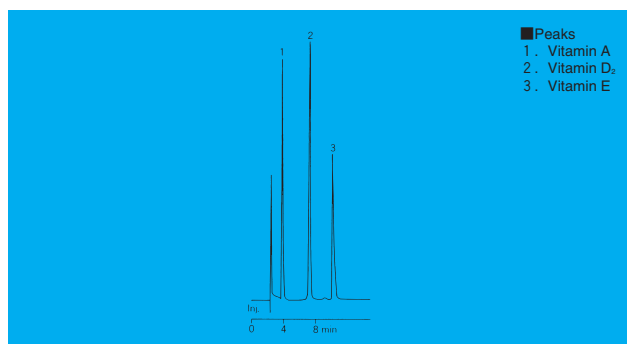


Fig. 118 Analysis of Oil-Soluble Vitamins

#### Operational Conditions

**Column:** ODS-H (4.0 mm i.d. × 15 cm)  
**Mobile phase:** Methanol  
**Column temperature:** 40°C  
**Flow rate:** 0.5 mL/min.  
**Detector:** UV spectrophotometric detector (260 nm)

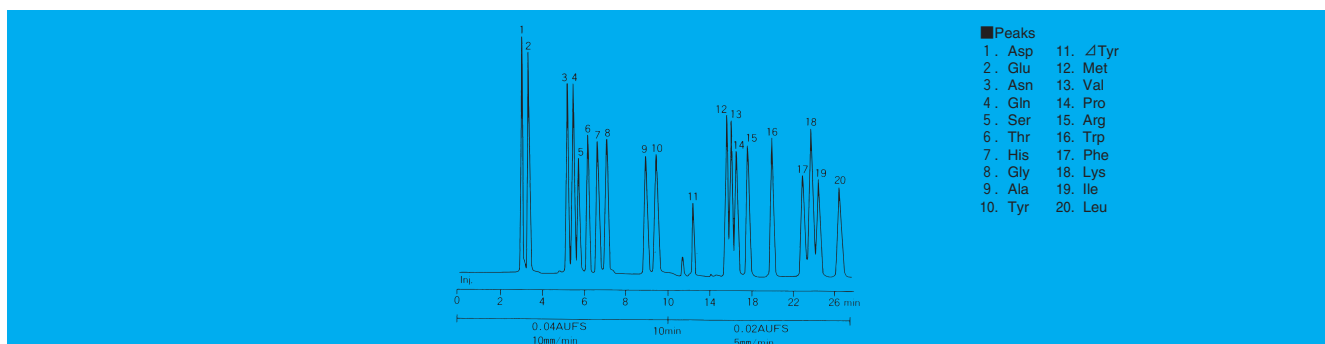


Fig. 119 Analysis of PHT-Amino Acids

#### Operational Conditions

**Column:** ODS-H (4.0 mm i.d. × 25 cm)  
**Mobile phase:** 10mM Sodium formate buffer solution (pH 5.50 and containing 0.070% SDS)/acetonitrile (60/40)  
**Column temperature:** 40°C  
**Flow rate:** 0.5 mL/min.  
**Detector:** UV spectrophotometric detector (269 nm)

## STR ODS-II

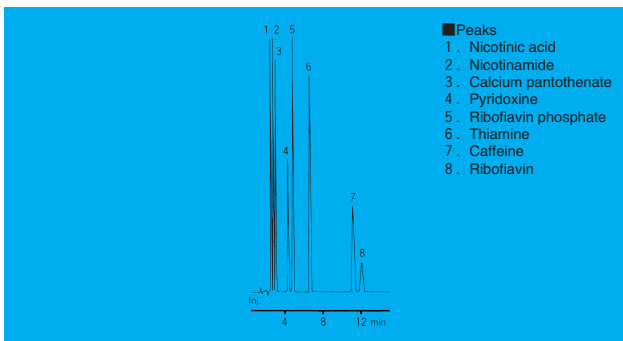


Fig. 120 Analysis of Water Soluble Vitamins

### Operational Conditions

**Column:** ODS-II (4.0 mm i.d.×15 cm)  
**Mobile phase:** 100mM phosphate buffer solution (pH 2.1) and 1.5mM Sodium octane sulfonate (9/1)  
**Column temperature:** 40°C  
**Flow rate:** 1.0 mL/min.  
**Detector:** UV spectrophotometric detector (210 nm)

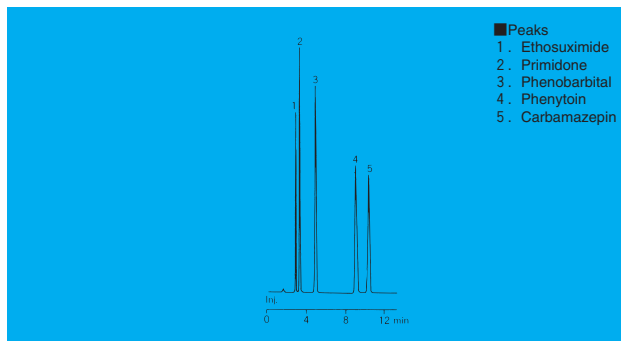


Fig. 121 Analysis of Anticonvulsant

### Operational Conditions

**Column:** ODS-II (4.0 mm i.d.×15 cm)  
**Mobile phase:** 100mM phosphate buffer solution (pH 2.1) /methanol/acetonitrile (4/2/1)  
**Column temperature:** 40°C  
**Flow rate:** 1.0 mL/min.  
**Detector:** UV spectrophotometric detector (210 nm)

## STR ODS-M

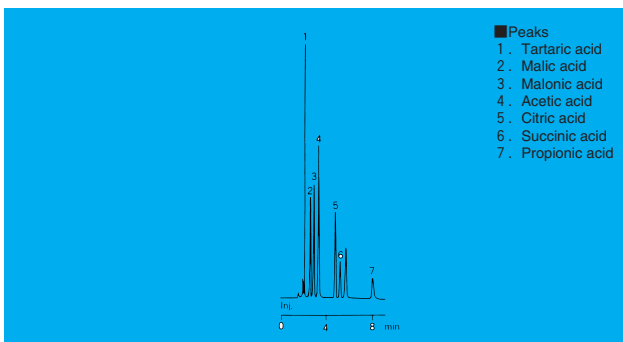


Fig. 122 Analysis of Organic Acids

### Operational Conditions

**Column:** ODS-M (4.6 mm i.d.×15 cm)  
**Mobile phase:** 10mM KH<sub>2</sub>PO<sub>4</sub> (adjusted to pH 2.3 with phosphoric acid)  
**Column temperature:** 40°C  
**Flow rate:** 1.0 mL/min.  
**Detector:** UV spectrophotometric detector (210 nm)

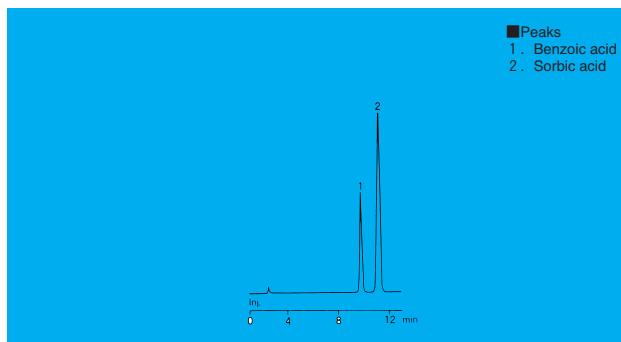


Fig. 123 Analysis of Food Preservatives

### Operational Conditions

**Column:** ODS-M (4.6 mm i.d.×15 cm)  
**Mobile phase:** 25mM Phosphate buffer solution (pH 3.5)/methanol (7/3)  
**Column temperature:** 50°C  
**Flow rate:** 0.8 mL/min.  
**Detector:** UV spectrophotometric detector (254 nm)

## STR PREP ODS-H

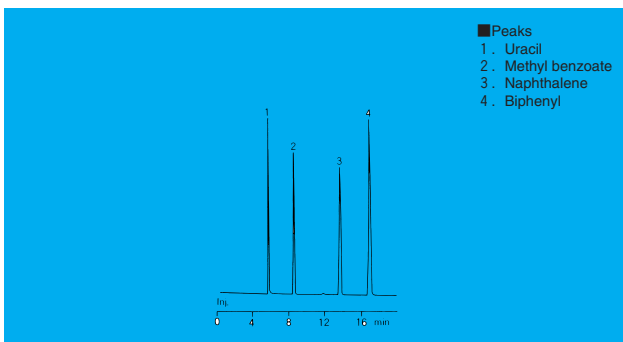


Fig. 124 Inspection Data

### Operational Conditions

**Column:** PREP ODS-H (20 mm i.d.×25 cm)  
**Mobile phase:** Methanol/water (85/15)  
**Column temperature:** Ambient  
**Flow rate:** 8.0 mL/min.  
**Detector:** 20 μL

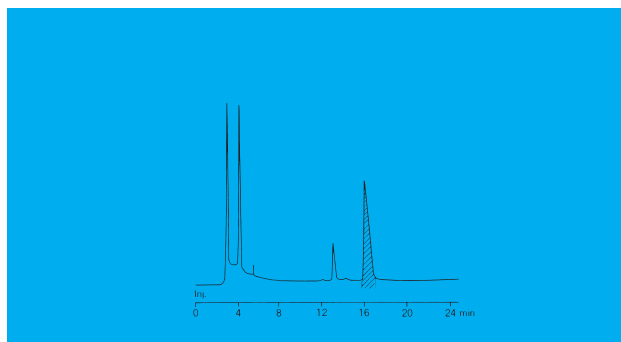


Fig. 125 Preparative LC Chromatogram of Paeoniflorin in Peony Extract

### Operational Conditions

**Column:** PREP ODS-H (20 mm i.d.×25 cm)  
**Mobile phase:** Water/acetonitrile (6/1)  
**Column temperature:** Ambient  
**Flow rate:** 15 mL/min.  
**Detector:** UV spectrophotometric detector (230 nm)  
**Sample size:** 100 mg of peony extract (2.0 mL of solution)

# 4 Application Data

## Food Industry

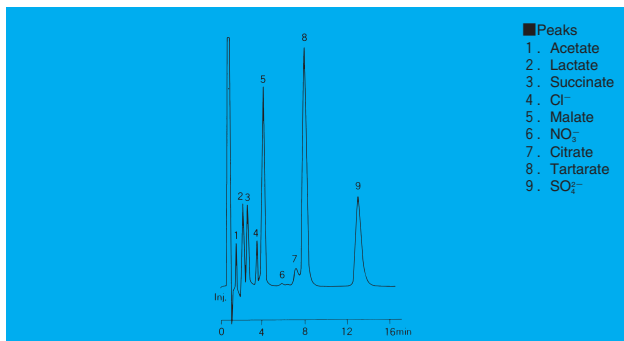


Fig. 126 Analysis of Anions in Wine

**Operational Conditions**  
**Column:** Shim-pack IC-A1 (4.6 mm i.d.×15 cm)  
**Mobile phase:** 1.2mM potassium biphthalate (pH 4.2)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector

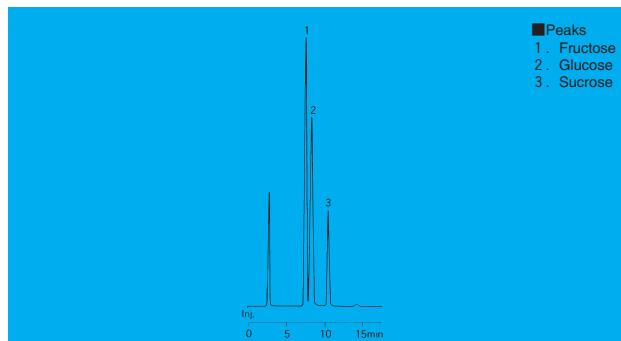


Fig. 127 Determination of Saccharides in Soft Drink

**Operational Conditions**  
**Column:** Shim-pack CLC-NH<sub>2</sub> (6.0 mm i.d.×15 cm)  
**Mobile phase:** Acetonitrile/water(7/3)  
**Flow rate:** 1.2 mL/min.  
**Column temperature:** Ambient  
**Detector:** Refractive index detector

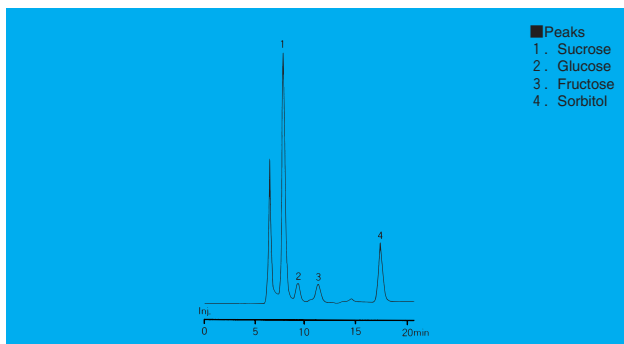


Fig. 128 Determination of Saccharides in Pickles

**Operational Conditions**  
**Column:** Shim-pack SCR-101C (7.9 mm i.d.×30 cm)  
**Mobile phase:** Water  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 85°C  
**Detector:** Refractive index detector

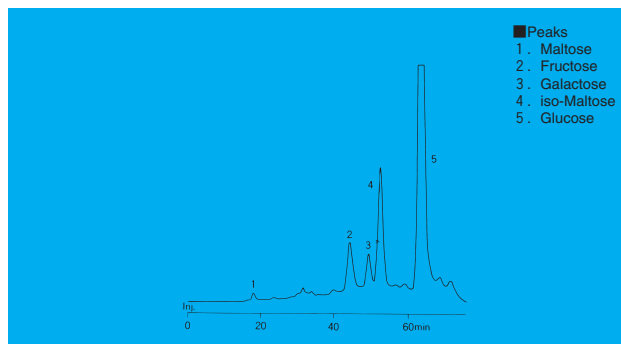


Fig. 129 Determination of Saccharides in Miso (Bean paste)

**Operational Conditions**  
**Column:** Shim-pack ISA-07 (4.0 mm i.d. × 25 cm)  
**Mobile phase:** Potassium borate buffer solution, gradient elution  
**Flow rate:** 0.6 mL/min.  
**Column temperature:** 65°C  
**Detector:** Fluorescence detector (Ex. 320nm, Em.430nm)  
**Method:** Post-column derivatization with arginine

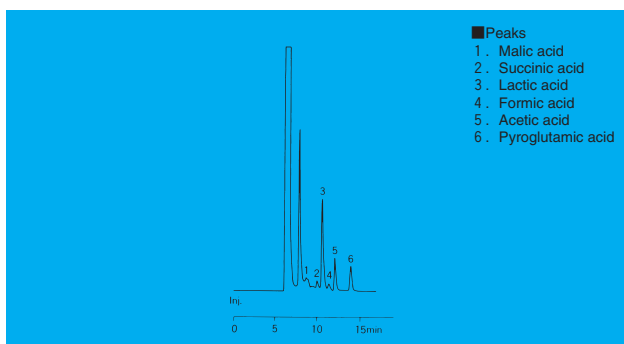


Fig. 130 Determination of Organic Acids in Soy Sauce

**Operational Conditions**  
**Column:** Shim-pack SCR-102H (8.0 mm i.d.×30 cm)  
**Mobile phase:** 5mM p-toluenesulfonic acid  
**Flow rate:** 0.8 mL/min.  
**Column temperature:** 45°C  
**Detector:** Conductivity detector  
**Method:** Post-column derivatization with bis-tris buffer

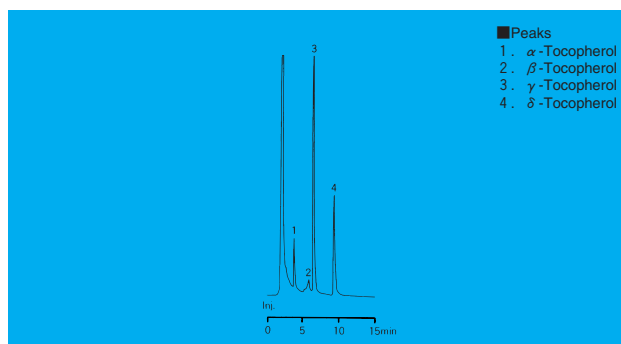


Fig. 131 Determination of Tocopherols in Soy Bean Oil

**Operational Conditions**  
**Column:** Shim-pack CLC-NH<sub>2</sub> (6.0mm i.d.×15 cm)  
**Mobile phase:** n-Hexane/isopropanol (25/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (297nm)

