# Silica-based for protein analysis:

TSKgel SW mAb TSKgel SW TSKgel SWxL **TSKgel SuperSW** 

**Polymer-based for desalting:** 

TSKgel BioAssist DS Columns

**Polymethacrylate-based for water-soluble** polymers analysis:

**TSKgel PW** TSKgel PWxL TSKgel PWxL-CP TSKgel SuperMultiporePW

# **Polymethacrylate-based for polar** organic-soluble polymers analysis:

**TSKgel Alpha** TSKgel SuperAW

**Polystyrene-divinylbenzene-based for** organic-soluble polymers analysis:

TSKgel HxL TSKgel Hhr TSKgel SuperH **TSKgel SuperHZ** TSKgel SuperMultiporeHZ

# About: TSKgel H Series Size Exclusion Columns

TSKgel H series columns are recommended for the analysis of organic-soluble polymers and are packed with spherical particles composed of polystyrene crosslinked with divinylbenzene (PS-DVB). This series includes TSKgel HxL, HHR, SuperH, Super HZ, and SuperMultiporeHZ columns. Each line of columns within this series differs in degree of inertness and operating temperature range.

The Super prefix designates short (15 cm) columns packed with particles as small as  $3 \mu m$ . The smaller particle allows for equivalent resolution to conventional TSKgel HxL columns, with 50% reduction in analysis time due to the shorter column length. The TSKgel Super series columns are an excellent choice for high throughput polymer analysis.

The TSKgel HxL columns are conventional GPC columns of 7.8 mm ID × 30 cm. The column line consists of eight columns with different pore sizes, TSKgel G1000HxL through TSKgel G7000HxL, and three columns with an extended linear range of the calibration curve, TSKgel GMHxL, TSKgel GMHxL-L and TSKgel MultiporeHxL-M. The 5 µm particles in the TSKgel MultiporeHxL-M column contain a broad range of pore sizes. This innovative approach essentially creates a linear calibration curve within each particle. As a result, columns with an extended linear calibration curve can now be prepared without mixing particles of different pore sizes.

The main characteristics of TSKgel HxL columns are: ultra-low sample adsorption, i.e., the columns show true size exclusion behavior for most polymers, limited solvent range, and a maximum operating temperature of 60 °C for TSKgel G1000HxL - G3000HxL, and 80 °C for the remaining columns in the TSKgel HxL column line.

• The TSKgel HHR column line consists of eight conventional GPC columns of 7.8 mm ID × 30 cm with different pore sizes, TSKgel G1000HHR through TSKgel G7000HHR, and seven mixed bed columns, in which particles with different pore sizes are blended to provide an extended linear calibration curve. The mixed bed columns feature increasing linear calibration ranges, from TSKgel GMHHR-L, GMHHR-N, GMHHR-M, to GMHHR-H. The main characteristic of these TSKgel HHR columns is a broad solvent range.

In addition, four TSKgel HHR mixed bed columns are available for ultra-high temperature analysis. The maximum operating temperature of these columns is 220 °C.

 The TSKgel SuperH column line consists of eight columns of 6.0 mm ID × 15 cm with different pore sizes, TSKgel SuperH1000 through TSKgel SuperH7000, and four mixed bed columns with an extended linear range of the calibration curve. The mixed bed columns feature increasing linear calibration ranges, from TSKgel SuperHM-L, SuperHM-N, SuperHM-M, to SuperHM-H. TSKgel SuperH columns are high efficiency/high throughput versions of the conventional TSKgel HHR columns. Both column types are based on the same bead chemistry.

The main characteristics of TSKgel SuperH columns are: a maximum operating temperature of 140 °C and the ability to use a broad range of solvents.

• The TSKgel SuperHZ column line consists of five columns of 4.6 mm ID × 15 cm and 6.0 mm ID × 15 cm with different pore sizes, TSKgel SuperHZ1000 through TSKgel SuperHZ4000, and three columns with an extended linear range of the calibration curve. The mixed bed columns feature increasing linear calibration ranges, from TSKgel SuperHZM-L, SuperHZM-N to SuperHZM-H.

The main characteristics of TSKgel SuperHZ columns are: developed for high throughput, high efficiency GPC applications such as those encountered in combinatorial chemistry experiments, ultra-low sample adsorption, limited solvent range, and a maximum operating temperature of 60 °C for TSKgel SuperHZ1000 - SuperHZ3000 and 80 °C for the remaining columns in the TSKgel SuperHZ line.

 The TSKgel SuperMultiporeHZ column line consists of three columns of 4.6 mm ID × 15 cm with particles sizes of 3, 4 and 6 µm. The particles in TSKgel SuperMultiporeHZ columns are monodisperse in size and exhibit a broad range of pore sizes. Each particle, by design, has an extended linear calibration curve, thereby greatly diminishing chromatograms with inflection points.

A comparison of TSKgel H series columns is detailed in Table 18. The cross-linking of the polystyrene particles in TSKgel H series columns ensures minimal shrinking and swelling of the column bed when the organic solvent is changed according to the solvent recommendations outlined in Table 19. Suggested flow rates for solvent exchange in TSKgel SuperH and HHR columns are outlined in Table 20. Table 21 lists the recommended solvents by application for TSKgel H series columns.



TSKgel series	SuperMultiporeHZ	SuperHZ	SuperH	HxL	Ннв
Application focus	Ultra-high performance with a low dead volume and a wide pore distribution in each particle for superior linearity	High throughput polymer analysis with ultra- low polymer adsorption, limited solvent compatibility range	High throughput polymer analysis with expanded solvent compatibility range	Conventional polymer analysis with ultra- low polymer adsorption, limited solvent compatibility range	Conventional polymer analysis with expanded solvent compatibility range
Particle size	3 µm, 4 µm, and 6 µm, depending on pore size	3 μm, 5 μm, and 10 μm, depending on pore size	3 μm and 5 μm, depending on pore size	5 μm, 6 μm, 9 μm, and 13 μm, depending on pore size	5 μm, 13 um, 20 μm, and 30 μm
Particle matrix	Polystyrene divinylbenzene (PS-DVB)	Polystyrene divinylbenzene (PS-DVB)	Polystyrene divinylbenzene (PS-DVB)	Polystyrene divinylbenzene (PS-DVB)	Polystyrene divinylbenzene (PS-DVB)
Number of solvent substitutions	None	One time only	Several <sup>1</sup>	One time only	Several <sup>1</sup>

### Table 18: Comparison of TSKgel H series columns

<sup>1</sup>After switching to a very polar solvent such as acetone, switching back to a nonpolar solvent is not recommended.

