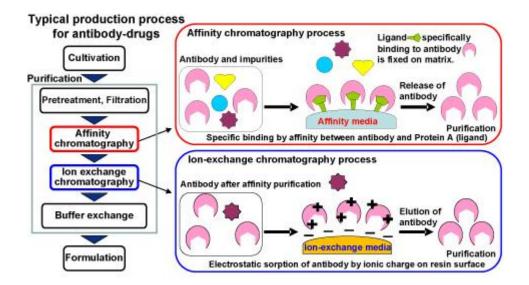


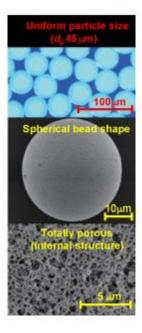


# Chromatographic resins

# MabSpeed & ChromSpeed Series

MabSpeed<sup>™</sup> and ChromSpeed<sup>™</sup> chromatographic resins are specially designed material for the purification of biopharmaceuticals (antibody-drugs, protein-drugs, etc.). They are based on hydrophilic polymethacrylate matrix with spherical, totally porous structure. The rigid, uniform-particle sized matrix enables high speed chromatographic operation with high production efficiency. Various product line can meet variety of demands for separation and purification.





# MabSpeed RP Series

## Spherical and monodisperse particles:

- Easy to pack
- · High packing reproducibility
- Low pressure drop

## Extremely high bed height available: > 50 cm

- Compatible with simple structured columns
- Enables equipment cost reduction

#### High flow rate available: > 600 cm/h

- Improves throughput
- Shortens process time suitable for higher efficacy of purified therapeutic mAb

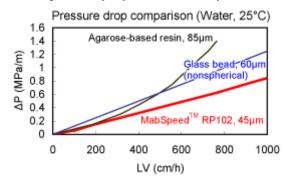
#### Rigid and durable matrix:

Non compressive bed

#### Variation in particle size:

Standard particle size: 45µm

#### Hydraulic properties of MabSpeed™



MCC 45µm data was calculated from the data measured for 60µm.

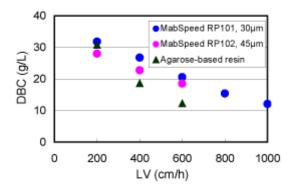
#### Cleaning in place - Low effect on Ligand leakage and Dynamic binding capacity

Ligand leakage		
cycles	ng PrA / mg IgG	
1	0.53	
20	0.23	
40	0.24	
60	0.25	
80	0.26	
100	0.25	

#### Characteristics of MabSpeed™

Product	MabSpeed™ RP102
Matrix	Crosslinked polymethacrylate
Ligand	recombinant Protein A
IgG-SBC	> 30 g/L
Particle size	45μm

#### Dynamic binding capacity comparison

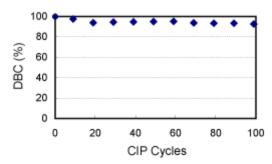


#### Conditions:

Column, 127 x 5 mml.D.;

Sample, 1.0 mg/mL human γ-globulin; Buffer, PBS (pH 7.4); Temp., 20°C.

The DBC was determined at 10% breakthrough.



#### Conditions:

Column, 50 x 5 mml.D.;
DBC of human γ-globulin was determined at 10% breakthrough.
Binding, Phosphate buffered saline (pH7.4);
Elution, 0.1M Sodium citrate (pH3.0);
CIP, 0.1M NaOH; Contact time, 15 min.

# **ChromSpeed S Series**

## Superior IgG binding capacity: > 100 g/L

- Suitable for therapeutic mAb production
- Reduces packing material cost

#### Extremely high bed height available: > 50 cm

- Compatible with simple structured columns
- Enables equipment cost reduction

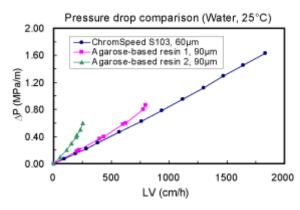
#### High flow rate available: > 600 cm/h

- Improves throughput
- Shorter process time suitable for higher efficacy of purified therapeutic mAb

#### High durability:

- Rigid spherical macroporous polymer
- · Chemically and mechanically stable

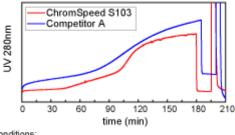
## Hydraulic properties of ChromSpeed™ S



## Characteristics of ChromSpeed™ S

Product	ChromSpeed™ S103
Matrix	Crosslinked polymethacrylate
Functionality	-SO <sub>3</sub>
Ion exchange capacity	> 0.05 eq/L
IgG-DBC	> 100 g/L
Particle size	60µm

IgG binding profile comparison (Mouse polyclonal IgG, pH: 5.2)



Conditions:

Column, 30 x 6.4 mm I.D.;

Adsorption, mouse polyclonal IgG, 1mg/mL in 20mM Na citrate + 15mM NaCl (pH5.2)

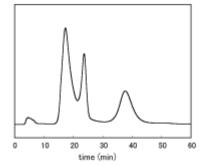
Washing, 20mM Na citrate (pH5.2)

Desorption, 20mM Na citrate + 1M NaCl (pH5.2)

Flow rate, 1.0 mL/min; Residence time, 1.0 min; Detector, UV 280 nm.