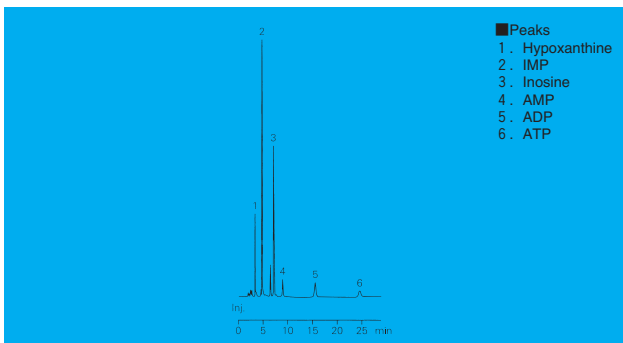


Fig. 132 Analysis of Adenine Derivative 6 Components

**Operational Conditions**  
**Column:** STR ODS-II (4.6 mm i.d.×15 cm)  
**Mobile phase:** 100mM phosphate (triethylammonium) buffer solution (pH 6.8) 100/acetonitrile  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (260nm)

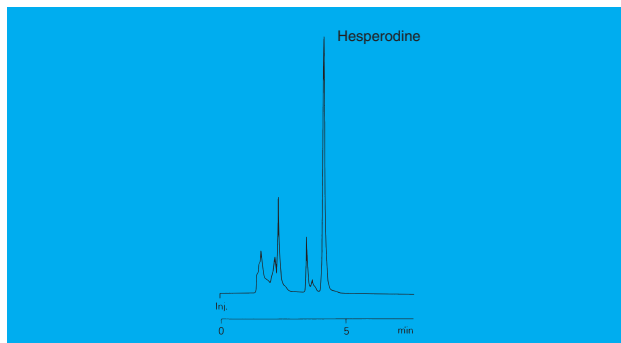


Fig. 133 Determination of Hesperidine in Orange Juice

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0 mm i.d.×15 cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1)/acetonitrile (3/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (265nm)

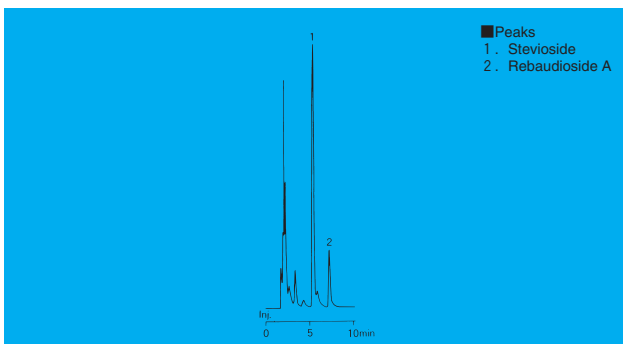


Fig. 134 Determination of Steviosides in Pickles

**Operational Conditions**  
**Column:** Shim-pack CLC-NH<sub>2</sub> (6.0 mm i.d.×15 cm)  
**Mobile phase:** Acetonitrile/water (3/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

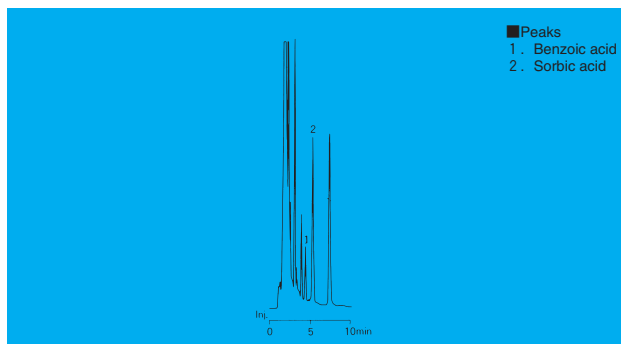


Fig. 135 Analysis of Preservatives in Food

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0 mm i.d.×15 cm)  
**Precolumn:** Shim-pack GRD-ODS  
**Mobile phase:** 10mM phosphate buffer solution (pH 6.9) /acetonitrile (20/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (225nm)

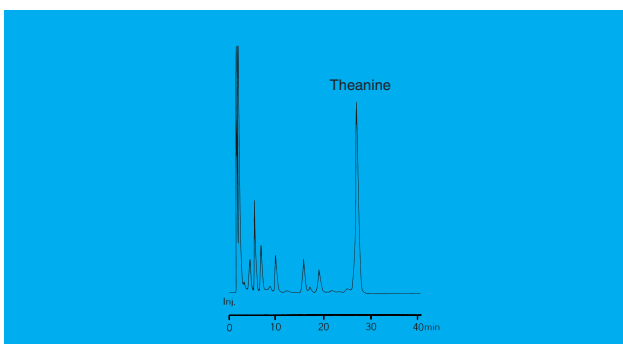


Fig. 136 Determination of Theanine in Tea Leaf

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0 mm i.d.×15 cm)  
**Precolumn:** Shim-pack GRD-ODS  
**Mobile phase:** 10mM phosphate buffer solution (pH 7.1)/acetonitrile (92/8)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** Ambient  
**Detector:** Fluorescence detector (Ex. 365nm, Em. 435nm)  
**Method:** Prelabelling with OPA

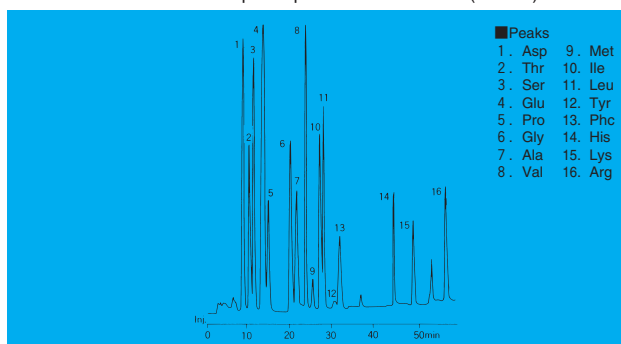


Fig. 137 Analysis of Amino Acids in Soy Sauce

**Operational Conditions**  
**Column:** Shim-pack ISC-07 (4.0 mm i.d.×15 cm)  
**Mobile phase:** Sodium citrate buffer solution, gradient elution  
**Flow rate:** 0.3 mL/min.  
**Detector:** Fluorescence detector (Ex. 348nm, Em. 450nm)  
**Method:** Prelabelling with OPA

## Medicines and Cosmetics

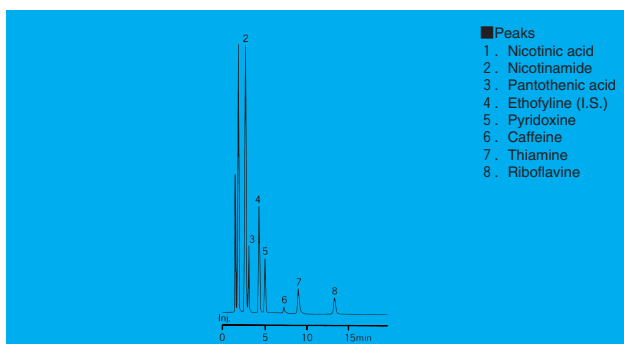


Fig. 138 Analysis of Vitamin Tablet

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0 mm i.d.×15 cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1) and 1.2mM sodium octane sulfonate/acetonitrile (9/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

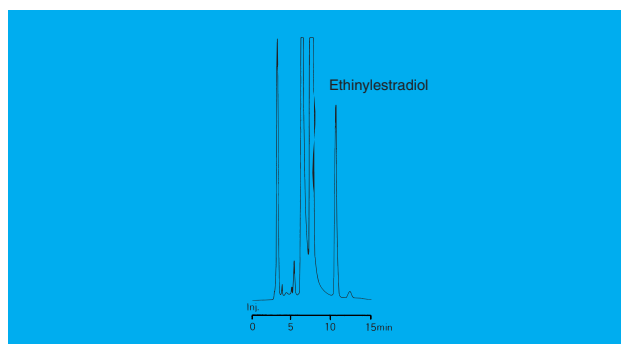


Fig. 139 Determination of Ethinylestradiol in Cream

**Operational Conditions**  
**Column:** Shim-pack CLC-SIL(M) (4.6mm i.d.×25cm)  
**Mobile phase:** n-Hexane/ethanol (20/1)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** Fluorescence detector (Ex. 285nm, Em. 315nm)

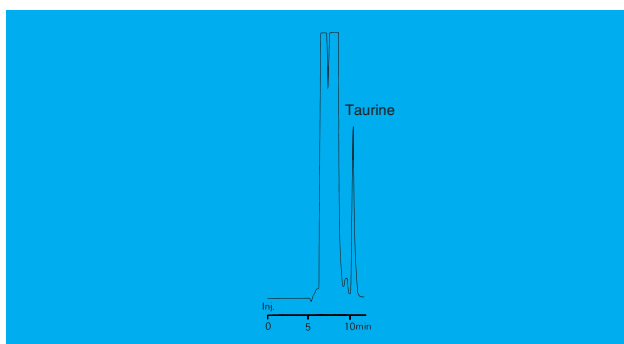


Fig. 140 Determination of Taurine in Soft Drink

**Operational Conditions**  
**Column:** Shim-pack SCR-101H (7.9mm i.d.×30cm)  
**Mobile phase:** 10mM aqueous solution of perchloric acid  
**Flow rate:** 0.8 mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

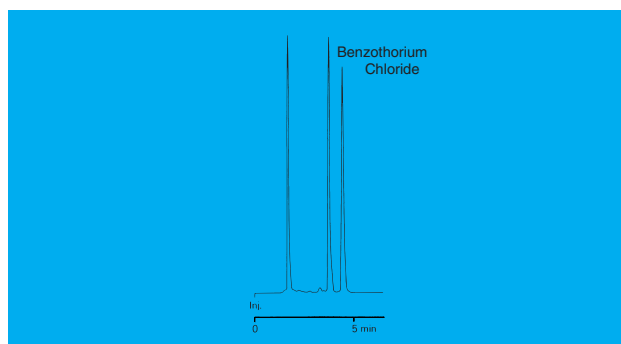


Fig. 141 Determination of Benzethonium Chloride in Gargle

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1)/acetonitrile (1/2)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (220nm)

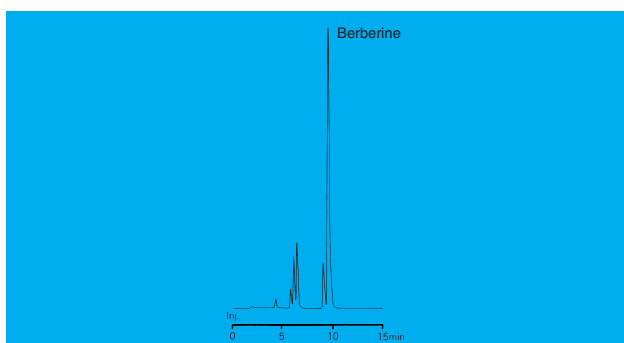


Fig. 142 Determination of Berberine in Coptis Japonica Powder

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) and 0.1M sodium perchlorate/acetonitrile (2/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (340nm)

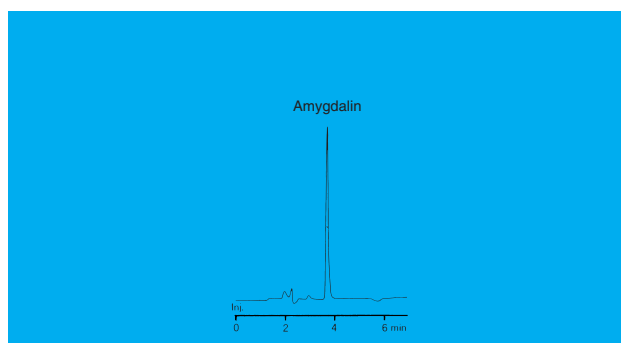


Fig. 143 Determination of Amygdalin in Apricot Seed

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Water/acetonitrile (4/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

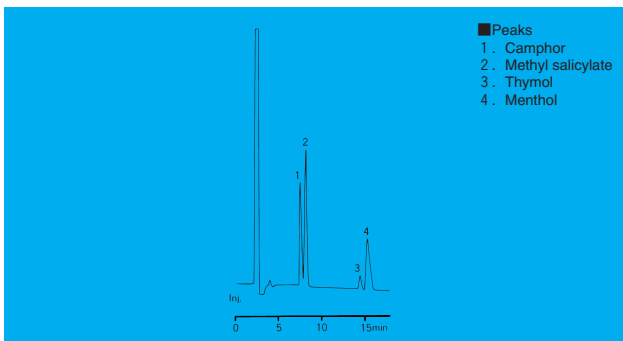


Fig. 144 Analysis of Antiphlogistic/Analgesic Paste

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Methanol/tetrahydrofuran/water (5/1/4)  
**Flow rate:** 1.2 mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

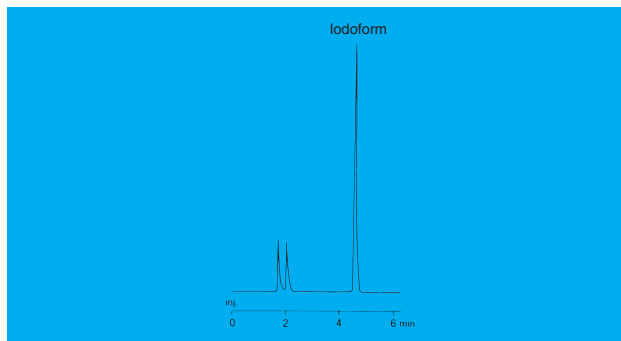


Fig. 145 Determination of Iodoform in Pulveres

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) /acetonitrile (1/2)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (335nm)

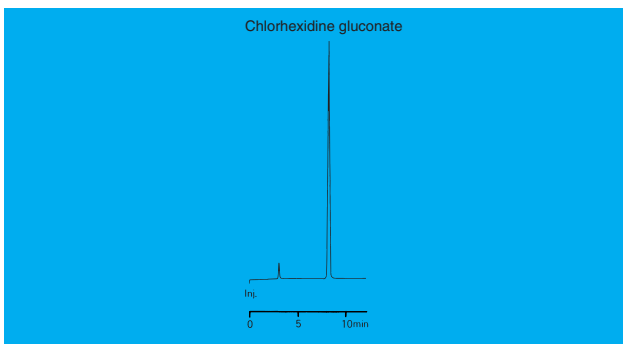


Fig. 146 Determination of Chlorhexidine Gluconate

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) and 0.1M sodium perchlorate/acetonitrile (3/2)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (240nm)

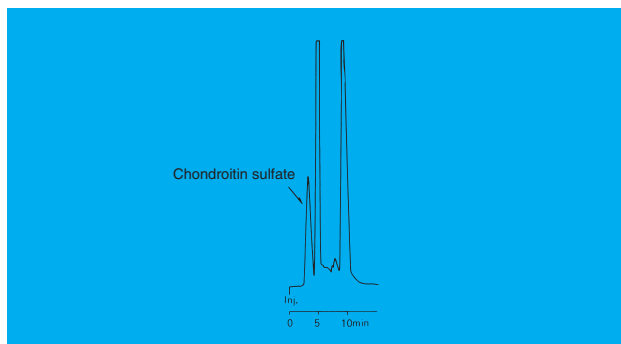


Fig. 147 Determination of Chondroitin sulfate in Eyewash

**Operational Conditions**  
**Column:** Shim-pack Diol-300 (7.9mm i.d.×25cm)  
**Mobile phase:** 50mM phosphate buffer solution (pH 3)  
**Flow rate:** 1.0 mL/min.  
**Detector:** UV spectrophotometric detector (210nm)

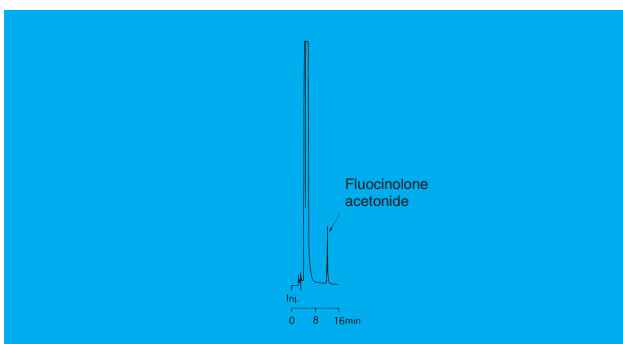


Fig. 148 Determination of Fluocinolone Acetonide in Cream

**Operational Conditions**  
**Column:** Zorbax CN (4.6mm i.d.×25cm)  
**Mobile phase:** n-Hexane/ethanol(4/1)  
**Flow rate:** 1.0mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector

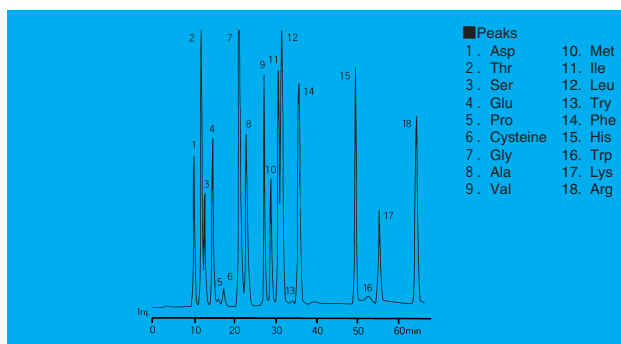


Fig. 149 Analysis of Amino Acid in Transfusion Liquid

**Operational Conditions**  
**Column:** Shim-pack ISC-07 (4.0mm i.d.×15cm)  
**Mobile phase:** Citric acid buffer solution, gradient elution  
**Flow rate:** 0.3 mL/min.  
**Column temperature:** 55°C  
**Detector:** Fluorescence detector (Ex. 348nm, Em. 450nm)  
**Method:** Post-column derivatization with OPA