TSKgel series	Shipping solvent*	Can be replaced with:	
	Tetrahydrofuran <sup>3,4</sup>	benzene, chloroform, toluene, xylene, dichloromethane, dichloroethane	
SuperHZ and Hx∟¹	Acetone**	carbon tetrachloride <sup>5</sup> , <i>o</i> -dichlorobenzene, dimethylformamide, dodecane, dimethyl sulfoxide, dioxane, ethylacetate, FC-113, hexane, pyridine, hexafluoroisopropanol/chloroform, methyl ethyl ketone, quinoline, cyclohexane	
	Chloroform**	<i>m</i> -cresol in chloroform, up to 10% hexafluoroisopropanol/chloroform	
	Dimethylformamide	dimethyl sulfoxide, dioxane, tetrahydrofuran, toluene	
SuperH and HHR <sup>2</sup>	Tetrahydrofuran <sup>3</sup>	acetone, ethanol, quinoline, benzene, <i>o</i> -dichlorobenzene, ethyl acetate, dodecane, FC-113, carbon tetrachloride <sup>5</sup> , dichloromethane, dichloroethane, trichloroethane, <i>n</i> -hexane, cyclohexane, xylene, tetrahydrofuran, chloroform, 1,4-dioxane, hexafluoroisopropanol, toluene, 1-chloronaphthalene, N,N-dimethylacetoacetamide, methyl ethyl ketone, trichlorobenzene, <i>m</i> -cresol, dimethylformamide, methylpyrrolidone, <i>o</i> -chlorophenol/chloroform, dimethyl sulfoxide, pyridine	
SuperMultiporeHZ	Tetrahydrofuran <sup>3</sup>	Cannot be replaced. TSKgel SuperMultiporeHZ columns can be used only in tetrahydrofuran	

Table 19: Solvent compatibility for TSKgel H series columns

<sup>1</sup> In case of TSKgel SuperHZ and HxL, keep flow rate as mentioned below during solvent change. Solvent can be changed one way/one time only.

TSKgel HxL: below <0.5 mL/min

TSKgel SuperHZ (4.6 mm ID): below <0.15 mL/min

TSKgel SuperHZ (6.0 mm ID): below <0.3 mL/min

<sup>2</sup> In case of TSKgel SuperH and H<sub>R</sub>, see Table 20 below for appropriate flow rates for solvent exchange. After switching to a very polar solvent, switching to a nonpolar solvent is not recommended.

<sup>3</sup> All TSKgel HxL, HHR, SuperHŽ, SuperH, SuperMultipore, and GMH analytical columns are shipped containing

tetrahydrofuran (THF), except the TSKgel high temperature columns, which contain o-dichlorobenzene (ODCB).

<sup>4</sup> THF in TSKgel G1000HxL columns cannot be replaced with dichloromethane or dichloroethane.

<sup>5</sup> Prolonged exposure to carbon tetrachloride can corrode the stainless steel parts of a column and an HPLC system. \* 100% methanol cannot be used with TSKgel H series columns; use this solvent with TSKgel SW or Alpha columns.

\*\* TSKgel H series columns may be specially ordered with this shipping solvent.

Table 20: Recommended flow rates (mL/min) for TSKgel SuperH and H\_{\rm HR} columns

Solvent	TSKgel SuperH, 6.0 mm ID × 15 cm	TSKgel H <sub>HR</sub> , 7.8 mm ID × 30 cm
<i>n</i> -Hexane	0.5	0.9
methyl ethyl ketone	0.4	0.7
dichloromethane, ethyl acetate	0.35	0.6
toluene, chloroform	0.3	0.5
dimethylformamide	0.2	0.4
carbon tetrachloride, pyridine	0.15	0.3
dimethyl sulfoxide, dioxane, ethanol, N-methylpyrrolidone, <i>o</i> -dichlorobenzene	0.1	0.2
quinoline, hexafluoroisopropanol, 1-chloronaphthalene	0.05	0.1

Table 21: Recommended solvents by application for TSKgel H series columns

Recommended solvent	Application
THF	polystyrene, epoxy resin, phenoxy resin, polycarbonate, polyisoprene, polyvinyl acetate, polyvinyl chloride, monoglycerides, fatty acids, polybutadiene, poly(methyl methacrylate), poly(styrene-butadiene), poly(styrene-acrylonitrile)
N,N-Dimethylformamide (DMF) + 5 mmol/L LiBr	polyvinyl chloride, polyvinyl fluoride, urea resins, polyurethane, polystyrene, polyester, polyimido ether, polyimido ester, polyphenol (aqueous solution), polyacrylonitrile
o-Dichlorobenzene (ODCB)	polyethylene, polypropylene
Chloroform	polycarboxylic ether, acrylic resin, epoxy resin, polystyrene
<i>m</i> -Cresol/Chloroform	nylon, polyester, polyamide, poly (ethylene terephthalate)
Toluene	polybutadiene, polysiloxane



# About: TSKgel HxL Size Exclusion Columns

TSKgel HxL columns are conventional GPC columns of 7.8 mm ID × 30 cm containing 5, 6, 9, or 13  $\mu$ m particles composed of PS-DVB. The TSKgel HxL column lines consists of eight columns with different pore sizes, TSKgel G1000HxL through TSKgel G7000HxL, and three columns with an extended linear range of the calibration curve, TSKgel GMHxL, TSKgel GMHxL-L and TSKgel MultiporeHxL-M.

The TSKgel HxL column line consists of the following columns:

- TSKgel G1000HxL
- TSKgel G2000HxL
- TSKgel G2500HxL
- TSKgel G3000HxL
- TSKgel G4000HxL
- TSKgel G5000HxL
- TSKgel G6000HxL
- TSKgel G7000HxL
- TSKgel GMHxL mixed bed
- TSKgel GMHxL-HT mixed bed
- TSKgel GMHxL-L mixed bed
- TSKgel MultiporeHxL-M

Three of the linear columns are mixed bed columns, in which particles with different pore sizes are blended to provide an extended linear calibration curve. The remaining column is a multi-pore column, in which each particle contains a range of pore sizes that provide a linear calibration curve. The innovative multi-pore approach, pioneered by Tosoh, is a synthetic chemistry answer to the question of how to obtain a column with an extended linear calibration curve, while mixed bed columns represent a mechanical way of obtaining a linear calibration curve. In general, Multipore columns have a smoother, more linear, calibration curve.

### **Attributes and Applications:**

Product attributes of all of the TSKgel HxL columns are shown in Table 22. These columns are for the use of conventional polymer analysis and show ultra-low polymer absorption, i.e., the columns show true size exclusion behavior for most polymers. TSKgel HxL columns are shipped in THF, with the exception of the TSKgel GMHxL-HT column, which is shipped in *o*-dichlorobenzene. These columns can be exchanged for a limited number of organic solvents. See the table within the TSKgel H series column overview for a listing of these solvents. Figures 65-66 show the calibration curves for the TSKgel HxL columns.

