

## **Pall Chromatography Media**



## Versatile offering for purification of biomolecules

- Size Exclusion (Gel Filtration)
- Ion Exchange
- Affinity Purification
- Mixed-Mode and **Hydrophobic Charge Induction Chromatography**
- Hydroxyapatite

Chromatography continues to be an essential technology for the purification of biomolecules. Pall offers an extensive portfolio of media for affinity, ion exchange, size exclusion, hydrophobic induction, and hydroxyaptite chromatography. We offer products to cover research needs, scale-up, and polishing. Depending on your specific application needs, Pall offers chromatography solutions in resin and/or membrane form. We offer flat sheet membranes, bulk resins, pre-packed columns, and media incorporated

Our chromatography medias can be used for purification of biomolecules (e.g., nucleic acids, proteins) and compounds on the research and process scale. Our resins are useful in proteomic sample preparation applications, as well as in laboratory bioprocessing development and scale-up work. For the full listing of chromatography technologies offered by Pall, please refer to the Pall Laboratory Filtration, Separation and Detection Products Catalog or visit www.pall.com/lab.

		into specific product housings.  Catalog or visit www.pall.com/lab.  Chromatography Product Chart								
	N.									
romatography pe	Product	Part Number	Size	Description	Particle Size	Capacity	Stability Range (pH)	Cleaning pH	Pressure Stability	Applications
ze Exclusion del Filtration) deparation by olecule Size ulk Resin	Ultrogel® AcA 34 Ultrogel AcA 44	23015-025 23015-019 23022-024 23022-015	100 mL Bottle 1000 mL Bottle 100 mL Bottle 1000 mL Bottle	Ultrogel AcA are polymeric sorbents for size exclusion composed of polyacrylamide and agarose, characterized by narrow particle	60-140 µm	NA 	3-10	3-10		Exclusion Limit   Fractionation Range
	Ultrogel AcA 54 Ultrogel AcA 202	23019-023 23019-011 24892-022 24892-010	100 mL Bottle 1000 mL Bottle 100 mL Bottle 1000 mL Bottle	size distribution and narrow pore size distribution. AcA 202 can be used for desalting.	Y	Desalting capacity 45% gel volume	*	<b>Y</b>		90 KD 5-70 KD 22 KD 1-15 KD
	Trisacryl® GF05 M  Trisacryl GF2000 LS	25914-060 25914-037 26065-045 26065-011	100 mL Bottle 1000 mL Bottle 100 mL Bottle 1000 mL Bottle	Trisacryl GF are highly hydrophilic copolymers designed for medium pressure gel filtration. GF05 can be used for desalting.	40-80 μm 80-160 μm	Desalting capacity 33% gel volume NA	1-11	1-11	Up to 3 bar (44 psi)	Exclusion Limit Fractionation Range 3 KD 200-2500 D Affinity chromatography, purification of macromolecules
n Exchange apparation to Charge ulk Resin	Q Ceramic HyperD® 20	20040-051 20040-044 20040-036 20040-028 20040-010	5 mL Bottle 25 mL Bottle 100 mL Bottle 500 mL Bottle 1000 mL Bottle	Strong anion exchanger. Ceramic HyperD 20 ion exchangers employ a high capacity hydrogel polymerized within the large pores of a rigid ceramic bead. High resolution of biomolecules from infinite variety of feedstocks.	20 µm (avg)	> 85 mg/mL¹	2-12	1-14	200 bar (3,000 psi)	Polypeptide and plasmid purification
	S Ceramic HyperD 20	20038-055 20038-048 20038-030 20038-022 20038-014	5 mL Bottle 25 mL Bottle 100 mL Bottle 500 mL Bottle 1000 mL Bottle	Strong cation exchanger. Ceramic HyperD 20 ion exchangers employ a high capacity hydrogel polymerized within the large pores of a rigid ceramic bead.	20 μm (avg)	> 85 mg/mL <sup>2</sup>	2-12	1-14	200 bar (3,000 psi)	Polypeptide purification
	Q Ceramic HyperD F	20066-098 20066-031 20066-023 20066-015	5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	Strong anion exchanger. Ceramic HyperD F ion exchangers employ a high capacity hydrogel polymerized within the large pores of a rigid ceramic bead.	50 μm (avg)	> 85 mg/mL¹	2-12	1-14	70 bar (1,000 psi)	Protein concentration, protein separation, contaminant removal