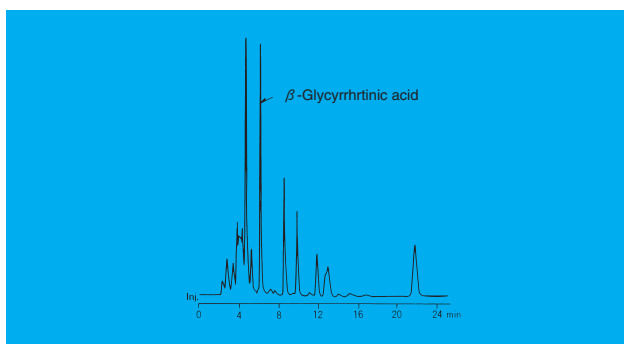


Fig. 150 Determination of  $\beta$ -Glycyrrhizic Acid in Cream

■ Operational Conditions

**Column:** CAPCELL PAK C<sub>18</sub> (5  $\mu$ m) (4.6mm i.d.  $\times$  25cm)  
**Mobile phase:** Acetonitrile/phosphoric acid (pH 2.5) (3/1)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (230nm)

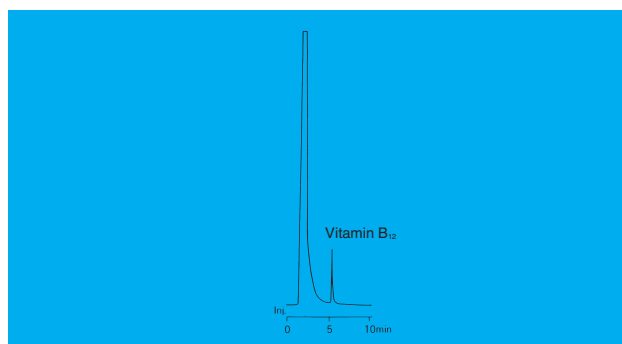


Fig. 151 Determination of Vitamin B<sub>12</sub> in Injection

■ Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.  $\times$  15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) /acetonitrile (7/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (278nm)

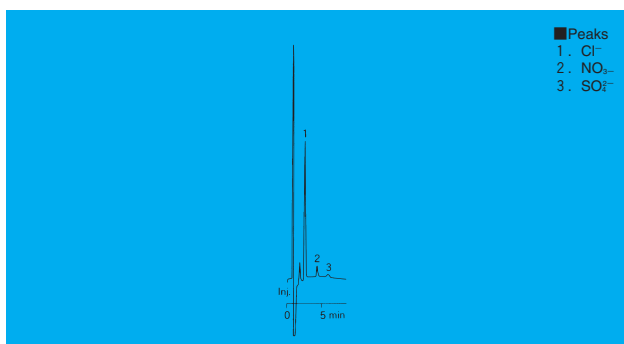


Fig. 152 Analysis of Inorganic Anions in Injection

■ Operational Conditions

**Column:** Shim-pack IC-A1 (4.6mm i.d.  $\times$  15cm)  
**Mobile phase:** 2.5mM phthalic acid and 2.4mM (tris) (hydroxymethyl) aminomethane (pH 4.0)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector

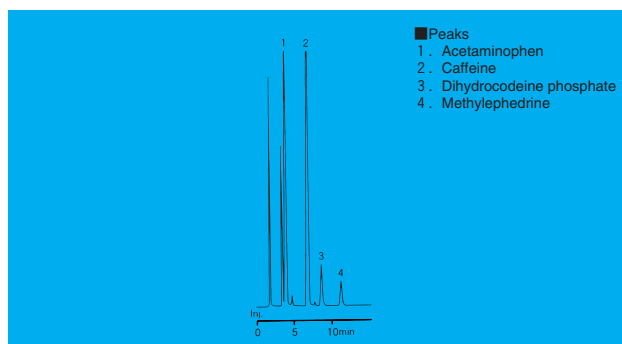


Fig. 153 Analysis of Cold Medicine

■ Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.  $\times$  15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) and 20mM sodium perchloric/acetonitrile (8/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

## Medical and Biochemical Field

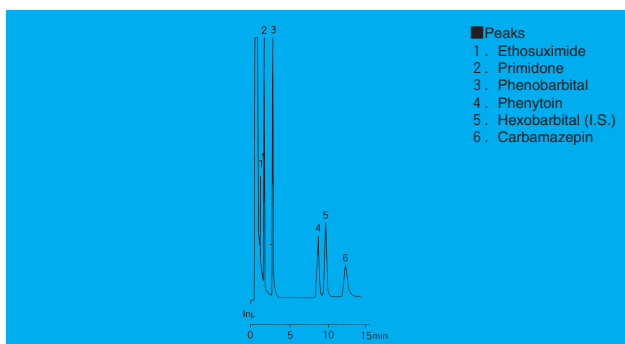


Fig. 154 Analysis of Anticonvulsants in Serum

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 5.5) /methanol(10/9)  
**Flow rate:** 1.2 mL/min.  
**Column temperature:** 55°C  
**Detector:** UV spectrophotometric detector (210nm)

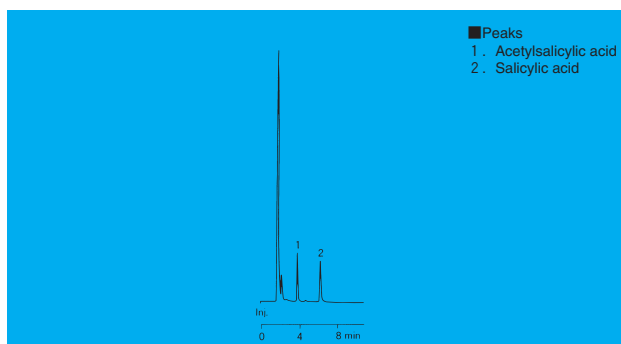


Fig. 155 Determination of Acetylsalicylic Acid and Salicylic Acid in Serum

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1)/methanol (1/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 55°C  
**Detector:** UV spectrophotometric detector (245nm)

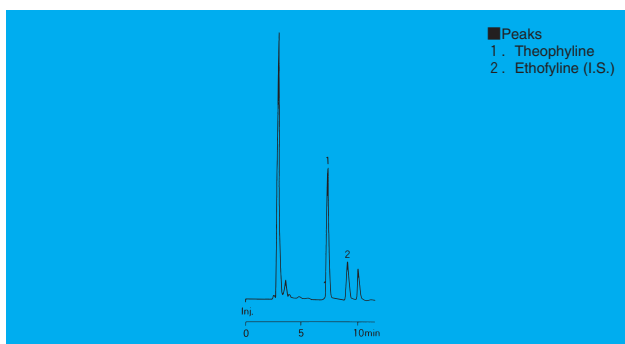


Fig. 156 Determination of Theophylline in Blood

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1)/acetonitrile (10/1)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (270nm)

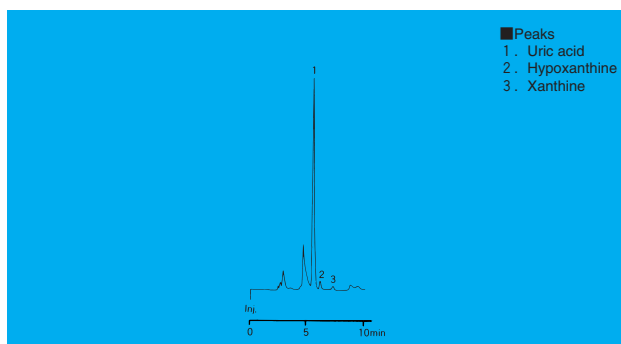


Fig. 157 Determination of Xanthine, Hypoxanthine, and Uric Acid in Urine

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 20mM phosphate buffer solution (pH 3)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (260nm)

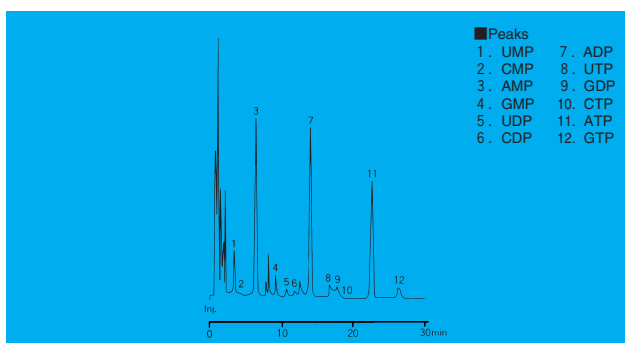


Fig. 158 Analysis of Nucleotides in Rat Liver Oil

**Operational Conditions**  
**Column:** Shim-pack WAX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** Phosphate buffer solution (pH 7), gradient elution  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 45°C  
**Detector:** UV spectrophotometric detector (260nm)

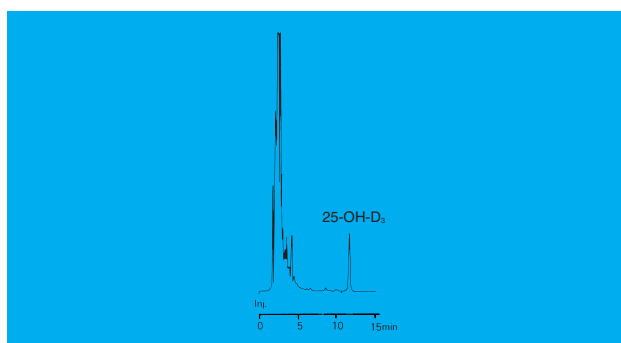


Fig. 159 Determination of Vitamin 25-OH-D<sub>3</sub> in Blood

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Methanol/water (5/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (265nm)



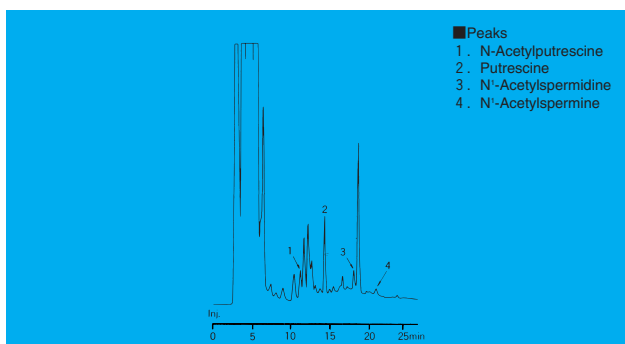


Fig. 160 Determination of Polyamines in Urine

■ Operational Conditions

**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Sodium perchlorate and sodium hexane sulfonate /acetonitril, gradient elution  
**Flow rate:** 1.1 mL/min.  
**Column temperature:** 50°C  
**Detector:** Fluorescence detector (Ex. 345nm, Em. 455nm)  
**Method:** Post-column derivatization with OPA

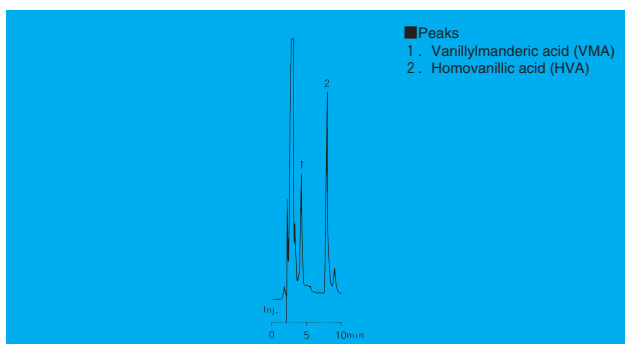


Fig. 161 Determination of HVA and VMA in Neonate Urine

■ Operational Conditions

**Column:** Shim-pack CLC-VMA (6.0mm i.d.×15cm)  
**Mobile phase:** Tartaric acid/acetonitrile (97/3) with a small amount of EDTA  
**Flow rate:** 1.5 mL/min.  
**Detector:** Conductivity detector

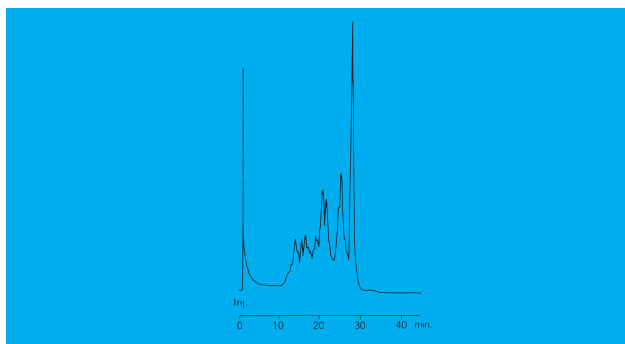


Fig. 162 Determination of  $\beta$ -Glucosidase

■ Operational Conditions

**Column:** Shim-pack HPC-C3 (4.0mm i.d.×5cm)  
**Mobile phase:** Ammonium sulfate/phosphate buffer solution (pH 7), gradient elution  
**Flow rate:** 0.5 mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (220nm)

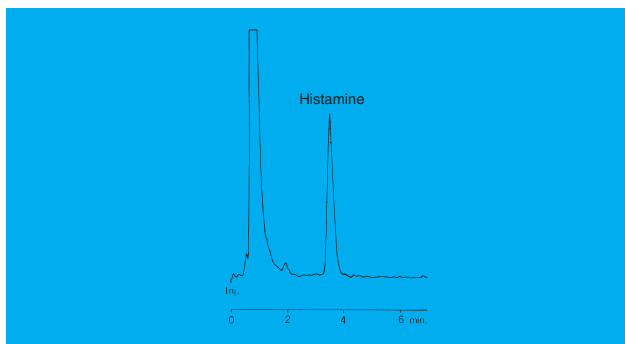


Fig. 163 Determination of Histamine in Plasma

■ Operational Conditions

**Column:** Shim-pack WCX-1 (4.0mm i.d.×5cm)  
**Mobile phase:** 40mM phosphate buffer solution (pH 6.9)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 50°C  
**Detector:** Fluorescence detector (Ex. 348nm, Em. 440nm)  
**Method:** Post-column derivatization with OPA

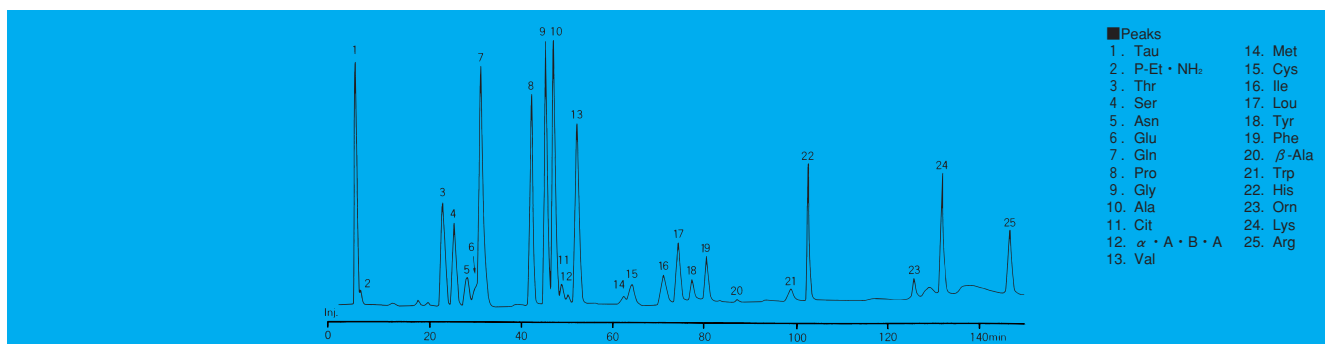


Fig. 164 Analysis of Amino Acids in Serum

■ Operational Conditions

**Column:** Shim-pack ISC-07 (4.0mm i.d.×15cm)  
**Mobile phase:** Lithium citrate buffer solution, gradient elution  
**Flow rate:** 0.4 mL/min.  
**Column temperature:** 38°C~58°C  
**Detector:** Fluorescence detector (Ex. 348nm, Em. 450nm)  
**Method:** Post-column derivatization with OPA

## Chemical Industry

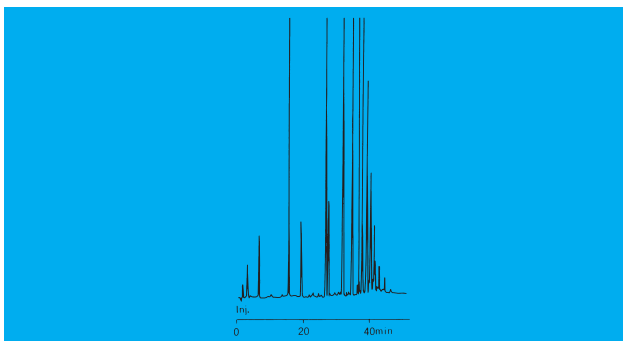


Fig. 165 Analysis of Epoxy Resin

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Water/acetonitrile, gradient elution  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (220nm)