TSKgel column	Particle size	Pore size	Exclusion limit	Max. temp.		
G1000HxL	5 µm	1.5 nm	1,000 Da	60 °C		
G2000HxL	5 µm	2 nm	1.0 × 104 Da	60 °C		
G2500HxL	5 µm	3 nm	2.0 × 104 Da	60 °C		
G3000HxL	5 µm	7.5 nm	6.0 × 10 <sup>4</sup> Da	60 °C		
G4000HxL	5 µm	20 nm	4.0 × 10⁵ Da	80 °C		
G5000HxL	9 µm	65 nm	4.0 × 10 <sup>6</sup> Da	80 °C		
G6000HxL	9 µm	>65 nm	4.0 × 10 <sup>7</sup> Da	80 °C		
G7000HxL	9 µm	>65 nm	4.0 × 10 <sup>8</sup> Da	80 °C		

mixed pore

mixed pore

mixed pore

distribution

of pore size

sizes

sizes

sizes

broad

in each

particle

4.0 × 10<sup>6</sup> Da

 $4.0 \times 10^8 \text{ Da}$ 

4.0 × 10<sup>6</sup> Da

2.0 × 10<sup>6</sup> Da

80 °C

140 °C

80 °C

60 °C

Table 22: Product attributes

**GMH**XL

GMHxL-HT

GMHxL-L

MultiporeHxL-M

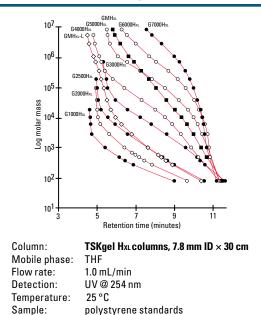
#### Figure 65: Calibration curves of TSKgel HxL columns

9 µm

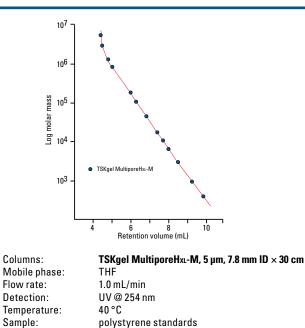
13 µm

5 µm

5 um



### Figure 66: Calibration curve of TSKgel MultiporeHxL-M column



### Phenol Resin

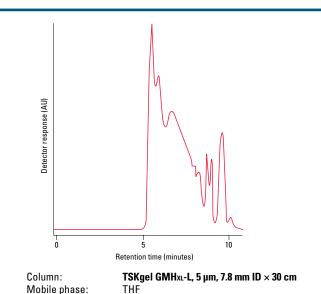
The TSKgel GMHxL-L column has been designed to provide a complete profile for high molar mass samples that contain low molar mass additives. The calibration curve for this mixed bed column is shallow in the low molar mass range of oligomers. Sample adsorption is not observed. For example, the complete profile of a phenol resin, with high resolution of the low molar mass components, is shown in Figure 68. Other applications for the TSKgel GMHxL-L column include analyses of paint materials, bond and adhesive components and synthetic polymer additives.

Figure 68: Separation of phenol resin

Flow rate:

Detection:

Sample:



1.0 mL/min

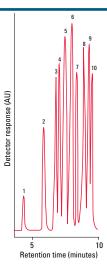
UV @ 254 nm

phenol resin

### **Phthalate Esters**

Figure 67 demonstrates the high efficiency separation on a TSKgel G1000HxL column for low molar mass phthalate esters. Resolution was close to baseline even though the molar masses of the esters differed by less than 50 Da.

Figure 67: High resolution of phthalate esters



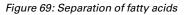
Column: Mobile phase: Flow rate: Detection: Samples:

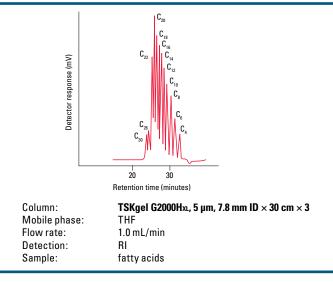
TSKgel G1000Hxt, 5 μm, 7.8 mm ID × 30 cm THF 1.0 mL/min UV @ 254 nm 1. polystyrene (1.0 × 10<sup>4</sup> Da) 2. dioctylphthalate (391 Da) 3. dibutylphthalate (278 Da) 4. diprophylphthalate (250 Da) 5. diethylphthalate (222 Da) 6. dimethylphthalate (194 Da) 7. n-propylbenzene (120 Da) 8. ethylbenzene (116 Da) 9. toluene (92 Da) 10. benzene (78 Da)



# **Fatty Acids**

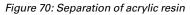
In Figure 69, two TSKgel G2000HxL columns in series separate a mixture of fatty acids ranging from C4 to C30.

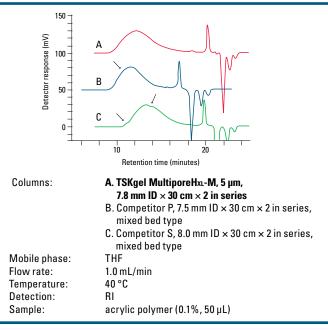




# **Acrylic Polymer**

Figure 70 shows the separation of an acrylic polymer on the TSKgel MultiporeHxL-M column compared with two commercially available mixed bed columns. The arrows illustrate the inflections seen in the chromatograms from mixed bed columns and the improvement achieved when using the TSKgel MultiporeHxL-M column.

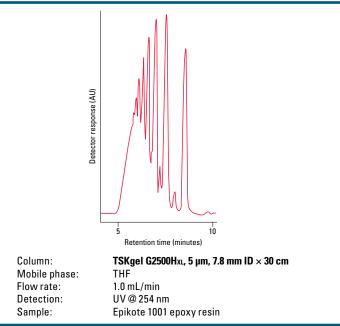




# **Epoxy Resin**

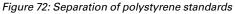
The analysis of a commercial epoxy resin, Epikote 1001, using a TSKgel G2500HxL column is shown in Figure 71.

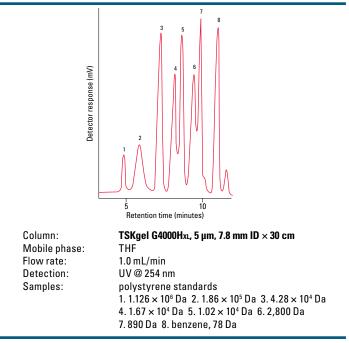
Figure 71: Separation of epoxy resin



# Polystyrene

For polymer mixtures that contain low and high molar mass compounds, a TSKgel G4000HxL column provides high resolution of samples ranging in size from benzene to  $1.126 \times 10^6$  Da as shown in Figure 72.





# About: TSKgel HHR Size Exclusion Columns

TSKgel HHR columns are conventional GPC columns with dimensions of 7.8 mm ID × 30 cm containing spherical particles composed of PS-DVB. The TSKgel HHR column line consists of eight columns with different pore sizes, TSKgel G1000HHR through TSKgel G7000HHR, and ten columns with an extended linear range of the calibration curve.

The TSKgel HHR column line consists of the following columns:

- TSKgel G1000HHR
- TSKgel G2000HHR
- TSKgel G2500HHR
- TSKgel G3000HHR
- TSKgel G4000HHR
- TSKgel G5000HHR
- TSKgel G6000Ннк
- TSKgel G7000HHR
- TSKgel GMHHR-H mixed bed
- TSKgel GMHHR-L mixed bed
- TSKgel GMHHR-M mixed bed
- TSKgel GMHHR-N mixed bed
- TSKgel G2000HHR HT
- TSKgel GMHHR-H(S) HT mixed bed
- TSKgel GMHHR-H HT mixed bed
- TSKgel G2000HHR HT2
- TSKgel GMHHR-H(S) HT2 mixed bed
- TSKgel GMHHR-H HT2 mixed bed

The linear, or mixed bed columns, contain particles with different pore sizes that are blended to provide an extended linear calibration curve. The mixed bed columns feature increasing linear calibration ranges, from TSKgel GMHHR-L, GMHHR-N, GMHHR-M, to GMHHR-H. All of the TSKgel high temperature mixed bed columns are shipped in ODCB (*o*-dichlorobenzene).