	S Ceramic HyperD F	20062-089 20062-030 20062-022 20062-014	5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	Strong cation exchanger. Ceramic HyperD F ion exchangers employ a high capacity hydrogel polymerized within the large pores of a rigid ceramic bead.	50 μm (avg)	> 75 mg/mL²	2-12	1-14	70 bar (1,000 psi)	Protein concentration, protein separation, contaminant removal
	DEAE Ceramic HyperD F	20067-070 20067-039 20067-021 20067-013	5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	Weak anion exchanger. Ceramic HyperD F ion exchangers employ a high capacity hydrogel polymerized within the large pores of a rigid ceramic bead.	50 μm (avg)	> 85 mg/mL¹	2-12	1-14	70 bar (1,000 psi)	Protein concentration, protein separation, contaminant removal
	CM Ceramic HyperD F	20050-084 20050-035 20050-027 20050-019	5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	Weak cation exchanger. Ceramic HyperD F ion exchangers employ a high capacity hydrogel polymerized within the large pores of a rigid ceramic bead.	50 μm (avg)	> 60 mg/mL <sup>3</sup>	2-12	1-14	70 bar (1,000 psi)	Protein concentration, protein separation, contaminant removal
roSep™ lumns	CM Ceramic HyperD F S Ceramic HyperD F Q Ceramic HyperD F DEAE Ceramic HyperD F IEX Variety Pack	20050-C001 20062-C001 20066-C001 20067-C001 IEXVP-C001	1 mL Column	Columns pre-packed with patented HyperD "gel-in-a-shell" resins for chromatographic separations. Offers extremely high dynamic binding capacities at fast flow rates.	50 μm (avg)	> 60 mg/mL <sup>3</sup> > 75 mg/mL <sup>2</sup> > 85 mg/mL <sup>1</sup> > 85 mg/mL <sup>1</sup> See above	2-12	1-14	3 bar (44 psi)	Protein concentration, protein separation, contaminant removal
vices	AcroPrep <sup>™</sup> 96 with Mustang® Q, 350 µL	5047	10/pkg	Strong anion exchange membrane in a 96-well filter plate configuration.	· ·		Y	'	· ·	Protein concentration, protein separation, contaminant removal
	AcroPrep 96 with Mustang Q, 1 mL AcroPrep 96 with	5062 5048	5/pkg 10/pkg	Strong cation exchange membrane in a 96-well filter plate configuration.						Contaminant removal
	Mustang S, 350 µL AcroPrep 96 with Mustang S, 1 mL Acrodisc® Mustang Q Chromatography Unit Acrodisc Mustang S Chromatography Unit	5063 MSTG25Q6 MSTG25S6	5/pkg 10/pkg 10/pkg	Strong anion exchange membrane in a syringe filter configuration. Strong cation exchange membrane in a syringe filter configuration.		DNA: 3.6 mg/Acrodisc un RSA: 10 mg/Acrodisc unit Lysozyme: 8 mg/ Acrodisc unit hu lgG: 11 mg/				Protein concentration, protein separation, contaminant removal
inity rification paration Using ecific Ligands IK Resin	Blue Trisacryl M	25896-C001 25896-051 25896-045 25896-010 25896-028	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	Blue Trisacryl M is an affinity chromatographic sorbent. The basic matrix is Trisacryl GF2000, a macroporous non-ionic sorbent on which Cibacron* blue is covalently	40-80 μm	Acrodisc unit  HSA: 10-15 mg/mL <sup>4</sup> ; BSA: 5-7 mg/mL <sup>4</sup>	1-10	1-10	3 bar (44 psi)	Albumin depletion, purification of enzymes and proteins, such as kinases, interferons, and some coagulation factors
in recom	Protein A Ceramic HyperD F	20078-C001 20078-036 20078-028 20078-010 20078-044	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	immobilized.  Protein A Ceramic HyperD F is an affinity sorbent prepared using a rigid proprietary ceramic bead. Recombinant Protein A is immobilized to a specially formulated hydrogel within the porous	V 50 μm (avg)	> 30 mg/mL <sup>s</sup>	2-11	2-13	3 bar (44 psi) 70 bar (1,000 psi)	lgG purification/depletion