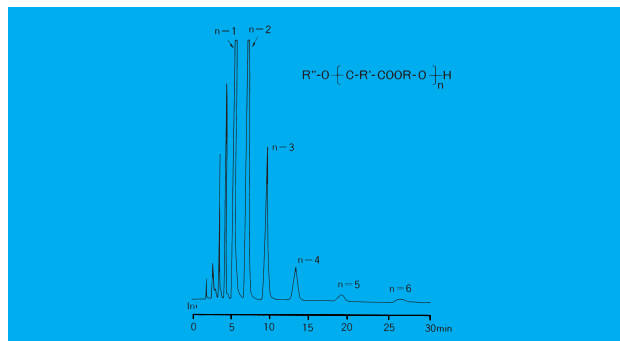


Fig. 166 Analysis of Ester Oligomers

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Methanol/water (4/1)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 50°C  
**Detector:** UV spectrophotometric detector (254nm)

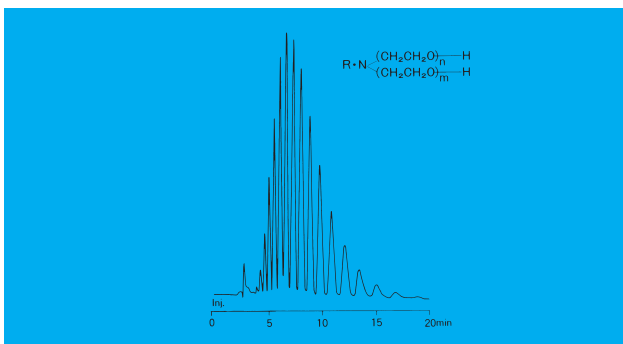


Fig. 167 Analysis of Alkylamine-Polyethylene Aducts

**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** Methanol/0.05% phosphoric acid (3/2)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

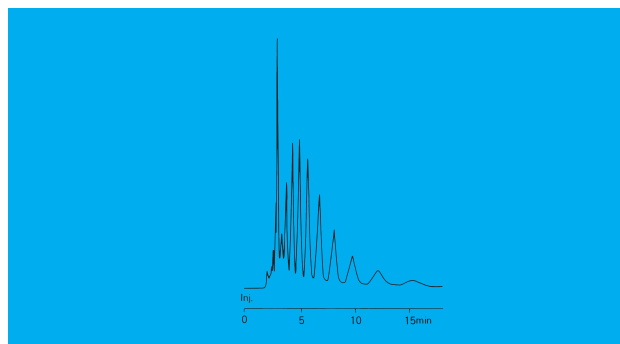


Fig. 168 Analysis of Polyoxyethylene Octylphenyl Ether

**Operational Conditions**  
**Column:** Shim-pack CLC-SIL (6.0mm i.d.×15cm)  
**Mobile phase:** n-Hexane/ethanol (2/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** Ambient  
**Detector:** UV spectrophotometric detector (250nm)

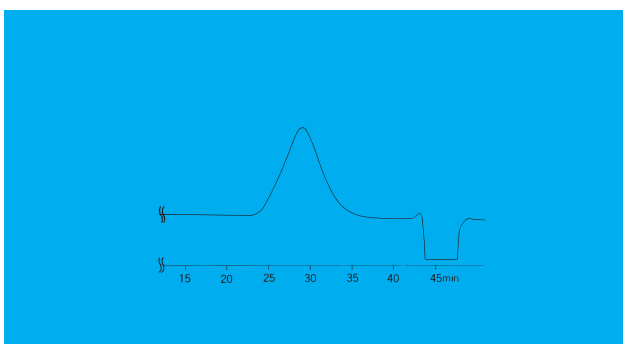


Fig. 169 Analysis of Polyvinylacetate

**Operational Conditions**  
**Column:** Shim-pack GPC-804, 803, and 802, (8.0mm i.d.×30cm, each)  
**Mobile phase:** Tetrahydrofuran  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

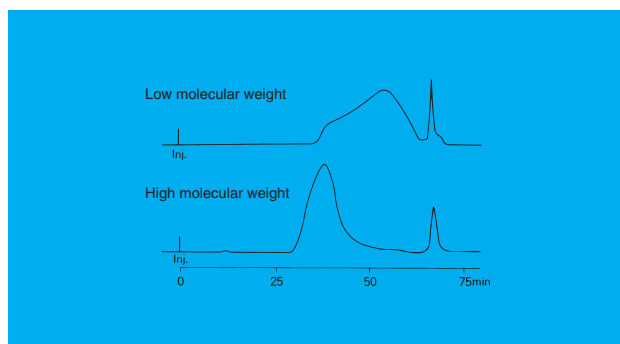


Fig. 170 Analysis of Carboxymethylcellulose

**Operational Conditions**  
**Column:** Asahipak GS-710 (two columns)  
**Mobile phase:** 50mM sodium nitrate  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 50°C  
**Detector:** Refractive index detector

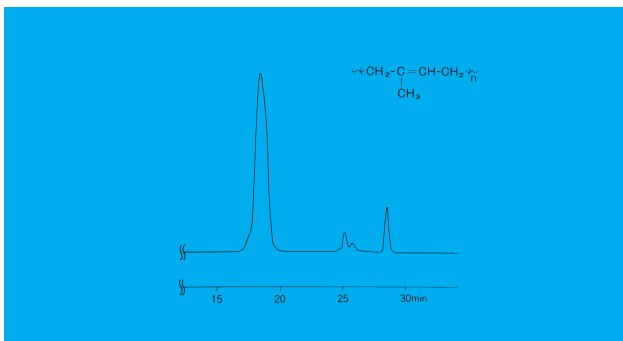


Fig. 171 Analysis of Isoprene Rubber

■Operational Conditions  
**Column:** Shim-pack GPC-804, 802, and 802 (8.0mm i.d.×30cm, each)  
**Mobile phase:** Tetrahydrofuran  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** Refractive index detector

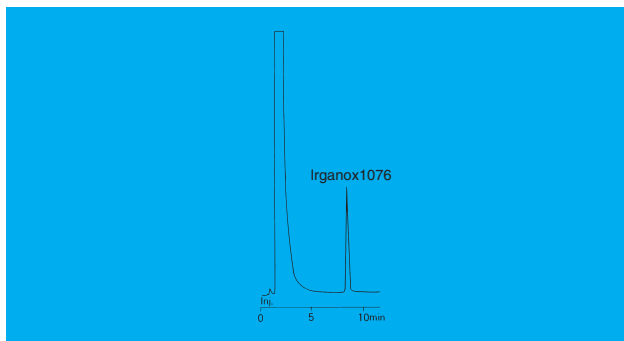


Fig. 172 Determination of Irganox 1076 in Rubber

■Operational Conditions  
**Column:** Zorbax ODS (4.6mm i.d.×15cm)  
**Mobile phase:** Methanol/tetrahydrofuran (9/1)  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (280nm)

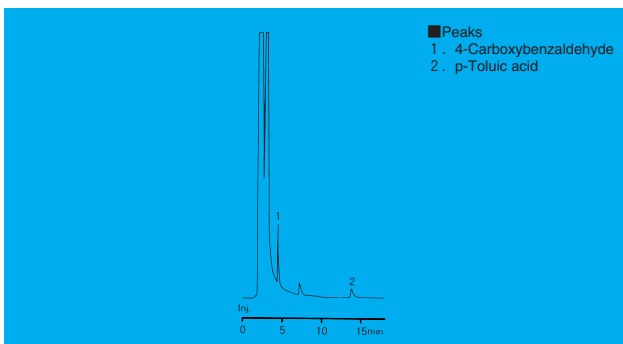


Fig. 173 Determination of Impurities in Terephthalic acid

■Operational Conditions  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 2.6) /acetonitrile (4/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 65°C  
**Detector:** UV spectrophotometric detector (230nm)

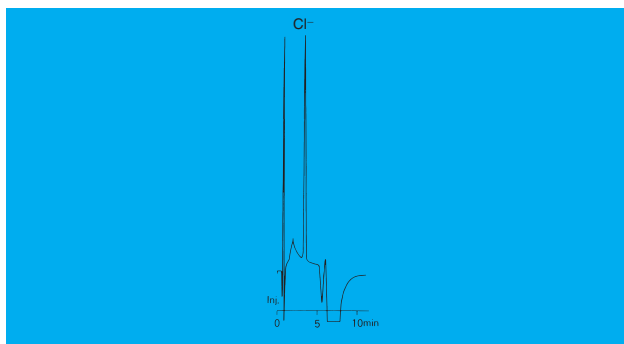


Fig. 174 Determination of Chlorine in Concrete Additive

■Operational Conditions  
**Column:** Shim-pack IC-A1 (4.6mm i.d.×10cm)  
**Mobile phase:** 2.0mM phthalic acid and 1.5mM tris (hydroxymethyl) aminomethane  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C

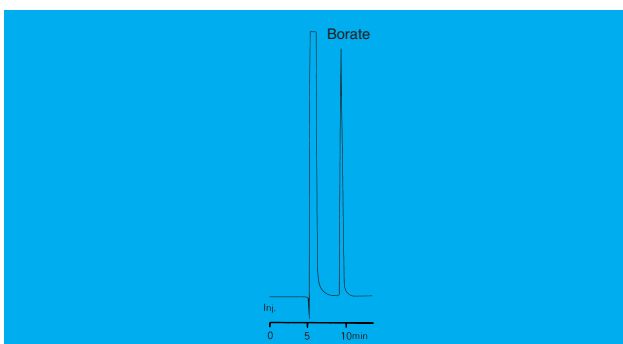


Fig. 175 Determination of Borate in Plating Solution

■Operational Conditions  
**Column:** Shim-pack SCR-101H (7.9mm i.d.×30cm)  
**Mobile phase:** 5mM perchloric acid aqueous solution  
**Flow rate:** 0.8 mL/min.  
**Column temperature:** 50°C  
**Detector:** Refractive index detector

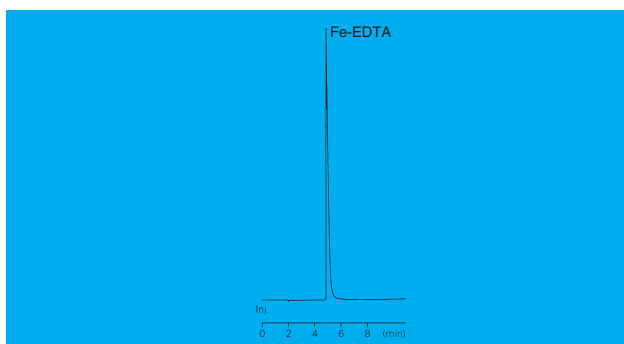


Fig. 176 Determination of EDTA

■Operational Conditions  
**Column:** STR ODS-II (4.6mm i.d.×15cm)  
**Mobile phase:** 15mM including tributylamines 10mM phosphate aqueous solution (pH 7) 5/acetonitrile 1  
**Flow rate:** 1.0 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (258nm)

## Environmental Pollutants

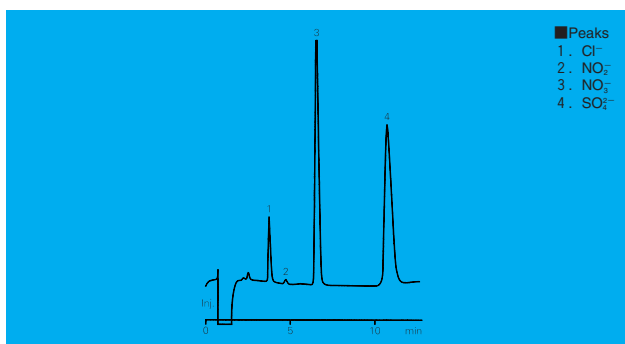


Fig. 177 Determination of Inorganic Anion in Rain Water

**Operational Conditions**  
**Column:** Shim-pack IC-A3  
**Mobile phase:** 8.0mM Hydroxyacetic benzoate  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector

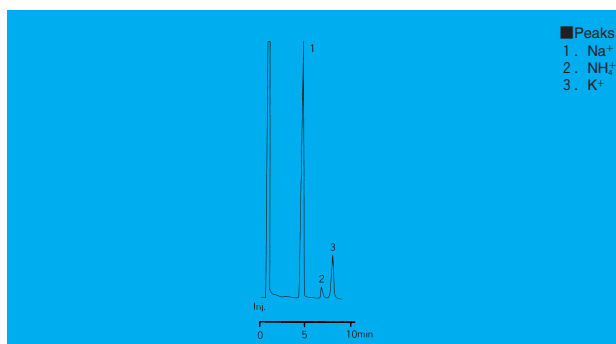


Fig. 178 Determination of Monovalent Cations in Soil Water

**Operational Conditions**  
**Column:** Shim-pack IC-C1 (5mm i.d.×15cm)  
**Mobile phase:** 5mM nitric acid aqueous solution  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** Conductivity detector

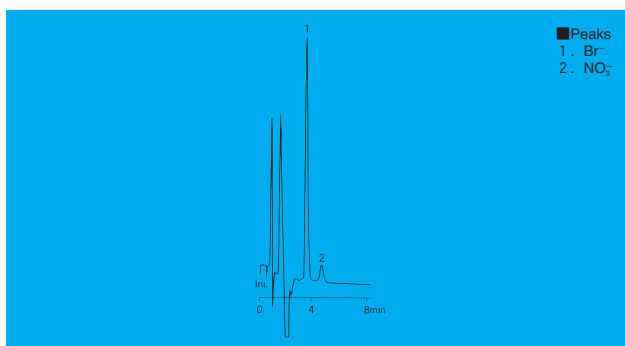


Fig. 179 Determination of Bromine and Nitric Acid

**Operational Conditions**  
**Column:** Shim-pack IC-A1 (4.6mm i.d.×15cm)  
**Mobile phase:** 10mM phosphate buffer solution (pH 6.8)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (210nm)

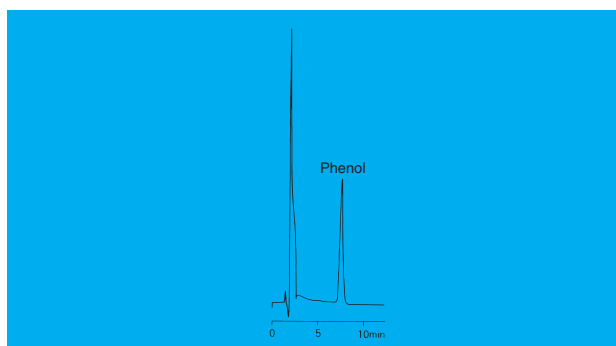


Fig. 180 Determination of Phenol in Water

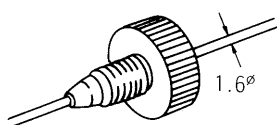
**Operational Conditions**  
**Column:** Shim-pack CLC-ODS (6.0mm i.d.×15cm)  
**Mobile phase:** 0.1M phosphate buffer solution (pH 2.1)/acetonitrile (3/1)  
**Flow rate:** 1.5 mL/min.  
**Column temperature:** 40°C  
**Detector:** UV spectrophotometric detector (265nm)  
**Sample pretreatment:** Concentrated with an automatic sample pretreatment system

# 5 Parts for Flow line

## Plumbing parts (Connecting Parts)

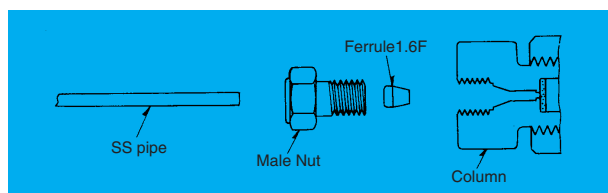
### Fingertightening connector

You can connect the column by tighening this connector, not by wrench. This PEEK connector which is superior in chemical compatibility and mechanical strength is suitable for biological LC. Available to 1.6mm i.d. pipe such as stainless, titanium, teflon, tefzel, PEEK etc. Maxium pressure is 250kgf/cm<sup>2</sup> when connecting to 0.5mm i.d. PEEK pipe.



Description	Cat. No.
Male Nut, PEEK (Inc. 5pcs.)	228-18565-84

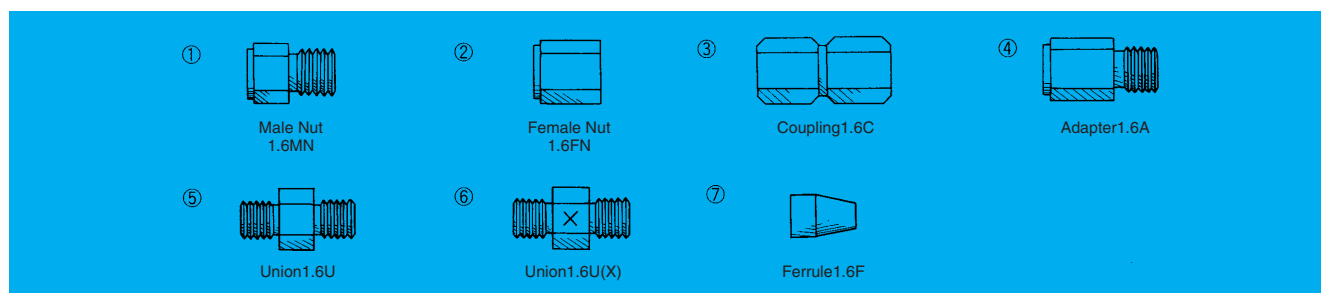
### ● Connecting by stainless plumbing parts



To connect this type of column, use the kit below.