Example of protein separation on ChromSpeed™ S packed in a 50cm height column - comparison with agarose-based resin packed in a 12.5cm height column

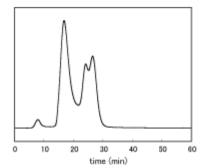
ChromSpeed S103 (pH6.0, 0-100%B over 30min) Hemoglobin/Cytochrome c/Lysozyme = 25/6.25/6.25mg/2.5mL Agarose-based resin (pH6.0, 0-100%B over 30min) Hemoglobin/Cytochrome c/Lysozyme = 25/6.25/6.25mg/2.5mL



#### Conditions:

Column, 500 x 8mml.D., 25mL Eluent A, 20mM Na phosphate (pH6.0); Eluent B, 20mM Na phosphate + 1M NaCl (pH6.0) Gradient, 0-100%B; Flow rate, 2.5mL/min;

SV=6; LV=300cm/h Detector, UV 280nm.



#### Conditions:

Column, 125 x 16mml.D., 25mL Eluent A, 20mM Na phosphate (pH6.0); Eluent B, 20mM Na phosphate + 1M NaCl (pH6.0) Gradient, 0-100%B; Flow rate, 2.5mL/min; SV=6; LV=75cm/h

Detector, UV 280nm.

# **ChromSpeed Q Series**

#### Superior IgG binding capacity: > 100 g/L

- Suitable for therapeutic mAb production
- Reduces packing material cost

#### Extremely high bed height available: > 50 cm

- Compatible with simple structured columns
- Enables equipment cost reduction

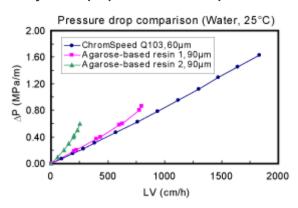
#### High flow rate available: > 600 cm/h

- Improves throughput
- Shorter process time suitable for higher efficacy of purified therapeutic mAb

#### High durability:

- · Rigid spherical macroporous polymer
- · Chemically and mechanically stable

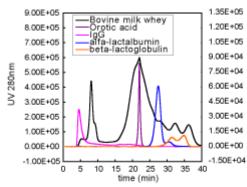
#### Hydraulic properties of ChromSpeed™ Q



#### Characteristics of ChromSpeed™ Q

Product	ChromSpeed™ Q103
Matrix	Crosslinked polymethacrylate
Functionality	-N(CH <sub>3</sub> ) <sub>3</sub> +
lon exchange capacity	> 0.05 eq/L
IgG-DBC	> 100 g/L
Particle size	60µm

# IgG binding profile comparison (Mouse polyclonal IgG, pH: 5.2)



Conditions: Column, 500 x 8mml.D. (ChromSpeed-Q103, 60µm);

Eluent A, 20mM sodium phosphate (pH7.0);

Eluent B, 20mM sodium phosphate + 1.0M NaCl (pH7.0);

Flow rate, 2.5ml/min (300cm/h);

Gradient, 0-30% B over 30min

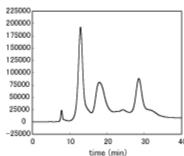
Orotic acid,40  $\mu g; IgG,\alpha\text{-lactalbumin}$  and  $\beta\text{-lactoglobulin},$ 

400µg; Whey, 400µl.

## Example of protein separation on ChromSpeed™ Q packed in a 50cm height column

#### - comparison with agarose-based resin packed in a 12.5cm height column

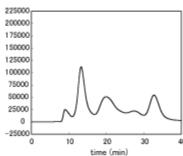
ChromSpeed<sup>™</sup>Q103 (pH8.0, 0-50%B over 30min) Myoglobin/Conalbumin/Trypsin inhibitor = 12.5/12.5/25mg/2.5mL



#### Conditions:

Column, 500 x 8mml.D., 25mL Eluent A, 20mM Tris-HCI (pH8.0); Eluent B, Tris-HCI + 1M NaCI (pH8.0) Gradient, 0-50%B; Flow rate, 2.5mL/min; SV=6; LV=300cm/h Detector, UV 280nm.

Agarose-based resin (pH8.0, 0-50%B over 30min) Myoglobin/Conalbumin/Trypsin inhibitor = 12.5/12.5/25mg/2.5mL



#### Conditions:

Column, 125 x 16mml.D., 25mL Eluent A, 20mM Tris-HCl (pH8.0); Eluent B, Tris-HCI + 1M NaCl (pH8.0) Gradient, 0-50%B; Flow rate, 2.5mL/min; SV=6; LV=75cm/h Detector, UV 280nm.