The TSKgel HHR HT2 mixed bed columns are available for ultra-high temperature analysis. Packed with PS-DVB beads, the maximum operating temperature of these columns is 220 °C.

The issue of shearing that occurs with the analysis of ultra-high molar mass polymers is overcome by the TSKgel GMH<sub>HR</sub>-M(S), TSKgel GMH<sub>HR</sub>-H(S), GMH<sub>HR</sub>-H(S) HT and GMH<sub>HR</sub>-H(S) HT2 columns. The (S) is a reference to this shearing effect.

### **Attributes and Applications:**

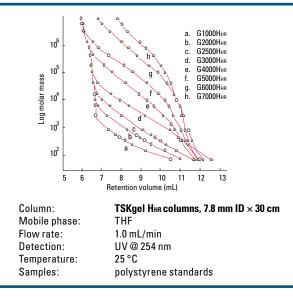
The product attributes for all of the TSKgel HHR columns is shown in Table 23. TSKgel HHR columns have a broad solvent range and are shipped in THF, except for the high temperature mixed bed columns, which are shipped in ODCB (*o*-dichlorobenzene). THF can be exchanged for a wide variety of organic solvents. See the table within the TSKgel H series column overview for a listing of these solvents. Figures 73-77 show the calibration curves for the TSKgel HHR columns.

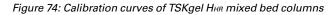
#### Table 23: Product attributes

	attributo		1	,
TSKgel column	Particle size	Pore size	Exclusion limit	Max. temp.
G1000Ннв	5 µm	1.5 nm	<1,000 Da	140 °C
G2000HHR	5 µm	2 nm	<1.0 × 10 <sup>4</sup> Da	140 °C
G2500Hнк	5 µm	3 nm	2.0 × 104 Da	140 °C
G3000Hнк	5 µm	7.5 nm	<6.0 × 104 Da	140 °C
G4000HHR	5 µm	20 nm	<4.0 × 10 <sup>5</sup> Da	140 °C
G5000Hнв	5 µm	65 nm	4.0 × 10 <sup>6</sup> Da	140 °C
G6000HHR	5 µm	>65 nm	4.0 × 10 <sup>7</sup> Da	140 °C
G7000Hнк	5 µm	>65 nm	4.0 × 10 <sup>8</sup> Da	140 °C
GMH <sub>HR</sub> -Н	5 μm, 13 μm, 20 μm, 30 μm	mixed pore sizes	4.0 × 10 <sup>8</sup> Da	80 °C
GMH <sub>HR</sub> -H(S) HT	13 µm	mixed pore sizes	4.0 × 10 <sup>8</sup> Da	140 °C
TSKgel G2000Ннв НТ	20 µm	2 nm	1 × 10 <sup>4</sup> Da	140 °C
GMH <sub>HR</sub> -H(20) HT	20 µm	mixed pore sizes	4 × 10 <sup>8</sup> Da	140 °C
GMHHR-H(30) HT	30 µm	mixed pore sizes	4 × 10 <sup>8</sup> Da	140 °C
GMHня-Н НТ	5 µm	mixed pore sizes	4 × 10 <sup>8</sup> Da	140 °C
GMH <sub>HR</sub> -L	5 µm	mixed pore sizes	4.0 × 10⁴ Da	80 °C
GMHнв-М	5 μm, 13 μm	mixed pore sizes	4.0 × 10 <sup>6</sup> Da	80 °C
GMH <sub>HR</sub> -N	5 µm	mixed pore sizes	<4.0 × 10⁵ Da	80 °C
TSKgel G2000Hhr HT2	20 µm	2 nm	1 × 10 <sup>4</sup> Da	220 °C
GMH <sub>HR</sub> -H(20) HT2	20 µm	mixed pore sizes	4 × 10 <sup>8</sup> Da	220 °C
GMH <sub>HR</sub> -H(30) HT2	30 µm	mixed pore sizes	4 × 10 <sup>8</sup> Da	220 °C
GMH <sub>HR</sub> -H(S) HT2	13 µm	mixed pore sizes	4 × 10 <sup>8</sup> Da	220 °C



Figure 73: Calibration curves of TSKgel HHR columns





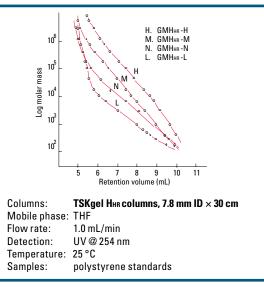


Figure 75: Calibration curves of TSKgel HHR columns

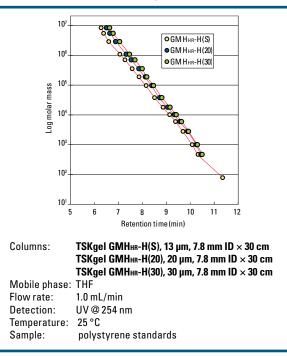
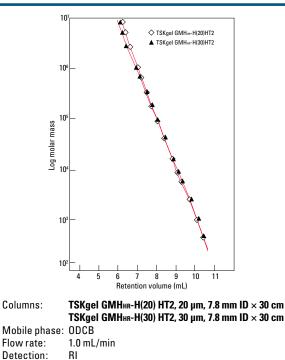


Figure 76: Calibration curves of TSKgel GMHHR-H HT2 columns

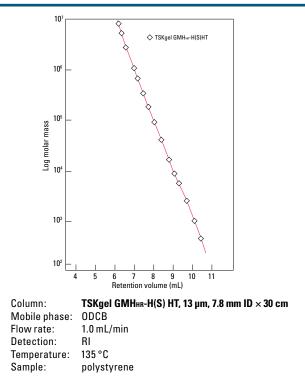


polystyrene standards

Temperature: 135 °C

Sample:

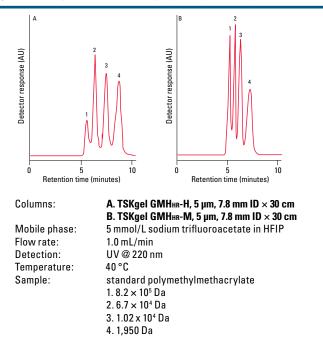
Figure 77: Calibration curve of TSKgel HHR HT column



### Polymethyl methacrylate

The effect of different pore size distributions in the mixed beds of TSKgel GMH<sub>HR</sub>-H and TSKgel GMH<sub>HR</sub>-M is illustrated in Figure 78. The TSKgel GMH<sub>HR</sub>-M produces sharper polymethyl methacrylate peaks in the  $8.0 \times 10^5$  to  $1.0 \times 10^4$  Da range.