Description	Cat. No.	Number
Column Connection Kit C ASSY (For new-new type connection)	228-16058-91	Pipe 0.3 i.d.×1.6 o.d.×25cm 1pc. Male Nut 1.6MN 2pcs. Ferrule 1.6F 2pcs.

### Connector Plumbing Parts

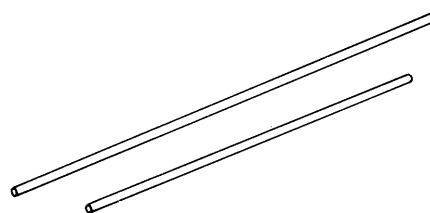


Description	Cat. No.	Number
① Male Nut 1.6MN	228-16001-84	10
② Female Nut 1.6FN	228-16002-84	10
③ Coupling 1.6C	228-16004-03	1
④ Adapter 1.6A	228-16005-03	1
⑤ Union 1.6U	228-00034	1
⑥ Union 1.6U(X)	228-01356	1
⑦ Ferrule 1.6F	228-16000-84	3
⑦ Ferrule 1.6F	228-16000-85	10

### Cut and sharpen not necessary

### Forged pipe

This 1.6mm o.d. pipe (SS 316) is cut at the size of easily use and mechanical-forged smoothly in both edges.



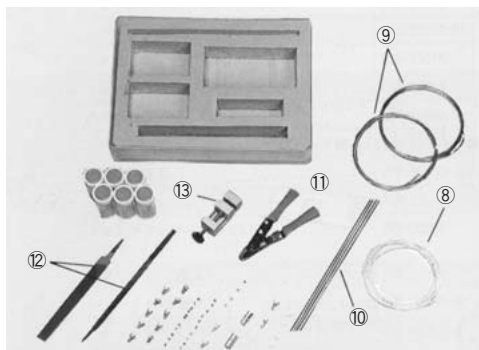
### Pre-cut Stainless Pipe

i.d.×o.d.×length (mm)	Cat. No
0.3×1.6×100	228-22301-00
0.3×1.6×200	228-22302-00
0.3×1.6×300	228-22303-00
0.3×1.6×400	228-22304-00
0.3×1.6×500	228-22305-00
0.8×1.6×100	228-22801-00
0.8×1.6×200	228-22802-00
0.8×1.6×300	228-22803-00
0.8×1.6×400	228-22804-00
0.8×1.6×500	228-22805-00

## LC plumbing part kit

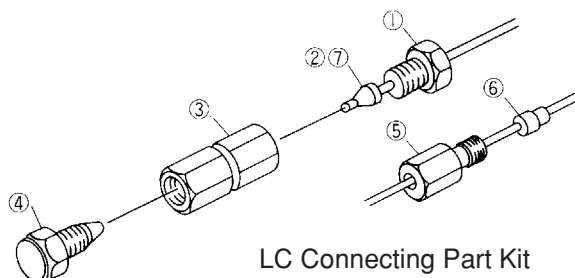
This kit put in a prastic case consists of pipes, connectors and tools needed to HPLC plumbing parts. It is possible to cut the pipe with the cutter attached as the length as you like. After cutting the pipe, hold this in the miniture vice and file the end of this smooth.

Description	Cat. No.
LC plumbing part kit	228-24209-91



[Kit]

Description	Cat. No.	Products Name	Number
Coupling 1.6C ASSY	228-16051-91	③ Coupling 1.6C	1
		⑦ Ferrule 1.6F	2
		① Male Nut 1.6MN	2
Adapter 1.6A ASSY	228-16052-91	④ Adapter 1.6A	1
		⑦ Ferrule 1.6F	2
		① Male Nut 1.6MN	1
		② Female Nut 1.6FN	1
Union 1.6U Newtype ASSY Type	228-16053-91	⑤ Union 1.6U	1
		⑦ Ferrule 1.6F	2
		② Female Nut 1.6FN	2
Union 1.6U(X) Newtype ASSY	228-16054-91	⑥ Union 1.6U(X)	1
		⑦ Ferrule 1.6F	2
		② Female Nut 1.6FN	2
Column connector kit	228-17943-92	① Male Nut 1.6MN	2
		⑦ Ferrule 1.6F	2



LC Connecting Part Kit

Mark	Description	Number	Notes
①	Male Nut, 1.6MN	10	
②	Ferrule, 1.6F	20	
③	Coupling, 1.6C-0.4	2	
④	Plug, 1.6P	2	
⑤	Longbush, 7010-011	2	} For Rheodyne injector
⑥	Ferrule, 7010-010	4	
⑦	PTFE ferrule, 1.6F-T	5	
⑧	PTFE tube, o.d. 1.6	2m	i.d. 0.4
⑨	SS pipe, o.d.1.6	Each 2m	i.d. 0.3 and 0.8
⑩	Pre-cut pipe, o.d. 1.6	4	i.d. 0.3, 250 length
⑪	Pipe cutter	1	
⑫	File (triangle, flat)	Each 1	
⑬	Miniture vice	1	

As if cutting a wire by wrench

### Cut the stainless pipe easily

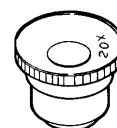
Holding and cutting the pipe like wrench by cutter in the LC plumbing part kit, we can prevent plugging the pipe. You can cut easily and the end of the pipe is smooth, comparing with the case of filing. (Do not forget to file the end of the pipe.) Possible to cut from 0.3mm i.d. to 1.6mm o.d.



Subscription	Cat. No.
Piping cutter	670-18802

### Checking the end of pipe

Use the pipe once filing the end of the stainless pipe smooth and checking with a magnifying glass carefully. We can look carefully with a High magnifying power glass made of non-colour lense.



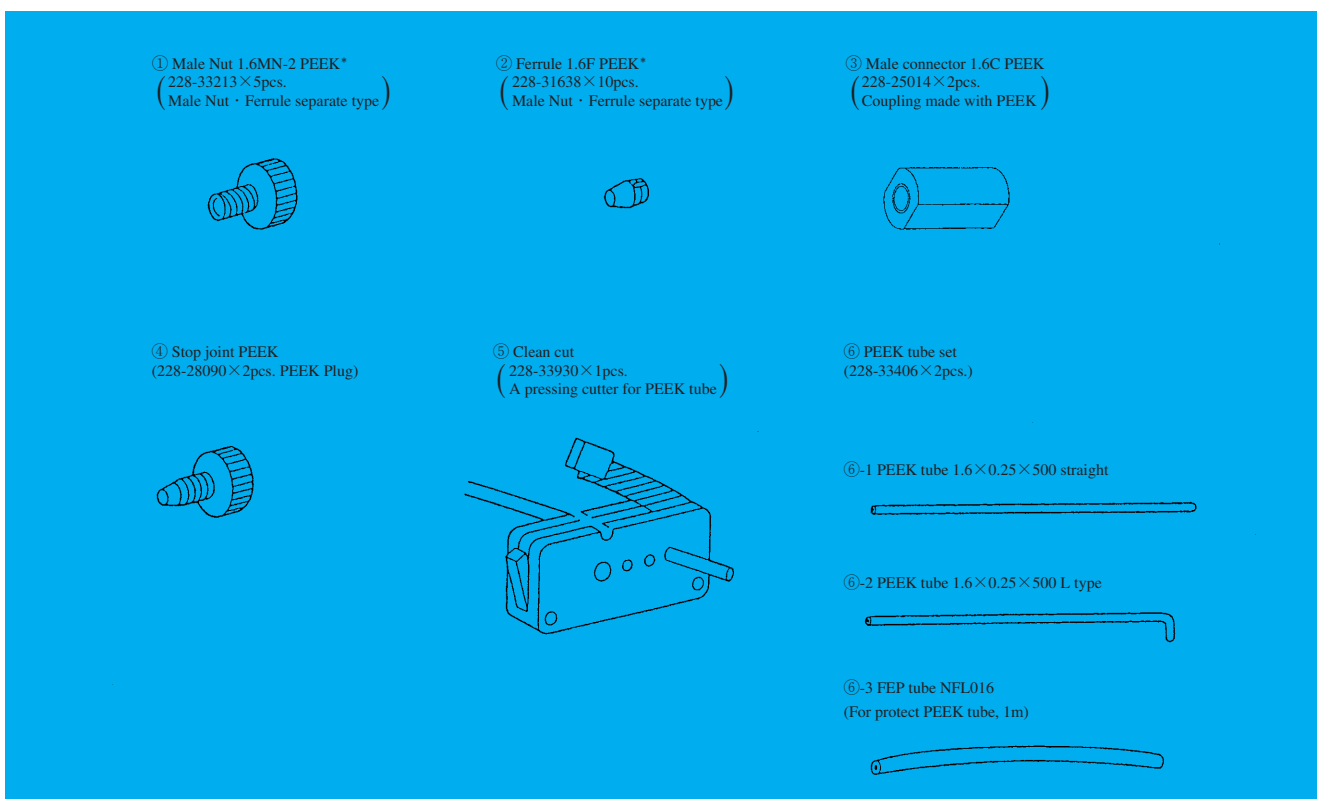
Description	Cat. No.
A magnifying glass, 20 times	670-12327

## PEEK plumbing parts

Use this parts for maintenance Non-metal LC system and system up.

Description	Cat. No.
Plumbing parts (Connecting parts) kit for inert LC	228-33285-91

### Component of Plumbing parts (Connecting parts) kit for Non-metal LC



\* There are two types: male nut · ferrule separation type and joint type. (Refer to P.83)  
 When using PEEK tube with organic solvent type mobile phase, use separation type.  
 When using PEEK tube with aqueous solution like IC, use joint type under normal pressure.

#### ● The solvent when using PEEK resin

Although PEEK resin has high mechanical strength and thus is considered as optimal plastic material for HPLC, there are some restrictions to use. Make sure that particularly the solvents below are NOT used.

Concentrated sulfuric acid, concentrated nitric acid, dichloroacetic acid, tetrahydrofuran (THF), dichloromethane, chloroform, dimethyl sulfoxide (DMSO), acetone

There is no problem of using methanol and acetonitriles, but use these solvents within the pressure below.

Methanol: Each diameter  $\sim 200 \text{ kgf/cm}^2$  (19.6MPa)

Acetonitriles: i.d.  $0.125 \phi / 0.25 \phi \rightarrow \sim 200 \text{ kgf/cm}^2$  (19.6MPa)

i.d.  $0.50 \phi \rightarrow \sim 150 \text{ kgf/cm}^2$  (14.7MPa)

i.d.  $0.75 \phi \rightarrow \sim 100 \text{ kgf/cm}^2$  (9.8MPa)

- Each PEEK tube showed in this page includes FEP tube for protection. Preparing for the case of PEEK tube is broken, be sure to put FEP tube for the security.

## PEEK plumbing parts (Not included in Plumbing Parts Kits)

Description	Cat. No.	Contents
Male Nut 1.6MN-2 PEEK	228-33213	Male nut • Ferrule separation type
Ferrule 1.6F PEEK (includes 3pcs.)	228-33513-91	Male nut • Ferrule separation type
Male connector 1.6C PEEK	228-25014	Coupling
3way joint 1.6C PEEK	228-25013	Female T joint* <sup>1</sup>
Stop joint PEEK	228-28090	Plug
Tubing cutter	228-32930-01	Press cutter for PEEK tube
Replacement blade	228-32930-02	
PEEK Tube set	228-33406-91	
Inline filter PEEK	228-32939	Filter for protecting a column (Put between injector and column)
Frit (1pcs.)	228-32744	Spare filter for inline filter
Frit (5pcs.)	228-33691-91	Spare filter for inline filter
Rheflex fitting set	228-32651-41	Inc. nut • ferrule each 5 (A spare for Rheodyne 9725/i)* <sup>2</sup>
PEEK tube (0.125 i.d.×1.6 o.d.×3000mm)	228-33833-91	
PEEK tube (0.25 i.d.×1.6 o.d.×3000mm)	228-33833-92	
PEEK tube (0.50 i.d.×1.6 o.d.×3000mm)	228-33833-93	
PEEK tube (0.75 i.d.×1.6 o.d.×3000mm)	228-33833-94	
Sample loop 100 μL	228-32651-16	Rheodyne injector for 9725/i
200 μL	228-32651-17	Rheodyne injector for 9725/i
500 μL	228-32651-18	Rheodyne injector for 9725/i
1mL	228-32651-19	Rheodyne injector for 9725/i

\* 1 Use this with chemical reaction detection system etc.

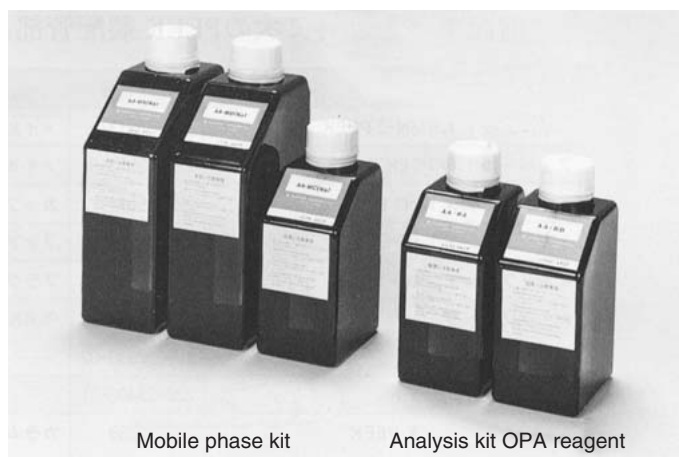
\* 2 Nut • Ferrule 5pcs. set for PEEK tube of Rheodyne 9725/i. Rheodyne 9725/i includes nut • ferrule 4pcs. so that this set is not necessary when installing.

"Rheflex" stands for Fex fitting made in Rheodyne.



## Shimadzu Amino Acid Analysis kit for amino acid analytical system

- As both mobile phase and reaction solutions are prepared to be optimized of analytical conditions, dilute and pH adjusting is not necessary.
- As strictly selected grade reagent is used, suitable for high sensitive analysis.



### Classification of analytical kit

Description	Cat. No.	Bottle name	Contents	Capacity	Usage
Amino acid mobile phase kit Na type	228-21195-94	AA-MA(Na)	0.2N-sodium citric acid	1L	A solution (No.1 buffer)
		AA-MB(Na)	0.6N-sodium citric acid	1L	B solution (No.2 buffer)
		AA-MC(Na)	0.2N-sodium hydroxide	500mL	C solution (column clean up solution)
Amino acid mobile phase kit A solution	228-21195-96	AA-MA(Na)	0.2N-sodium citric acid	1L×3pcs.	A solution (No.1 buffer)
Amino acid mobile phase kit Li type	228-21195-95	AA-MA(Li)	0.15N-lithium citric acid	1L	A solution (No.1 buffer)
		AA-MB(Li)	0.3N-lithium citric acid	1L	B solution (No.2 buffer)
		AA-MC(Li)	0.2N-lithium hydroxide	500mL	C solution (column clean up solution)
Amino acid mobile phase Li type A solution	228-21195-97	AA-MA(Li)	0.15N-lithium citric acid	1L×3pcs.	A solution (No.1 buffer)
Amino acid analytical kit OPA reagent	228-21195-93	AA-RA	Basic buffer	500mL	Reaction reagent A
		AA-RB	O-phthalic aldehyde	500mL	Reaction reagent B

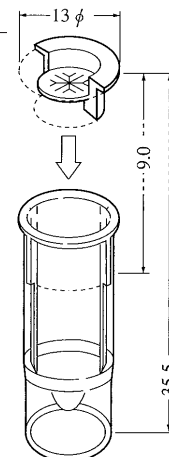
- Amino acid mobile phase kit Na type consists of mobile phase for amino acid analysis composed of protein (2 pcs.) and column clean up solution.
- Amino acid mobile phase kit Li type consists of amino acid mobile phase and column clean up solution.
- Analytical kit OPA reagent is a set of reaction reagent (2 kinds) and can be used in case of Na type and Li type.

### Disposable Vial

- Mass products made by plastic  
Suitable for the disposable of many samples
- Septa is attached in a cap  
Not necessary to set up septa
- Special patterned ditch on the septa  
The trouble such as the tears of the septa and the plugging of the syring hole doesn't occur.
- V bottom which solutions don't leave in  
The amount of the sample is available in the range of 50  $\mu$ l to 1ml. Also, the shape of the bottom is arranged not to leave air bubbles.