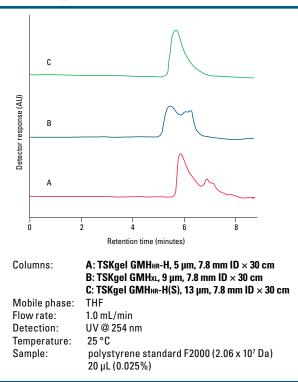
Figure 78: Comparison of standard polymethyl methacrylate mixture



### **Shear Degradation**

Shear degradation is observed especially when ultra-high molar mass compounds are analyzed. It tends to occur when analysis is carried out at high flow rates using a microparticle size packing material. Figure 79 demonstrates the relationship between shear degradation and particle size of the packing material, when TSKgel GMH columns were used. When the flow rate is 1.0 mL/min, normal elution of an ultra-high molar mass sample (2.06 × 10<sup>7</sup> Da) is only possible with the TSKgel GMH<sub>HR</sub>-H(S) column, which has a large particle size. However, with the TSKgel GMH<sub>XL</sub> and GMH<sub>HR</sub>-H columns, shear degradation does take place and new peaks appear in the chromatogram on the smaller molar mass side.

Figure 79: Shear degradation comparison

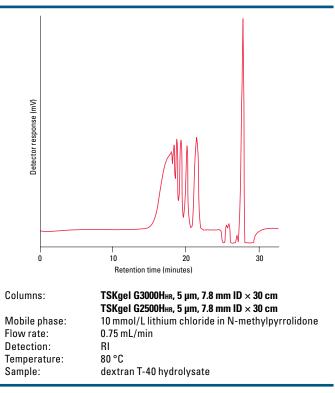




# **Dextran T-40 Hydrolysate**

The analysis of dextran T-40 hydrolysate is shown using TSKgel G3000HHR and G2000HHR columns in series in Figure 80 below.

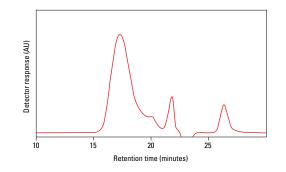
#### Figure 80: Dextran T-40 hydrolysate analysis



# **Polyphenylene Sulfide**

The analysis of PPS (polyphenylene sulfide) is shown using two TSKgel GMH\_{HR}-H(S) HT2 columns in series in Figure 81 below.

Figure 81: Polyphenylene sulfide analysis

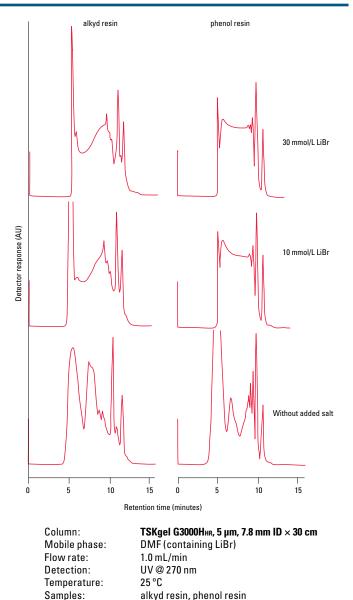


Column:TSKgel GMH<sub>HR</sub>-H(S) HT2, 13 µm, 7.8 mm ID x 30 cm x 2Mobile phase:1-chrloronapthalaneFlow rate:1.0 mL/minDetection:RITemperature:220 °CInjection vol.:300µLSample:PPS (polyphenylene sulfide), 2 g/L

### **Effect of Adding Salt**

Using the TSKgel G3000HHR column, Figure 82 shows the elution behavior of alkyd resin and phenol resin analyzed using a DMF solvent, as well as the effects of adding LiBr to DMF. With the DMF solvent, both resins eluted abnormally early from the column due to a static electric interaction. However, by adding LiBr to the DMF solvent, a normal chromatogram is obtained. Normal elution behavior of alkyd resins is possible when the LiBr concentration is about 30 mmol/L, and with phenol resins, when the concentration of the salt is around 10 mmol/L.





# About: TSKgel SuperH Size Exclusion Columns

TSKgel SuperH columns are conventional GPC columns with dimensions of 6.0 mm ID × 15 cm containing spherical particles composed of PS-DVB. The TSKgel SuperH column line consists of eight columns with different pore sizes, TSKgel SuperH1000 through TSKgel SuperH7000, and four columns with an extended linear range of the calibration curve.

TSKgel SuperH columns are high efficiency/high throughput versions of the conventional TSKgel HHR columns. Both column types are based on the same bead chemistry.

The TSKgel SuperH line consists of the following columns:

- TSKgel SuperH1000
- TSKgel SuperH2000
- TSKgel SuperH2500
- TSKgel SuperH3000
- TSKgel SuperH4000
- TSKgel SuperH5000
- TSKgel SuperH6000
- TSKgel SuperH7000
- TSKgel SuperHM-H mixed bed
- TSKgel SuperHM-L mixed bed
- TSKgel SuperHM-M mixed bed
- TSKgel SuperHM-N mixed bed

The TSKgel SuperH product line contains four linear or mixed bed columns, in which particles with different pore sizes are blended to provide an extended linear calibration curve. The mixed bed columns feature increasing linear calibration ranges, from TSKgel SuperHM-L, SuperHM-M, SuperHM-N, to SuperHM-H.

The volume of a 6 mm ID × 15 cm TSKgel SuperH column is 3.4 times smaller than that of a conventional 7.8 mm ID × 30 cm column. As a result, peak volumes will be proportionally smaller on TSKgel SuperH columns compared to a corresponding TSKgel HHR column. Thus, your HPLC system may require optimization of components that can give rise to extra-column band broadening, such as connecting tubing, injector, injection volume, detector cell volume, and detector time constant.

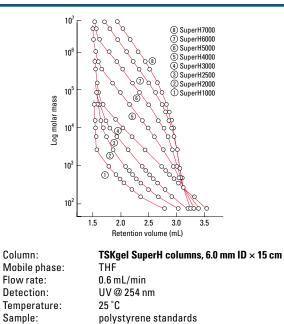
### **Attributes and Applications:**

Table 24 shows product attributes of TSKgel SuperH columns. The maximum operating temperature for TSKgel SuperH columns is 140 °C. All TSKgel SuperH columns are shipped in THF, which can be exchanged for a wide variety of organic solvents. See the table within the TSKgel H series column overview for a listing of these solvents. Figures 83 and 84 show the calibration curves for the TSKgel SuperH columns.