#### Specification

<b>Injector</b>	SIL-10A, SIL-9A, SIL-6B, SIL-10AXL
<b>Maxium amount</b>	1mL
<b>Minium amount of sample</b> (10 $\mu$ l inject)	50 $\mu$
<b>Materials</b>	Polyethylene (Cap, Septa) Polypropylene (Case)
<b>Cat. No.</b>	228-31600-91












Caution) When using disposable vial with a sample of organic solvent type, be sure to check whether there are disturbing peak or not to do blanktest.

## Disposable Filter

This disposable filter makes it possible to eliminate precipitate materials and minute particles in a sample. You can prevent the plugging of the column, the change of the property, and the trouble of the injector and the micro syringe. Therefore the column you use can be kept long.

### Features

- Small disk shape disposable filter.
- Rear lock attached and you can filtrate with easily operation and no trouble.
- The amount of the liquid left in this filter is quite small.
- Filtration can be with high flow and low pressure.
- There are many kinds of filter including ion chromatography as follows.

Series	Description	Shape	Diameter (mm)	Pore Size ( $\mu\text{m}$ )	Cat. No.	Features
Aqueous type	Filter, 4A		4	0.45	670-12540-01	<ul style="list-style-type: none"> <li>● Composed of a hydrophilic membrane and a polypropylene housing (blue).</li> <li>● Suitable for the pretreatment of the sample for HPLC analysis and protein analysis which are an object of aqueous and alcohol solution.</li> <li>● Sterilization, eliminated particles.</li> </ul>
	Filter, 13A		13	0.45	670-12540-02	
	Filter, 25A		25	0.45	670-12540-03	
Ion chromatography type	Filter, 4AI		4	0.45	670-12540-11	<ul style="list-style-type: none"> <li>● Composed of a de-ionized hydrophilic membrane and a polypropylene housing (light blue)</li> <li>● Suitable for the pretreatment of the sample for ion chromatography analysis.</li> <li>● As divided into each 10pcs, clean can be kept.</li> </ul>
	Filter, 13AI		13	0.45	670-12540-12	
	Filter, 25AI		25	0.45	670-12540-13	
Organic type	Filter, 4N		4	0.45	670-12540-21	<ul style="list-style-type: none"> <li>● Composed of PTFE membrane and a polypropylene housing. (half-transparent)</li> <li>● Superiority is solvent and chemical compatibility</li> <li>● Suitable for the pretreatment of semi-analytical sample which are an object of solvent, strong acid and strong alkaline.</li> <li>● It is good for air vent filter.</li> </ul>
	Filter, 13N		13	0.45	670-12540-22	
	Filter, 25N		25	0.45	670-12540-23	

(Inc. each 100pcs.)

## Chemical Compatibility

○ Can use △ Can use for short time × Can NOT use

	Chemical name	Aqueous & Ion chromatography type	Organic type
Acid	Glacial acetic acid	△	△
	Acetic acid 90%	△	△
	//    30%	○	○
	//    10%	○	○
	Hydrochloric acid Conc	×	×
	//    6N	×	△
	Sulfuric acid Conc	×	×
//    6N	×	△	
Nitric acid Conc	×	×	
//    6N	×	△	
Alkaline	Ammonia water 6N	△	△
	//    3N	○	○
	Calcium hydroxide 3N	○	○
	Sodium hydroxide 5N	△	○
//    3N	○	○	
Alcohols	Methanol	○	○
	Ethanol	○	○
	Propanol	○	○
	Isopropanol	○	○
	Butanol	○	○
	Amyl alcohol	○	○
	Ethylene glycol	○	○
	Propylene glycol	○	○
	Glycerine	○	○
Ether	Ethyl ether	○	○
	Isopropyl ether	○	○
	Dioxane	×	○
	Tetrahydrofuran	×	○
Esters	Methyl esters	×	○
	Ethyl esters	×	○
	Isopropyl esters	×	○
	Butyl esters	×	○
	Amyl esters	×	○
	Serosolve esters	×	○

	Chemical name	Aqueous & Ion chromatography type	Organic type
Ketones	Acetone	×	○
	Cyclohexane	×	○
	Methyl ethyl ketone	×	○
	Methyl isobutyl ketone	×	○
Hydrogen carbide	Benzene	×	○
	Toluene	×	○
	Xylenes	×	○
Halogenation hydrogen carbide	Ethane dichloride	×	○
	Methylene chloride	×	○
	Chloroform	×	○
	Carbon tetrachloride	×	○
	Perchloroethylene	×	○
	Trichloroethylene	×	○
	Freon TF	×	○
	Freon TMC	×	○
Oil group	Cotton-seed oil	○	○
	Lubrication oil	×	○
	Peanut oil	○	○
	Sesame oil	○	○
Others	Acetonitrile	×	○
	Aniline	×	○
	Gasoline	○	○
	Kerosene	○	○
	Dimethylsulfoxide	×	○
	Dimethylsulfoxide	×	○
	Terpene oil	○	○
	Pyridine	×	○
	Phenols (liquid)	×	○
	Hexane (dry)	○	○
	Formaldehyde 37%	○	○

## Specification

Item	Type	4mm			13mm			25mm		
		aqueous type	Ion chromatography type	Organic type	aqueous type	Ion chromatography type	Organic type	aqueous type	Ion chromatography type	Organic type
		4A	4AI	4N	13A	13AI	13N	25A	25AI	25N
Pore size	μm	0.45	0.45	0.45	0.45	0.45	0.45	0.45	0.45	0.45
Stuff*1	Filter	OP	OP	PTFE	OP	OP	PTFE	OP	OP	PTFE
	Housing	PP	PP	PP	PP	PP	PP	PP	PP	PP
Size (φ × L)	mm	8 × 18	8 × 18	8 × 18	18 × 22	18 × 22	18 × 22	29 × 24	29 × 24	29 × 24
Available filtration area	cm <sup>2</sup>	0.07	0.07	0.07	0.8	0.8	0.8	4	4	4
Maximum pressure	kg/cm <sup>2</sup>	7	7	7	5	5	5	6	6	6
Maximum temperature	°C	100	100	100	100	100	100	100	100	100
Amount of the liquid left	μL	Less than 10	Less than 10	Less than 10	Less than 30	Less than 30	Less than 30	Less than 100	Less than 100	Less than 100
Connecting parts	In	Ruar lock	Ruar lock	Ruar lock	Ruar lock	Ruar lock	Ruar lock	Ruar lock	Ruar lock	Ruar lock
	Out	Ruar slip	Ruar slip	Ruar slip	Ruar slip	Ruar slip	Ruar slip	Ruar slip	Ruar slip	Ruar slip
Sterilization method*2		E	E	A,E	E	E	A,E	E	E	A,E
Number(pcs./)case		100	100	100	100	100	100	100	100	100

\* 1 OP=Olefines type polymer, PP=Polypropylene, PTFE=Polytetrafluoroethylene

\* 2 S=AUTOclave, E=Ethylene oxide gas

## 6 Shim-pack Columns

Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack Amino	Shim-pack Amino-Na	6.0mm i.d.×10cm	228-18837-91	P.33
	Shim-pack Amino-Li	6.0mm i.d.×10cm	228-18837-91	
Shim-pack BIO (T)	Shim-pack CLC-ODS (T)	4.6mm i.d.×15cm	228-18062-91	P.57
	Shim-pack WAX-1 (T)	4.6mm i.d.×5cm	228-18257-91	
	Shim-pack WAX-2 (T)	4.6mm i.d.×5cm	228-18258-91	
	Shim-pack WCX-1 (T)	4.6mm i.d.×5cm	228-18259-91	
	Shim-pack HPC-C2 (T)	4.6mm i.d.×5cm	228-18260-91	
	Shim-pack HPC-C3 (T)	4.6mm i.d.×5cm	228-18261-91	
Shim-pack CLC	Shim-pack CLC-SIL	6.0mm i.d.×15cm	228-00807-91	P.24
	Shim-pack CLC-SIL (M)	4.6mm i.d.×15cm	228-17872-91	
	Shim-pack CLC-SIL (M)	4.6mm i.d.×25cm	228-17872-92	
	Shim-pack CLC-ODS	6.0mm i.d.×15cm	228-00808-91	
	Shim-pack CLC-ODS (M)	4.6mm i.d.×15cm	228-17873-91	
	Shim-pack CLC-ODS (M)	4.6mm i.d.×25cm	228-17873-92	
	Shim-pack CLC-C <sub>8</sub>	6.0mm i.d.×15cm	228-00809-91	
	Shim-pack CLC-C <sub>8</sub> (M)	4.6mm i.d.×15cm	228-17874-91	
	Shim-pack CLC-C <sub>8</sub> (M)	4.6mm i.d.×25cm	228-17874-92	
	Shim-pack CLC-TMS	6.0mm i.d.×15cm	228-00810-91	
	Shim-pack CLC-TMS (M)	4.6mm i.d.×15cm	228-17875-91	
	Shim-pack CLC-TMS (M)	4.6mm i.d.×25cm	228-17875-92	
	Shim-pack CLC-CN	6.0mm i.d.×15cm	228-00810-91	
	Shim-pack CLC-CN (M)	4.6mm i.d.×15cm	228-17876-91	
	Shim-pack CLC-CN (M)	4.6mm i.d.×25cm	228-17876-92	
	Shim-pack CLC-Phenyl	6.0mm i.d.×15cm	228-00812-91	
	Shim-pack CLC-Phenyl (M)	4.6mm i.d.×15cm	228-17877-91	
	Shim-pack CLC-Phenyl (M)	4.6mm i.d.×25cm	228-17877-92	
	Shim-pack CLC-NH <sub>2</sub>	6.0mm i.d.×15cm	228-16725-91	
	Shim-pack CLC-NH <sub>2</sub> (M)	4.6mm i.d.×15cm	228-17878-91	
Shim-pack CLC-NH <sub>2</sub> (M)	4.6mm i.d.×25cm	228-17878-92		
Shim-pack CLC-VMA	6.0mm i.d.×15cm	228-17255-91	P.51	
Shim-pack CLC-ODS/H	4.6mm i.d.×25cm	228-00808-92		

Column series	Column name	Dimension	Cat.No.	Reference		
Shim-pack Diol	Shim-pack Diol-150	7.9mm i.d.×25cm	228-14775-91	P.41		
	Shim-pack Diol-150	7.9mm i.d.×50cm	228-14775-92			
	Shim-pack Diol-300	7.9mm i.d.×25cm	228-14776-91			
	Shim-pack Diol-300	7.9mm i.d.×50cm	228-14776-92			
Shim-pack FLC	Shim-pack FLC-ODS	4.6mm i.d.×5cm	228-13375-91	P.30		
	Shim-pack FLC-SIL	4.6mm i.d.×5cm	228-13375-92			
	Shim-pack FLC-CN	4.6mm i.d.×5cm	228-13694-91			
	Shim-pack FLC-C <sub>8</sub>	4.6mm i.d.×5cm	228-13695-91			
	Shim-pack FLC-NH <sub>2</sub>	4.6mm i.d.×5cm	228-13696-91			
Shim-pack G	Shim-pack G-SIL (4)	4.0mm i.d.×1cm	228-18270-91	P.24, 29		
	Shim-pack G-SIL (8)	8.0mm i.d.×1.5cm	228-18270-92			
	Shim-pack GK-SIL	30mm i.d.×7.5cm	228-18338-91			
	Shim-pack GL-SIL	50mm i.d.×5cm	228-18339-91			
	Shim-pack G-ODS (4)	4.0mm i.d.×1cm	228-18246-91			
	Shim-pack G-ODS (8)	8.0mm i.d.×1.5cm	228-18246-92			
	Shim-pack GK-ODS	30mm i.d.×7.5cm	228-18321-91			
	Shim-pack GL-ODS	50mm i.d.×5cm	228-18322-91			
	Shim-pack G-C <sub>8</sub> (4)	4.0mm i.d.×1cm	228-18248-91			
	Shim-pack G-C <sub>8</sub> (8)	8.0mm i.d.×1.5cm	228-18248-92			
	Shim-pack G-TMS (4)	4.0mm i.d.×1cm	228-18262-91			
	Shim-pack G-TMS (8)	8.0mm i.d.×1.5cm	228-18262-92			
	Shim-pack G-CN (4)	4.0mm i.d.×1cm	228-18266-91			
	Shim-pack G-CN (8)	8.0mm i.d.×1.5cm	228-18266-92			
	Shim-pack G-Phenyl (4)	4.0mm i.d.×1cm	228-18264-91			
	Shim-pack G-Phenyl (8)	8.0mm i.d.×1.5cm	228-18264-92			
	Shim-pack G-NH <sub>2</sub> (4)	4.0mm i.d.×1cm	228-18268-91			
	Shim-pack G-NH <sub>2</sub> (8)	8.0mm i.d.×1.5cm	228-18268-92			
	Shim-pack GPRC	Shim-pack GPRC-SIL	8.0mm i.d.×1.5cm		228-23462-92	P.54
		Shim-pack GPRC-ODS	8.0mm i.d.×1.5cm		228-23465-92	
Shim-pack GPRC-C <sub>8</sub>		8.0mm i.d.×1.5cm	228-24386-92			
Shim-pack GPRC-TMS		8.0mm i.d.×1.5cm	228-24387-92			

Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack G	Shim-pack GPRC-NH <sub>2</sub>	8.0mm i.d.×1.5cm	228-24388-92	P.54
	Shim-pack GPRC-CN	8.0mm i.d.×1.5cm	228-24389-92	
	Shim-pack GPRC-SIL (K)	3.0mm i.d.×7.5cm	228-23462-93	
	Shim-pack GPRC-SIL (L)	50mm i.d.×5cm	228-23462-92	
	Shim-pack GPRC-ODS (K)	30mm i.d.×7.5cm	228-23465-93	
	Shim-pack GPRC-ODS (L)	50mm i.d.×5cm	228-23465-94	
Shim-pack GPC	Shim-pack GPC-801	8.0mm i.d.×30cm	228-20803-91	P.38
	Shim-pack GPC-802	8.0mm i.d.×30cm	228-20804-91	
	Shim-pack GPC-8025	8.0mm i.d.×30cm	228-20805-91	
	Shim-pack GPC-803	8.0mm i.d.×30cm	228-20806-91	
	Shim-pack GPC-804	8.0mm i.d.×30cm	228-20807-91	
	Shim-pack GPC-805	8.0mm i.d.×30cm	228-20808-91	
	Shim-pack GPC-806	8.0mm i.d.×30cm	228-20809-91	
	Shim-pack GPC-80M	8.0mm i.d.×30cm	228-20810-91	
	Shim-pack GPC-807	8.0mm i.d.×30cm	228-20811-91	
	Shim-pack GPC-800P	4.6mm i.d.×1cm	228-20812-91	
	Shim-pack GPC-801C	8.0mm i.d.×30cm	228-20803-92	
	Shim-pack GPC-802C	8.0mm i.d.×30cm	228-20804-92	
	Shim-pack GPC-8025C	8.0mm i.d.×30cm	228-20805-92	
	Shim-pack GPC-803C	8.0mm i.d.×30cm	228-20806-92	
	Shim-pack GPC-804C	8.0mm i.d.×30cm	228-20807-92	
	Shim-pack GPC-805C	8.0mm i.d.×30cm	228-20808-92	
	Shim-pack GPC-806C	8.0mm i.d.×30cm	228-20809-92	
	Shim-pack GPC-80MC	8.0mm i.d.×30cm	228-20810-92	
	Shim-pack GPC-807C	8.0mm i.d.×30cm	228-20811-92	
	Shim-pack GPC-800CP	4.6mm i.d.×1cm	228-20812-92	
	Shim-pack GPC-801D	8.0mm i.d.×30cm	228-20803-93	
Shim-pack GPC-802D	8.0mm i.d.×30cm	228-20804-93		
Shim-pack GPC-8025D	8.0mm i.d.×30cm	228-20805-93		
Shim-pack GPC-803D	8.0mm i.d.×30cm	228-20806-93		
Shim-pack GPC-804D	8.0mm i.d.×30cm	228-20807-93		

Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack GPC	Shim-pack GPC-805D	8.0mm i.d.×30cm	228-20808-93	P.38
	Shim-pack GPC-806D	8.0mm i.d.×30cm	228-20809-93	
	Shim-pack GPC-80MD	8.0mm i.d.×30cm	228-20810-93	
	Shim-pack GPC-807D	8.0mm i.d.×30cm	228-20811-93	
	Shim-pack GPC-800DP	4.6mm i.d.×1cm	228-20812-93	
	Shim-pack GPC-2001	20.0mm i.d.×30cm	228-23342-91	
	Shim-pack GPC-2002	20.0mm i.d.×30cm	228-23342-92	
	Shim-pack GPC-20025	20.0mm i.d.×30cm	228-23342-93	
	Shim-pack GPC-2003	20.0mm i.d.×30cm	228-23342-94	
	Shim-pack GPC-2000P	8.0mm i.d.×30cm	228-20812-94	
	Shim-pack GPC-2001C	20.0mm i.d.×30cm	228-23343-91	
	Shim-pack GPC-2002C	20.0mm i.d.×30cm	228-23343-92	
	Shim-pack GPC-20025C	20.0mm i.d.×30cm	228-23343-93	
	Shim-pack GPC-2003C	20mm i.d.×30cm	228-23343-94	
	Shim-pack GPC-2000CP	8.0mm i.d.×30cm	228-20812-95	
Shim-pack GRD	Shim-pack GRD-ODS	4.0mm i.d.×25cm	228-16557-91	P.50
Shim-pack HPC	Shim-pack HPC-C2	4.0mm i.d.×5cm	228-17792-91	P.47
	Shim-pack HPC-C3	4.0mm i.d.×5cm	228-17793-91	
Shim-pack HRC	Shim-pack HRC-SIL	4.6mm i.d.×15cm	228-23460-91	P.21
		4.6mm i.d.×25cm	228-23460-92	
		6.0mm i.d.×15cm	228-23460-93	
	Shim-pack HRC-ODS	4.6mm i.d.×15cm	228-24463-91	
		4.6mm i.d.×25cm	228-23463-92	
		6.0mm i.d.×15cm	228-23463-93	
	Shim-pack HRC-C <sub>8</sub>	4.6mm i.d.×15cm	228-24376-92	
		4.6mm i.d.×25cm	228-24376-92	
		6.0mm i.d.×15cm	228-24376-92	
	Shim-pack HRC-TMS	4.6mm i.d.×15cm	228-24337-91	
		4.6mm i.d.×25cm	228-24377-92	
		6.0mm i.d.×15cm	228-24377-93	
Shim-pack HRC-NH <sub>2</sub>	4.6mm i.d.×15cm	228-24378-91		

Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack HRC	Shim-pack HRC-NH <sub>2</sub>	4.6mm i.d.×25cm	228-24378-92	P.21
		6.0mm i.d.×15cm	228-24378-93	
	Shim-pack-HRC-CN	4.6mm i.d.×15cm	228-24379-91	
		4.6mm i.d.×25cm	228-24379-92	
		6.0mm i.d.×15cm	228-24379-93	
Shim-pack IC	Shim-pack IC-A1	4.6mm i.d.×10cm	228-17733-91	P.46
	Shim-pack IC-A2	5.0mm i.d.×5cm	228-17735-91	
	Shim-pack IC-C1	5.0mm i.d.×15cm	228-17737-91	
	Shim-pack IC-GA1	4.6mm i.d.×1cm	228-17734-91	
	Shim-pack IC-GA2	4.0mm i.d.×1cm	228-17736-91	
	Shim-pack IC-GC1	4.0mm i.d.×1cm	228-17738-91	
	Shim-pack IC-PCI	8.0mm i.d.×5cm	228-17744-91	
	Shim-pack IC-A1 (S)	2.0mm i.d.×15cm	228-33400-91	
	Shim-pack IC-A3 (S)	2.0mm i.d.×15cm	228-33366-91	
	Shim-pack IC-C3 (S)	2.0mm i.d.×10cm	228-33367-91	
	Shim-pack IC-CI PEEK	4.6mm i.d.×10cm	228-33497-91	
	Shim-pack IC-GC1 PEEK	4.6mm i.d.×9cm	228-33497-92	
Shim-pack ISA	Shim-pack ISA-07/S2504	4.0mm i.d.×25cm	228-09699-91	P.33
	Guard Column ISA	4.0mm i.d.×5cm	228-00823-91	
Shim-pack ISC	Shim-pack ISC-05/S0504	4.6mm i.d.×3.8cm	228-09700-91	P.33
	Shim-pack ISC-07/S1504	4.0mm i.d.×15cm	228-09328-91	
	Shim-pack ISC-07/S1504Li	4.0mm i.d.×15cm	228-00796-91	
	Shim-pack ISC-30/S0504	4.0mm i.d.×5cm	228-14206-91	
	Shim-pack ISC-30/S0504Li	4.0mm i.d.×5cm	228-00821-91	
	Guard Column ISC-07	4.0mm i.d.×5cm	228-00802-91	
	Guard Column ISC-07Li	4.0mm i.d.×5cm	228-00797-91	



Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack MBC	Shim-pack MBC-ODS	1.0mm i.d.×25cm	228-12812-02	P.29
	Shim-pack MBC-ODS	1.0mm i.d.×50cm	228-12812-05	
	Shim-pack MBC-SIL	1.0mm i.d.×50cm	228-12811-05	
	Shim-pack MBC-SIL	1.0mm i.d.×100cm	228-12811-10	
	Shim-pack MBC-ACN	1.0mm i.d.×25cm	228-12813-02	
	Shim-pack MBC-ACN	1.0mm i.d.×50cm	228-12813-05	
	Shim-pack MBC-C <sub>8</sub>	1.0mm i.d.×25cm	228-12814-02	
	Shim-pack MBC-C <sub>8</sub>	1.0mm i.d.×50cm	228-12814-05	
Shim-pack-MRC	Shim-pack MRC-SIL	6.0mm i.d.×25cm	228-23461-92	P.21
	Shim-pack MRC-ODS	6.0mm i.d.×25cm	228-23464-92	
	Shim-pack MRC-C <sub>8</sub>	6.0mm i.d.×25cm	228-24381-92	
	Shim-pack MRC-TMS	6.0mm i.d.×25cm	228-24382-92	
	Shim-pack MRC-NH <sub>2</sub>	6.0mm i.d.×25cm	228-24383-92	
	Shim-pack MRC-CN	6.0mm i.d.×25cm	228-24384-92	
Shim-pack PA	Shim-pack PA-DEAE	8.0mm i.d.×10cm	228-20758-91	P.33
	Shim-pack PA-DEAE	20mm i.d.×10cm	228-20758-92	
	Shim-pack PA-QA	8.0mm i.d.×10cm	228-20759-91	
	Shim-pack PA-QA	20mm i.d.×10cm	228-20759-92	
	Shim-pack PA-CM	8.0mm i.d.×10cm	228-20760-91	
	Shim-pack PA-CM	20mm i.d.×10cm	228-20760-92	
	Shim-pack PA-SP	8.0mm i.d.×10cm	228-20761-91	
	Shim-pack PA-SP	20mm i.d.×10cm	228-20761-92	
Shim-pack PAG	Shim-pack PAG-DEAE	8.0mm i.d.×1cm	228-20762-91	P.33
	Shim-pack PAG-QA	8.0mm i.d.×1cm	228-20763-91	
	Shim-pack PAG-CM	8.0mm i.d.×1cm	228-20764-91	
	Shim-pack PAG-SP	8.0mm i.d.×1cm	228-20765-91	
Shim-pack-PRC	Shim-pack PRC-SIL	20mm i.d.×25cm	228-23461-93	P.21
	Shim-pack PRC-SIL (K)	30mm i.d.×25cm	228-23461-94	
	Shim-pack PRC-SIL (L)	50mm i.d.×25cm	228-23461-95	
	Shim-pack PRC-SIL (H)	20mm i.d.×25cm	228-23461-91	
	Shim-pack PRC-ODS	20mm i.d.×25cm	228-23464-93	

Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack-PRC	Shim-pack PRC-ODS (K)	30mm i.d.×25cm	228-23464-94	P.21
	Shim-pack PRC-ODS (L)	50mm i.d.×25cm	228-23464-95	
	Shim-pack PRC-ODS (H)	20mm i.d.×25cm	228-23464-91	
	Shim-pack PRC-C <sub>8</sub>	20mm i.d.×25cm	228-24381-93	
	Shim-pack PRC-C <sub>8</sub> (H)	20mm i.d.×25cm	228-24381-91	
	Shim-pack PRC-TMS	20mm i.d.×25cm	228-24382-93	
	Shim-pack PRC-TMS (H)	20mm i.d.×25cm	228-24382-91	
	Shim-pack PRC-NH <sub>2</sub>	20mm i.d.×25cm	228-24383-93	
	Shim-pack PRC-NH <sub>2</sub> (H)	20mm i.d.×25cm	228-24383-91	
	Shim-pack PRC-CN	20mm i.d.×25cm	228-24384-93	
	Shim-pack PRC-CN (H)	20mm i.d.×25cm	228-24384-91	
Shim-pack PREP	Shim-pack PREP-SIL	20mm i.d.×25cm	228-00814-91	P.28
	Shim-pack PREP-SIL (H)	20mm i.d.×25cm	228-17880-91	
	Shim-pack PREP-SIL (H) ·Kit	4.6mm i.d.×25cm 20mm i.d.×25cm	228-17887-91	
	Shim-pack PREP-ODS	20mm i.d.×25cm	228-00815-91	
	Shim-pack PREP-ODS (H)	20mm i.d.×25cm	228-17881-91	
	Shim-pack PREP·ODS (H) ·Kit	4.6mm i.d.×25cm 20mm i.d.×25cm	228-17888-91	
	Shim-pack PREP-C <sub>8</sub>	20mm i.d.×25cm	228-00816-91	
	Shim-pack PREP-C <sub>8</sub> (H)	20mm i.d.×25cm	228-17882-91	
	Shim-pack PREP-TMS	20mm i.d.×25cm	228-00817-91	
	Shim-pack PREP-TMS (H)	20mm i.d.×25cm	228-17883-91	
	Shim-pack PREP-CN	20mm i.d.×25cm	228-00818-91	
	Shim-pack PREP-CN (H)	20mm i.d.×25cm	228-17884-91	
	Shim-pack PREP-Phenyl	20mm i.d.×25cm	228-00819-91	
	Shim-pack PREP-Phenyl (H)	20mm i.d.×25cm	228-17885-91	
	Shim-pack PREP-NH <sub>2</sub>	20mm i.d.×25cm	228-17879-91	
	Shim-pack PREP-NH <sub>2</sub> (H)	20mm i.d.×25cm	228-17886-91	
	Shim-pack PREP-SIL (K)	30mm i.d.×25cm	228-18273-91	
	Shim-pack PREP-ODS (K)	30mm i.d.×25cm	228-18319-91	
Shim-pack PREP-SIL (L)	50mm i.d.×25cm	228-18274-91		
Shim-pack PREP-ODS (L)	50mm i.d.×25cm	228-18320-91		

Column series	Column name	Dimension	Cat.No.	Reference
Shim-pack SBC	Shim-pack SBC-ODS	2.5mm i.d.×15cm	228-17268-91	P.30
	Shim-pack SBC-C <sub>8</sub>	2.5mm i.d.×15cm	228-17269-91	
	Shim-pack SBC-SIL	2.5mm i.d.×15cm	228-17270-91	
Shim-pack SCR	Shim-pack SCR-101N	7.9mm i.d.×30cm	228-07730-92	P.41
	Shim-pack SCR-101C	7.9mm i.d.×30cm	228-17889-91	
	Shim-pack SCR-101P	7.9mm i.d.×30cm	228-17890-91	
	Shim-pack SCR-101H	7.9mm i.d.×30cm	228-07730-93	
	Shim-pack SCR-102H	8.0mm i.d.×30cm	228-17893-91	
	Guard Column SCR (N)	4.0mm i.d.×5cm	228-09619-92	
	Guard Column SCR (C)	4.0mm i.d.×5cm	228-17891-91	
	Guard Column SCR (P)	4.0mm i.d.×5cm	228-17892-91	
	Guard Column SCR (H)	4.0mm i.d.×5cm	228-09619-93	
	Guard Column SCR-102 (H)	6.0mm i.d.×5cm	228-17924-91	
Shim-pack SPC	Shim-pack SPC-RP3	4.0mm i.d.×3cm	228-33713-91	P.49
	Shim-pack SPC-RP2	4.6mm i.d.×1cm	228-18838-91	
	Shim-pack SPC-AE1	4.0mm i.d.×1cm	228-17990-91	
Shim-pack WAX	Shim-pack WAX-1	4.0mm i.d.×5cm	228-16225-91	P.33
	Shim-pack WAX-2	4.0mm i.d.×5cm	228-16365-91	
Shim-pack WCX	Shim-pack WCX-1	4.0mm i.d.×5cm	228-16366-91	P.33

# 7 Sample and data

Compound	Page	Figure
$\alpha$ - AAA	37	51
$\alpha$ · A · B · A	79	164
$\gamma$ - ABA	37	51
Acenaphthene	16	3
Acetaldehyde	18	11
Acetaldehyde	71	117
Acetaminophen	77	153
Acetate	73	126
Acetic acid	43	64
Acetic acid	72	122
Acetic acid	73	130
Acetone	18	11
Acetylsalicylic acid	26	21
Acetylsalicylic acid	78	155
Aceumlnophen	26	21
Acrolein	18	11
Adenine	25	16
Adenosine	25	16
Adenosine	32	34
Adenylate kinase	44	66
ADP	18	6
ADP	35	42
ADP	74	132
ADP	78	158
$\beta$ - AiBA	37	51
Ala	37	50
Ala	37	51
Ala	71	119
Ala	74	137
Ala	76	149
Ala	79	164
$\beta$ - Ala	37	51
$\beta$ - Ala	79	164
Albumin	44	67
Albumin	48	76
Amitriptyline	16	3
AMP	18	6
AMP	35	42
AMP	74	132
AMP	78	158
Amygdalin	75	143
$\alpha$ - AnBA	37	51
Ans	37	51
Arabinose	43	63
Arg	37	50
Arg	37	51
Arg	71	119
Arg	73	137
Arg	76	149
Arg	79	164
Ascorbic acid	27	27
Ascorbic acid	68	113
Asn	37	51
Asn	71	119
Asn	79	164
Asp	37	50
Asp	37	51
Asp	71	119
Asp	74	137
Asp	76	149
Aspartame	17	4
ATP	18	6
ATP	35	42
ATP	74	132
ATP	78	158
Atropine	25	17
Bacitracin	61	104
Benzaldehyde	18	11
Benzanilide	21	
Benzene	21	
Benzene	40	54
Benzoic acid	17	4
Benzoic acid	68	114
Benzoic acid	72	123

Compound	Page	Figure
Benzoic acid	74	135
Benzothorium Chloride	75	141
Berberine	18	9
Berberine	75	142
Big gastrin	60	96
Biotin	68	113
Biphenyl	72	124
Blue dextran 2000	44	65
Borate	81	175
Bovine serum albumin	36	45
Br <sup>-</sup>	46	70
Br <sup>-</sup>	82	179
Bradykinin	61	104
Bromovalerylurea	26	21
Bromovalerylurea	32	37
BSA	68	115
Butanol	60	94
2 - butanone	18	11
Butylaldehyde	71	117
Butyraldehyde	18	11
Ca <sup>2+</sup>	46	72
Caffeine	17	4
Caffeine	18	7
Caffeine	25	13
Caffeine	26	21
Caffeine	32	37
Caffeine	68	113
Caffeine	72	120
Caffeine	75	138
Caffeine	77	153
Calcium pantothenate	68	113
Calcium pantothenate	72	120
Camphor	76	144
Car	37	51
Carbamazepin	72	121
Carbamazepin	78	154
Carbonic anhydrase	36	45
Carbonic anhydrase	58	90
4 - Carboxybenzaldehyde	81	173
Cd <sup>2+</sup>	46	73
CDP	35	42
CDP	78	158
Cellobiose	37	49
Chlorhexidine	18	8
Chlorhexidine gluconate	76	146
Chondroitin sulfate	76	147
Chlorpheniramine maleate	26	18
Chymotrypsin	44	65
$\alpha$ - Chymotrypsinogen A	35	38
$\alpha$ - Chymotrypsinogen A	36	46
$\alpha$ - Chymotrypsinogen A	36	47
$\alpha$ - Chymotrypsinogen A	36	48
$\alpha$ - Chymotrypsinogen A	48	74
$\alpha$ - Chymotrypsinogen A	48	76
$\alpha$ - Chymotrypsinogen A	58	91
$\alpha$ - Chymotrypsinogen A	61	101
$\alpha$ - Chymotrypsinogen A	64	107
Cinchonine	61	100
cis - and trans - Isoadhumulone	51	80
cis - and trans - Isocohumulone	51	80
cis - and trans - Isohumulone	51	80
Cit	37	51
Cit	79	164
Citrate	73	126
Citric acid	43	64
Citric acid	72	122
Cl	46	70
Cl <sup>-</sup>	73	126
Cl <sup>-</sup>	77	152
Cl <sup>-</sup>	81	174
Cl <sup>-</sup>	82	177
CMP	35	42
CMP	78	158
Co <sup>2+</sup>	46	73