#### Table 24: Product attributes

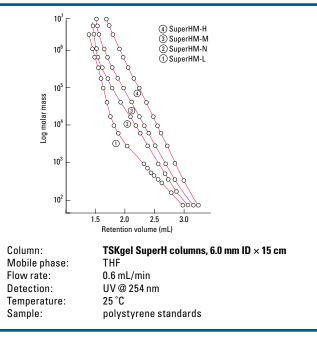
TSKgel column	Particle size (mean)	Pore size (mean)	Exclusion limit
SuperH1000	3 µm	1.5 nm	1,000 Da
SuperH2000	3 µm	2 nm	1.0 × 10 <sup>4</sup> Da
SuperH2500	3 µm	3 nm	2.0 × 104 Da
SuperH3000	3 µm	7.5 nm	6.0 × 10 <sup>4</sup> Da
SuperH4000	3 µm	20 nm	4.0 × 10⁵ Da
SuperH5000	3 µm	65 nm	4.0 × 10 <sup>6</sup> Da
SuperH6000	5 µm	>65 nm	4.0 × 10 <sup>7</sup> Da
SuperH7000	5 µm	>65 nm	4.0 × 10 <sup>8</sup> Da
SuperHM-H	3 µm	mixed pore sizes	4.0 × 10 <sup>8</sup> Da
SuperHM-L	3 µm	mixed pore sizes	4.0 × 10 <sup>6</sup> Da
SuperHM-M	3 µm	mixed pore sizes	4.0 × 10 <sup>6</sup> Da
SuperHM-N	3 µm	mixed pore sizes	4.0 × 10⁵ Da

#### Figure 83: Calibration curves for TSKgel SuperH columns

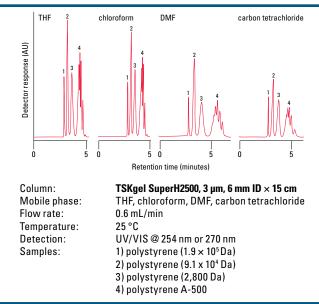


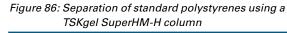


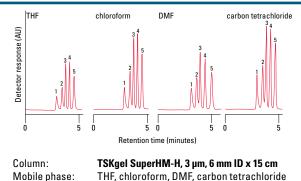
### Figure 84: Calibration curves for TSKgel SuperH mixed bed columns

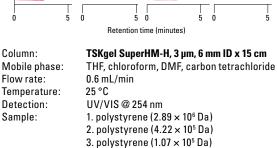


#### Figure 85: Separation of standard polystyrenes using a TSKgel SuperH2500 column









- 4. polystyrene  $(1.67 \times 10^4 \text{ Da})$
- 5. polystyrene (2,800 Da)

# **Polystyrene Mixtures**

Figure 85 compares chromatograms of standard polystyrene mixtures separated using a TSKgel SuperH2500 column with various organic solvents (THF, CHCl<sub>2</sub>, DMF, and CCl<sub>4</sub>) and Figure 86 compares chromatograms of standard polystyrene mixtures separated using a TSKgel SuperHM-H column with various organic solvents.

Due to the interaction between the packing material and standard polystyrene when using DMF as the mobile phase, the elution volume of standard polystyrenes is greater than it is with "good" solvents such as THF and CHCI,. This effect is particularly noticeable with TSKgel SuperH2500, a column for the analysis of low molar mass samples. Under these circumstances, polyethylene oxide (PEO) is recommended as the standard sample, as this reacts very little with the packing material.

### **Band Broadening in the Detector**

Table 25 compares the number of theoretical plates for a low molar mass sample (DCHP) using a TSKgel SuperH2500 column with various types of UV detectors and different flow cell volumes. Figure 87 compares the separation performance of each of these using standard polystyrene A-500 and epoxy resin samples. Based on these results, it is clear that the number of theoretical plates and the separation performance of the TSKgel SuperH column are significantly affected by band broadening in the detector, including the size of the flow cell. In analyses performed with a TSKgel SuperH column, a UV-8020 microcell (or an equivalent device) with reduced dead volume must be used in the detector.

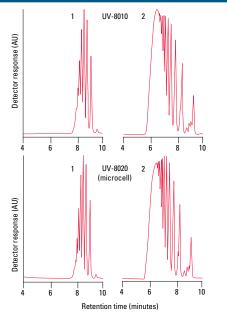
#### Table 25: Comparison of number of theoretical plates with various detectors

Number of theoretical plates (TP/15 cm)				
UV-8020*1		UV-8010*2		UV-8010*3
28,100		23,860		17,890
Column:	TSKgel SuperH2500, 3 µm, 6 mm ID × 15 cm			
Mobile phase:	THF			
Flow rate:	0.6 r	mL/min		
Temperature:	25 °(	C		
Detection:	UV@	254 nm		
Sample:	DCH	Ρ 0.1%, 2 μL		
*1 flow cell volume:	2 µL mi	crocell		

 $^{*2}$  flow cell volume: 10 µL low dead volume type

\*<sup>3</sup> flow cell volume: 10 μL

Figure 87: Dependence of separation performance on band spreading in detector



 Column:
 TSKgel SuperH2500, 3 μm, 6 mm ID × 15 cm × 2

 Mobile phase:
 THF

 Flow rate:
 0.6 mL/min

 Detection:
 UV/VIS @ 254 nm

 Temperature:
 25 °C

 Samples:
 1. standard polystyrene A-500 (0.1%), 10 μL

 2. Epikote 1004 (0.1%), 10 μL

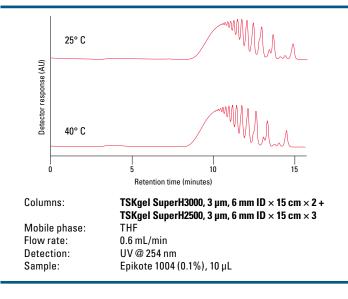
### **Column Temperature**

The following advantages are gained by conducting analysis at high temperature:

- Peaks become sharper as separation performance is increased. This is especially noticeable at higher flow rates.
- Viscosity of the mobile phase is lowered and operating pressure is decreased. This is an especially effective method with high-viscosity solvents such as DMSO, DMF, HFIP, etc.

Figure 88 demonstrates the temperature dependence of the separation of epoxy resin and a standard polystyrene mixture in TSKgel SuperH3000 and SuperH2500 columns.







# About: TSKgel SuperHZ Size Exclusion Columns

The TSKgel SuperHZ column line consists of five columns of 4.6 mm ID and 6.0 mm ID  $\times$  15 cm containing spherical particles composed of PS-DVB, TSKgel Super HZ1000 – 4000. Each column consists of a different pore size packing material. Subsequently, a unique separation range for each column exists, allowing researchers to choose a column that is designed for the sample type being analyzed.

The TSKgel SuperHZ column line also contains three linear, or mixed bed columns in which particles with different pore sizes are blended to provide an extended linear calibration curve. The mixed bed columns feature increasing linear calibration ranges, from TSKgel SuperHZM-M to SuperHZM-N to SuperHZM-H. The mixed bed columns are also available in 4.6 mm ID and 6.0 mm ID x 15 cm.

The following eight columns are available within the TSKgel SuperHZ column line:

- TSKgel SuperHZ1000
- TSKgel SuperHZ2000
- TSKgel SuperHZ2500
- TSKgel SuperHZ3000
- TSKgel SuperHZ4000
- TSKgel SuperHZM-H mixed bed
- TSKgel SuperHZM-M mixed bed
- TSKgel SuperHZM-N mixed bed

TSKgel SuperHZ column dimensions are 6 mm ID × 15 cm and 4.6 mm ID × 15 cm versus 7.8 mm ID × 30 cm for conventional GPC columns. The smaller column dimensions translate to a reduction of peak volume by a factor of 3.4 (6 mm ID) and a factor of 5.8 (4.6 mm ID) versus the same component eluting from a corresponding TSKgel HxL column. Thus, your HPLC system may require optimization of components that can give rise to extra-column band broadening, such as connecting tubing, injector, injection volume, detector cell volume, and detector time constant.

# **Attributes and Applications:**

TSKgel SuperHZ columns have been developed for high throughput, high efficiency GPC applications such as those encountered in combinatorial chemistry experiments. These columns feature ultra-low sample adsorption, i.e., the columns show true size exclusion behavior for most polymers.

TSKgel SuperHZ1000 – 4000 columns are capable of measuring monomers, polymer additives, oligomers and polymers up to a molar mass of several hundred thousand with proper selection of pore size. Ultra-fine particles (3  $\mu$ m) have been developed to provide high resolution over the entire molar mass range. This is especially important for the separation of low molar mass compounds.

Additionally, the mixed bed columns (TSKgel SuperHZM-N, M-M, and M-H) are capable of measuring oligomers and polymers with molar masses up to tens of millions with proper selection of the pore size. The various particle sizes of the mixed bed packing materials have been optimized to ensure resolution in the low molar mass range while avoiding shear degradation of polymers in the high molar mass region. The columns are shipped in THF, which can be exchanged for a limited number of organic solvents as shown in the table within the TSKgel H series column overview.

Table 26 shows the product attributes of TSKgel SuperHZ columns, while Table 27 lists the features of the TSKgel SuperHZ column line and the corresponding benefits. The calibration curves for the TSKgel SuperHZ columns are shown in Figures 89 and 90.

Table 26: Produ	ict attributes
-----------------	----------------

TSKgel column	Particle size	Pore size	Exclusion limit	Max. temp.
SuperHZ1000	3 µm	1.5 nm	1,000 Da	60 °C
SuperHZ2000	3 µm	2 nm	1.0 × 104 Da	60 °C
SuperHZ2500	3 µm	3 nm	2.0 × 104 Da	60 °C
SuperHZ3000	3 µm	7.5 nm	6.0 × 10 <sup>4</sup> Da	60 °C
SuperHZ4000	3 µm	20 nm	4.0 × 10⁵ Da	80 °C
SuperHZM-N	3 µm	mixed pore sizes	7.0 × 10⁵ Da	80 °C
SuperHZM-M	3 µm	mixed pore sizes	4.0 × 10 <sup>6</sup> Da	80 °C
SuperHZM-H	10 µm	mixed pore sizes	4.0 × 10 <sup>8</sup> Da	80 °C

Table 27: Features and benefits of TSKgel SuperHZ columns

Feature	Benefit
Ultra-fine particles used in packing material	<ul> <li>Short measurement time is achieved.</li> <li>Resolution equivalent to conventional columns (30 cm) can be obtained in ½ measurement time</li> <li>Resolution does not deteriorate even under a high flow rate.</li> </ul>
Semi-micro columns (4.6 mm ID and 6.0 mm ID)	<ul> <li>Reduction in solvent consumption (running costs, effluent processing costs) 1/6 to 1/3 solvent consumption compared to conventional columns</li> </ul>
Optimization of particle size in the packing materials	• Shear degradation in polymers with high molar mass can be prevented
Adoption of low- adsorption packing materials	• Applicable to wide range of samples

### Figure 89: Calibration curves for TSKgel SuperHZ columns

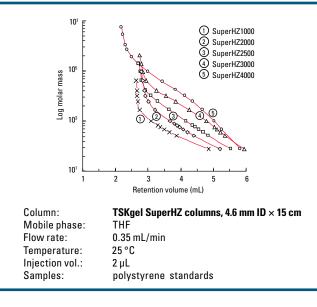
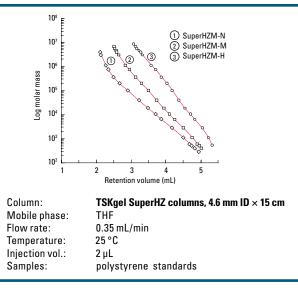
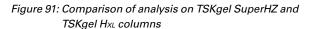


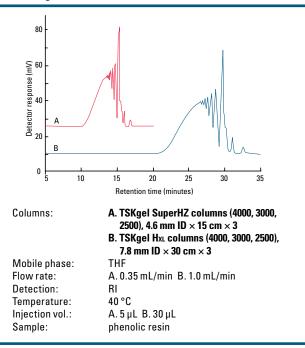
Figure 90: Calibration curves for TSKgel SuperHZ mixed bed columns



### Faster Analysis

TSKgel SuperHZ1000-SuperHZ4000 columns are packed with 3  $\mu$ m particles. The ultra-fine particles allow for high efficiency separations of low molar mass substances such as oligomers. These columns have theoretical plate values (per unit length) which are twice those of the conventional 5  $\mu$ m columns. As a result, equal resolution can be obtained within half the analysis time. An example showing the analysis of phenolic resin is demonstrated in Figure 91.

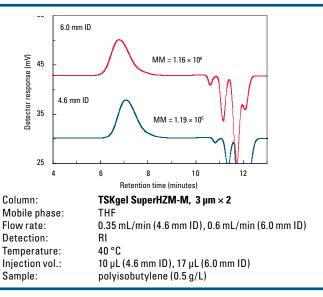




# Polyisobutylene

The chromatogram in Figure 92 shows the analysis of polyisobutylene using two TSKgel SuperHZM-M columns in series

Figure 92: Analysis of polyisobutylene





# About: TSKgel SuperMultiporeHZ Size Exclusion Columns

TSKgel SuperMultiporeHZ columns represent a new strategy for the separation of polymers with a wide range of molar masses. These columns are packed with particles of a uniform size, with each particle having a very broad pore size distribution. This innovative multi-pore approach, pioneered by Tosoh Bioscience, essentially creates a linear calibration curve within each particle. The spherical monodisperse, 3, 4 or 6  $\mu$ m particles consist of cross-linked polystyrene/divinylbenzene copolymer. This base material, coupled with the semi-micro column dimensions (4.6 mm ID × 15 cm), offers users high speed and low solvent consumption analyses with precise results. Three columns are available within the TSKgel SuperMultiporeHZ series, each with a different particle size and separation range.

The TSKgel SuperMultiporeHZ columns offered include:

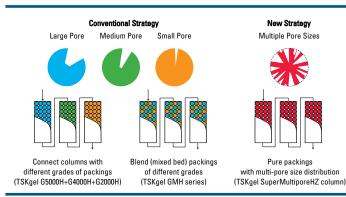
- TSKgel SuperMultiporeHZ-N
- TSKgel SuperMultiporeHZ-M
- TSKgel SuperMultiporeHZ-H

### About: Multi-pore Technology

Prior to the introduction of TSKgel SuperMultiporeHZ columns, scientists separating polymers with a wide range of molar masses were left with two options. One option was to use multiple columns of different pore sizes linked together in series. A second was to use a column packed with a mixed bed resin of different pore sizes at an optimized mix ratio. However, problems can occur with both of these methods, which include distortion of the chromatogram or deviations between the actual calibration curve and the calibration curve approximated from data obtained from the molar mass standards.

As is shown in Figure 93, a novel approach to solve this problem was developed by Tosoh scientists and is incorporated in TSKgel SuperMultiporeHZ columns. Small particles of uniform size are synthesized with a broad distribution of pore sizes. This novel approach creates a linear calibration curve within each particle. Therefore, columns with an extended linear calibration curve can now be prepared without mixing particles of different pore sizes.