Compound	Page	Figure
Conalbumin	36	45
Creatinine	60	97
Crotonaldehyde	18	11
Cs <sup>+</sup>	46	71
CTP	35	42
CTP	78	158
Cu <sup>2+</sup>	46	73
Cys	79	164
Cysteine	76	149
Cystine	37	50
Cystine	37	51
Cytidine	25	16
Cytidine	32	34
Cytochrome C	36	43
Cytochrome C	44	65
Cytochrome C	44	66
Cytochrome C	48	74
Cytochrome C	48	76
Cytochrome C	61	101
Cytochrome C	68	115
Cytosine	25	16
1-Decanol	60	94
Dehydroacetic acid	68	114
Dexamethasone	27	24
Dibutyl phthalate	26	22
Dibutylphthalate	56	87
Didecylphthalate	56	87
Diethyl phthalate	26	22
Diheptylphthalate	56	87
Dihydrocodeine phosphate	77	153
Dimethyl phthalate	26	22
Dioctyl phthalate	26	22
1-Dodecanol	60	94
DOMA	51	81
DOPAC	51	81
Dulcoside A	31	32
$\beta$ -Endorfin	60	95
$\beta$ -Endorfin	60	96
Enolase	44	66
Ergosterol	26	23
Erythorbic acid	27	27
Estradiol	32	35
Estriol	32	35
Estrone	32	35
Ethanol	43	62
Ethenzamide	26	21
Ethenzamide	32	37
Ethinylestradiol	75	139
Ethofyline (L.S.)	75	138
Ethofyline (L.S.)	78	156
Ethosuximide	72	121
Ethosuximide	78	154
Ethyl acetate	25	15
Ethyl butyrate	25	15
Ethyl caproate	25	15
Ethyl propionate	25	15
Ethyl valerate	25	15
Ethylbenzoate	32	33
Ethylene glycol	60	94
Ethylenglycol	44	69
Ethylparaben	68	114
F <sup>-</sup>	46	70
Fe-EDTA	81	176
Fe <sup>2+</sup>	46	73
Fe <sup>3+</sup>	46	73
Fluocinolone acetonide	76	148
Folic acid	68	113
Formaldehyde	18	11
Formaldehyde	71	117
Formic acid	43	64
Formic acid	73	130
Fructose	27	26
Fructose	37	49
Fructose	43	62

Compound	Page	Figure
Fructose	65	109
Fructose	65	111
Fructose	73	127
Fructose	73	128
Fructose	73	129
Fumaric acid	43	64
Galactose	37	49
Galactose	43	63
Galactose	73	129
GDP	35	42
GDP	78	158
Glucose	37	49
Gln	37	51
Gln	71	119
Gln	79	164
Globulin	64	108
Glu	37	50
Glu	37	51
Glu	71	119
Glu	74	137
Glu	76	149
Glu	79	164
Glucose	27	26
Glucose	43	61
Glucose	43	62
Glucose	43	63
Glucose	65	109
Glucose	65	110
Glucose	65	111
Glucose	65	112
Glucose	73	127
Glucose	73	128
Glucose	73	129
Glutamate dehydrogenase	44	66
Gly	37	50
Gly	37	51
Gly	71	119
Gly	74	137
Gly	76	149
Gly	79	164
Gly-Gly-Ala	61	103
Gly-Gly-Leu	61	103
Gly-Tyr	44	65
Glycerol	27	26
Glycerol	43	62
Glycerol	43	63
$\beta$ -Glycyrhrrtinic acid	77	150
GMP	35	42
GMP	78	158
GTP	35	42
GTP	78	158
Guaninc	25	16
Guanosine	25	16
Guanosine	32	34
Hemoglobin A	35	39
Hemoglobin A	36	44
Hemoglobin C	35	39
Hemoglobin F	35	39
Hemoglobin F	36	44
Hemoglobin S	35	39
Hemoglobin S	36	44
Hesperidin	17	5
Hesperodine	74	133
Hexanaldehyde	18	11
Hexanol	60	94
Hexobarbital (L.S.)	78	154
His	37	50
His	37	51
His	71	119
His	74	137
His	76	149
His	79	164
Histamine	79	163
Homovanillic acid (HVA)	79	161

Compound	Page	Figure
Human serum albumin	44	65
HVA	51	81
Hydrocortisone acetate	27	24
HyLys	37	51
Hypoxanthine	18	6
Hypoxanthine	74	132
Hypoxanthine	78	157
Hypro	37	51
IgG	44	67
IgG	58	92
Ile	37	50
Ile	37	51
Ile	71	119
Ile	73	137
Ile	76	149
Ile	79	164
IMP	18	6
IMP	74	132
Inosine	18	6
Inosine	74	132
Insulin	60	95
Insulin	60	96
Insulin	61	104
Insulin B chain	61	104
Insulin b chain	60	95
iso-Butylparaben	68	114
iso-Maltose	73	129
Isomaltoheptaose	65	112
Isomaltohexaose	65	112
Isomaltononaose	65	112
Isomaltooctaose	65	112
Isomaltopentaose	65	112
Isomaltose	65	112
Isomaltotetraose	65	112
Isomaltotriose	65	112
K <sup>+</sup>	46	71
K <sup>+</sup>	46	72
K <sup>+</sup>	82	178
$\alpha$ -Ketoglutaric acid	43	64
$\alpha$ -Lactalbumin	36	45
$\alpha$ -Lactalbumin	58	90
Lactate	73	126
Lactate dehydrogenase	44	66
Lactic acid	43	64
Lactic acid	73	130
$\beta$ -Lactoglobulin	44	65
$\beta$ -Lactoglobulin A	36	45
$\beta$ -Lactoglobulin B	36	45
Lactose	27	26
Lactose	37	49
Lactose	65	109
Leu	37	50
Leu	37	51
Leu	71	119
Leu	73	137
Leu	76	149
Leu-Enkephalin	61	103
Leu-Enkephalin	61	104
Levulinic acid	43	64
Li <sup>+</sup>	46	71
Iodoform	76	145
Lou	79	164
Irganox1076	81	172
Lys	37	50
Lys	37	51
Lys	71	119
Lys	74	137
Lys	76	149
Lys	79	164
Lys-Bradykinin	61	104
Lysophosphatidylcholine (Lysolecithin)	25	12
Lysozyme	35	38
Lysozyme	36	46
Lysozyme	36	47

Compound	Page	Figure
Lysozyme	36	48
Lysozyme	44	65
Lysozyme	44	68
Lysozyme	48	76
Lysozyme	58	91
Lysozyme	61	101
Lysozyme	61	104
Lysozyme	64	107
Lysozyme	68	115
m-tolualdehyde	18	11
Malate	73	126
Malic acid	43	64
Malic acid	72	122
Malic acid	73	130
Malonic acid	72	122
Maltoheptaose	65	110
Maltohexaose	65	110
Maltopentaose	65	110
Maltose	27	26
Maltose	37	49
Maltose	43	61
Maltose	43	62
Maltose	65	110
Maltose	65	111
Maltose	73	129
Maltotetraose	65	110
Maltotriose	65	110
Mannitol	43	63
Mannose	37	49
Mastoparan	61	104
1-Me-His	37	51
3-Me-His	37	51
Menthol	76	144
Mercaptoalbumin	64	108
Met	37	50
Met	37	51
Met	71	119
Met	74	137
Met	76	149
Met	79	164
Met-Enkephalin	61	103
Met-Enkephalin	61	104
Methacrolein	18	11
Methylparabon	68	114
Methyl benzoate	60	97
Methyl benzoate	72	124
Methyl salicylate	76	144
Methylbenzoate	32	33
Methylephedrine	77	153
Mg <sup>2+</sup>	46	72
MHPG	51	81
Mn <sup>2+</sup>	46	73
Moltotriose	43	61
Myoglobin	35	38
Myoglobin	36	46
Myoglobin	36	47
Myoglobin	36	48
Myoglobin	44	65
Myoglobin	48	74
Myoglobin	48	76
Myoglobin	58	90
Myoglobin	58	91
Myoglobin	61	104
Myoglobin	64	107
Myoglobin	68	115
N1-Acetylspermidine	79	160
N1-Acetylspermine	79	160
N-acetylprocinamide	15	1
N-Acetylputrescine	79	160
n-Butylbenzoate	32	33
n-Butylparaben	68	114
n-Hexyl benzoate	60	97
n-Propyl benzoate	60	97
n-Propylbenzene	39	53

Compound	Page	Figure
n-Propylbenzene	40	54
n-Propylbenzoate	32	33
n-Propylparaben	68	114
Na <sup>+</sup>	46	71
Na <sup>+</sup>	46	72
Na <sup>+</sup>	82	178
NAD	26	19
NADH	26	19
Naphazoline chloride	26	18
Naphthalene	72	124
Neostigmine sulfate	26	18
Neurotensin	61	104
NH <sub>4</sub> <sup>+</sup>	46	71
NH <sub>4</sub> <sup>+</sup>	46	72
NH <sub>4</sub> <sup>+</sup>	82	178
NH <sub>3</sub> + Et · NH <sub>2</sub>	37	51
Ni <sup>2+</sup>	46	73
Nicotinamide	18	7
Nicotinamide	25	13
Nicotinamide	68	113
Nicotinamide	72	120
Nicotinamide	75	138
Nicotinic acid	25	13
Nicotinic acid	72	120
Nicotinic acid	75	138
NO <sub>2</sub>	46	70
NO <sub>2</sub> <sup>-</sup>	82	177
NO <sub>3</sub> <sup>-</sup>	46	70
NO <sub>3</sub> <sup>-</sup>	73	126
NO <sub>3</sub> <sup>-</sup>	82	177
NO <sub>3</sub> <sup>-</sup>	82	179
NO <sub>3</sub> <sup>-</sup>	77	152
Nonmercaptoalbumin	64	108
Octanol	60	94
25-OH-D <sub>3</sub>	78	159
Orn	37	51
Orn	79	164
Ovalbumin	35	38
Ovalbumin	36	43
Ovalbumin	44	65
Ovalbumin	44	68
Ovalbumin	58	91
Ovalbumin	61	101
Ovalbumin	68	115
Oxine	16	2
Oxine	18	10
p-Bromoacetanilide	21	
p-Et · NH <sub>2</sub>	79	164
p-Et · NH <sub>2</sub>	37	51
p-Hydroxy benzoic acid	68	114
p-Ser	37	51
p-Toluic acid	81	173
Paeoniflorin	55	82
Paeonol	55	84
Pantothenic acid	25	13
Pantothenic acid	75	138
PEG 1000	44	69
PEG 4000	43	62
PEG 4000	44	69
PEG 6000	44	69
Pentanal	71	117
Phc	74	137
Phe	37	50
Phe	71	119
Phe	76	149
Phe	79	164
Phenacetin	32	37
Phenobarbital	72	121
Phenobarbital	78	154
Phenol	15	1
Phenol	82	180
β-Phenylchalcone (I.S.)	51	80
Phenytoin	72	121
Phenytoin	78	154

Compound	Page	Figure
Phosphatidylcholine (Lecithin)	25	12
Phosphoric acid	43	64
Phridoxine	18	7
PO <sub>4</sub> <sup>3-</sup>	46	70
Polyethylene glycol	63	105
Polyethylene glycol	63	106
Polysaccharides	43	61
Polystyrene 1,100K	39	53
Polystyrene 1.35K	39	53
Polystyrene 115K	39	53
Polystyrene 33K	39	53
Polystyrene 410K	39	53
Polystyrene 7.6K	39	53
Polystyrene M.w. 9,120K	39	53
Polystyrene Mw 2,800	40	54
Polystyrene Mw 300	40	54
Polystyrene Mw 42,800	40	54
Pre	37	51
Primidone	72	121
Primidone	78	154
Pro	37	50
Pro	37	51
Pro	71	119
Pro	74	137
Pro	76	149
Pro	79	164
Propionaldehyde	18	11
Propionic acid	72	122
Propranolol	16	3
Propylaldehyde	71	117
Pullulan	63	105
Pullulan	63	106
Putrescine	79	160
Pyridoxine	25	13
Pyridoxine	72	120
Pyridoxine	75	138
Pyridoxine hydrochloride	68	113
Pyroglutamic acid	43	64
Pyroglutamic acid	73	130
Pyruvic acid	43	64
Quinine	61	100
Rafinose	65	111
Ramnose	65	111
Rb <sup>+</sup>	46	71
Rebaudioside A	31	32
Rebaudioside A	74	134
Rebaudioside C	31	32
Rhamnose	37	49
Riboflavin	72	120
Riboflavin phosphate	72	120
Riboflavin	25	13
Riboflavin	68	113
Riboflavin phosphate	18	7
Riboflavin phosphate	25	13
Riboflavin	75	138
Ribonuclease	61	101
Ribonuclease A	35	38
Ribonuclease A	36	46
Ribonuclease A	36	47
Ribonuclease A	36	48
Ribonuclease A	44	65
Ribonuclease A	48	74
Ribonuclease A	58	91
Ribonuclease A	64	107
Ribonuclease A	68	115
Ribose	37	49
Saccharides	42	60
Saccharin	25	14
Salicylamide	26	21
Salicylic acid	68	114
Salicylic acid	78	155
Sar	37	51
Scopolamine	25	17
Ser	37	50

Compound	Page	Figure
Ser	37	51
Ser	71	119
Ser	73	137
Ser	76	149
Ser	79	164
SO <sub>4</sub> <sup>2-</sup>	73	126
SO <sub>4</sub> <sup>2-</sup>	46	70
SO <sub>4</sub> <sup>2-</sup>	77	152
SO <sub>4</sub> <sup>2-</sup>	82	177
Sorbic acid	25	14
Sorbic acid	68	114
Sorbic acid	72	123
Sorbic acid	74	135
Sorbitol	43	62
Sorbitol	43	63
Sorbitol	73	128
Sphingomyelin	25	12
Stevioside	31	32
Stevioside	74	134
Substance P	60	95
Substance P	61	104
Succinate	73	126
Succinic acid	72	122
Succinic acid	73	130
Succinic acid	43	64
Sucrose	27	26
Sucrose	37	49
Sucrose	43	63
Sucrose	65	109
Sucrose	65	111
Sucrose	73	127
Sucrose	73	128
Tartarate	73	126
Tartaric acid	72	122
Tau	37	51
Tau	79	164
Taurine	75	140
1-Tetradecanol	60	94
Theanine	74	136
Theophylline	78	156
Thiamine	18	7
Thiamine	25	13
Thiamine	68	113
Thiamine	72	120
Thiamine	75	138
Thr	71	119
Thr	74	137
Thr	76	149
Thr	79	164
Thy	37	50
Thy	37	51
Thymidine	32	34
Thymine	25	16
Thymol	76	144
Thyroglobulin	44	65
Tocol	32	36
$\alpha$ -Tocopherol	27	25
$\alpha$ -Tocopherol	31	31
$\alpha$ -Tocopherol	32	36
$\alpha$ -Tocopherol	73	131
$\beta$ -Tocopherol	27	25
$\beta$ -Tocopherol	31	31
$\beta$ -Tocopherol	32	36
$\beta$ -Tocopherol	73	131
$\gamma$ -Tocopherol	27	25
$\gamma$ -Tocopherol	31	31
$\gamma$ -Tocopherol	32	36
$\gamma$ -Tocopherol	73	131
$\delta$ -Tocopherol	27	25
$\delta$ -Tocopherol	31	31
$\delta$ -Tocopherol	32	36
$\delta$ -Tocopherol	73	131
Transferrin	44	65
Transferrin	48	74

Compound	Page	Figure
Transferrin	58	90
Trp	71	119
Trp	76	149
Trp	79	164
Try	76	149
Trypsin inhibitor	36	43
Trypsin inhibitor	58	90
Tyr	37	50
Tyr	37	51
Tyr	71	119
Tyr	74	137
Tyr	79	164
$\Delta$ Tyr	71	119
UDP	35	42
UDP	78	158
UMP	35	42
UMP	78	158
Undine	32	34
Unknown	25	16
Uracil	25	16
Uracil	72	124
Urease	35	40
Uric acid	64	108
Uric acid	78	157
Uridine	25	16
UTP	35	42
UTP	78	158
Val	37	50
Val	37	51
Val	71	119
Val	73	137
Val	76	149
Val	79	164
Vanillylmanderic acid (VMA)	79	161
Veraldehyde	18	11
Vitamin A	71	118
Vitamin A acetate	26	20
Vitamin A palmitate	26	20
Vitamin B <sub>6</sub>	61	99
Vitamin B <sub>1</sub>	61	99
Vitamin B <sub>12</sub>	61	99
Vitamin B <sub>12</sub>	77	151
Vitamin B <sub>2</sub>	61	99
Vitamin D <sub>2</sub>	26	23
Vitamin D <sub>2</sub>	60	98
Vitamin D <sub>2</sub>	71	118
Vitamin D <sub>3</sub>	60	98
Vitamin E	26	20
Vitamin E	71	118
Vitamin E acetate	26	20
Vitamin K <sub>2</sub>	26	20
VLA	51	81
VMA	51	81
Xanthine	78	157
Xylose	27	26
Xylose	37	49
Xylose	43	63
$\gamma$ -Globulin	44	65
Zn <sup>2+</sup>	46	73





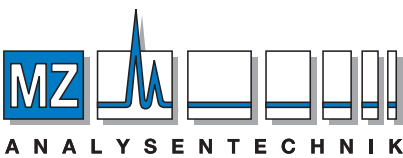


JQA-0376

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