Figure 93: Graphical representations illustrate the multi-pore particle synthesis technology



# **Attributes and Applications:**

Product attributes for the TSKgel SuperMultiporeHZ columns are listed in Table 28. Table 29 lists features and benefits of these columns. TSKgel SuperMultiporeHZ columns can be utilized for the analysis of polymers with a wide MM distribution range. The columns are shipped in THF, which cannot be replaced for any other organic solvent. Figure 94 shows the calibration curves for the TSKgel SuperMultiporeHZ columns.

### Table 28: Product attributes

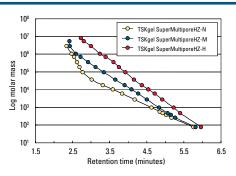
TSKgel column	SuperMultipore HZ-N	SuperMultipore HZ-M	SuperMultipore HZ-H
Base material	PS-DVB	PS-DVB	PS-DVB
Particle size	3 µm*	4 µm*	6 µm*
Pore size	8 nm	14 nm	>14 nm
Exclusion limit (PST/ THF)	1.2 × 10⁵ Da	2.0 × 10⁵ Da	4.0 × 10 <sup>7</sup> Da
Separation range	300 ~ 5.0 × 10⁴ Da	500 ~ 1.0 × 10º Da	1,000 ~ 1.0 × 10 <sup>7</sup> Da
Theoretical plates/15 cm column	20,000	16,000	11,000

\* Particle size distribution is monodisperse.

Table 29: Features and benefits

Feature	Benefit
Multi-pore packing material (wide range of pores contained in single particle)	<ul> <li>Calibration curves with superior linearity</li> <li>No observable distortion of chromatograms</li> <li>Improved accuracy and repeatability of molar mass data</li> <li>Capable of rapid analysis with high separation performance</li> </ul>
Smaller particle size (monodisperse particles)	<ul> <li>Capable of achieving the same separation performance as conventional columns (30 cm) in half the analysis time</li> <li>No reduction in separation performance even for analysis at high flow rates</li> <li>Improved robustness of column performance</li> </ul>
Semi-micro column	<ul> <li>Reduced solvent consumption</li> <li>1/6th the consumption of conventional (30 cm) columns</li> </ul>
Low adsorption packing material	• Can be used for a wide variety of samples

Figure 94: Calibration curves for TSKgel SuperMultiporeHZ columns

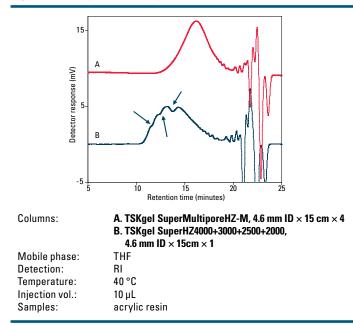


Columns:	TSKgel SuperMultiporeHZ-N, 3 μm, 4.6 mm ID ×15 cm TSKgel SuperMultiporeHZ-M, 4 μm, 4.6 mm ID × 15 cm TSKgel SuperMultiporeHZ-H, 6 μm, 4.6 mm ID × 15 cm
Mobile phase:	THF
Flow rate:	0.35 mL/min
Detection:	UV @ 254 nm
Temperature:	25 °C
Samples:	PStQuick polystyrene standards

# **Acrylic Resin**

Figure 95 demonstrates that inflection points are no longer observed with columns packed from particles prepared by multi-pore technology.

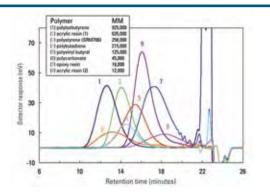
Figure 95: Comparison for separation of acrylic resin



### **Various Polymers**

Various polymers were analyzed on four TSKgel SuperMultiporeHZ-M columns in series. The superimposed chromatograms in Figure 96 clearly demonstrate that these new GPC columns can be utilized for the analysis of polymers with a wide molar mass distribution range.

Figure 96: Separation of various polymers



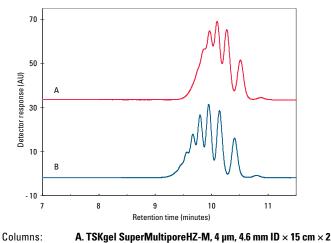
Columns:SuperMultiporeHZ-M, 4 μm, 4.6 mm ID × 15 cm × 4Mobile phase:THFFlow rate:0.35 mL/minDetection:RITemperature:25 °CInjection vol.:10 μLSample conc.:0.3%



# **Standard Polystyrene**

Figure 97 compares separation on the TSKgel SuperMultiporeHZ-N column versus the TSKgel SuperMultiporeHZ-M column in the low molar mass region (standard polystyrene A-500). The calibration curve for the TSKgel SuperMultiporeHZ-N column is not as steep and better separation is provided in the low molar mass region due to the smaller particle size (higher number of theoretical plates) of the TSKgel SuperMultiporeHZ-N column.

### Figure 97: Analysis of standard polystyrene



 B. TSKgel SuperMultiporeHZ-N, 3 μm, 4.6 mm ID × 15 cm × 2

 B. TSKgel SuperMultiporeHZ-N, 3 μm, 4.6 mm ID × 15 cm × 2

 Mobile phase:
 THF

 Flow rate:
 0.35 mL/min

 Detection:
 UV @ 254 nm

 Temperature:
 25 °C

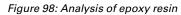
 Injection vol.:
 5 μL

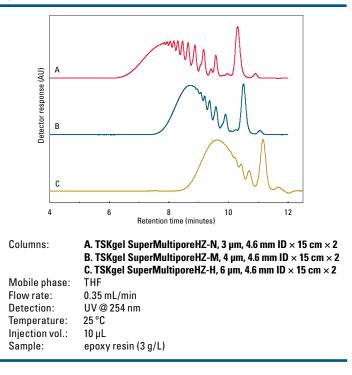
 Sample:
 standard polystyrene oligomer

 (TSKgel standard polystyrene A-500) (5 g/L)

### **Epoxy Resin**

Figure 98 is a chromatogram of an epoxy resin (approximately 6,000 Da) created using the TSKgel SuperMultiporeHZ columns. The best separation performance is shown by the TSKgel SuperMultiporeHZ-N column, the particle size used for low molar mass samples, and it is clear that the TSKgel SuperMultiporeHZ-H column does not provide adequate separation performance.

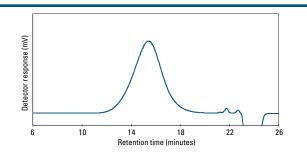




# Polyvinylbutyral

The analysis of polyvinylbutyral using a TSKgel SuperMultiporeHZ-H column is show in Figure 99. A smooth chromatogram without any distortion is obtained.

Figure 99: Analysis of polyvinylbutyral



Column:TSKgel SuperMultiporeHZ-H, 6 μm, 4.6 mm ID × 15 cm × 4Mobile phase:THFFlow rate:0.35 mL/minDetection:RITemperature:40 °CInjection vol.:10 μLSample:polyvinylbutyral (3 g/L)