	Heparin HyperD M	20029-062 20029-039 20029-021 20029-013	5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	ceramic bead.  Heparin HyperD M is composed of a porous rigid mineral bead containing heparin (porcine) bound hydrogel filled pores.	80 µm (avg)	> 25 mg/mL <sup>6</sup>	3-13	3-13	70 bar (1,000 psi)	Purification of coagulation factors, lipoproteins, growth hormones, growth factors, nucleic acid binding enzymes
	IMAC HyperCel™	20029-013 20093-C001 20093-069 20093-010 20093-028	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle	IMAC HyperCel uses tridentate IDA as a chelating agent. This ligand is immobilized on the HyperCel base sorbent.	80-100 μm	Metal ion capacity: 40-70 μmol Cu <sup>++</sup> /mL Ionic capacity: 90-140 μeg/mL	<u> </u>	<u> </u>	3 bar (44 psi)	Purification of HIS-tagged proteins, antibodies, and prefractionation of complex protein mixtures
	Lysine HyperD	20059-058 20059-036 20059-028 20059-010	5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	Lysine HyperD is comprised of a porous rigid mineral bead containing lysine (L-lysine) bound hydrogel filled pores.	70 µm (avg)	о по рофине	3-13	3-13	70 bar (1,000 psi)	Purification of glycoproteins and other biological molecules that bind to Lysine
ts	Enchant <sup>™</sup> Albumin Depletion Kit	5300-ALBDEP	25 purifications	Convenient spin column format using Cibacron blue dye. Includes all buffer and devices needed for 25 purifications.		> 2 mg albumin				Albumin depletion from serum or plasma
	Enchant Multi-protein Affinity Separation Kit	5300-AFFMPS	24 purifications	Includes all of the components necessary to fractionate albumin and IgG from human serum or plasma. The HSA and IgG fractionation resins utilize unique ligands coupled to a solid support for highly specific removal of target protein(s).		> 97% removal of human albumin and IgG from 50 µL of serum/plasma diluted 1:5				Facilitates biomarker discovery by removing > 80% of the total protein content (sample complexity reduction)
xed Mode & drophobic luction (HCIC) lk Resin	MEP HyperCel	12035-C001 12035-069 12035-010 12035-028 12035-036	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	MEP HyperCel (4-mercapto-ethyl- pyridine) sorbent is a high capacity, high selectivity sorbent. In contrast to protein A sorbents, IgG binding on MEP HyperCel is essentially independent of subclass or species. "Weakly binding" variants (e.g., murine IgG, rat IgG) are well retained.	80-100 µm	> 20 mg/mL <sup>7</sup>	3-12 Adsorption: 7.0-9.0 Elution: 4.0-5.8	3-14	< 3 bar (44 psi)	Purification/depletion of polyclonal and monoclonal antibodies from most species
	SDR HyperD	20033-C001 20033-065 20033-031 20033-023 20033-015	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	SDR HyperD is a "mixed mode" of size exclusion, normal phase, and reversed phase. SDR HyperD is a composite sorbent that combines a silica bead moiety filled with long chain aliphatic polymers that are cross-linked to provide a 3D mesh.	40-100 μm	60-80 mg/mL <sup>s</sup>	2-12	2-12	3 bar (44 psi) 70 bar (1,000 psi)	Solvent and detergent removal while recovering NATIVE protein
	HEA HyperCel	20250-C001 20250-012 20250-026 20250-033 20250-041	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	'Mixed-mode' chromatography based on a combination of electrostatic and hydrophobic properties of the protein and ligands. Has an aliphatic (n-hexylamine) ligand which has lower hydrophobicity than PPA.	80-100 µm (avg)	40-60 mg/mL <sup>9</sup>	By gradient or step-elution, e.g. pH 7.0-2.6 adsorption 7.0-9.0	1-14	< 3 bar (44 psi)	Unique industry-scalable sorbents designed for protein capture and impurity removal in a biopharmaceutical environment. Provides different selectivites not accessible with traditional ion exchange or HIC.
	PPA HyperCel	20260-C001 20260-015 20260-025 20260-030 20260-040	1 mL Column 5 mL Bottle 25 mL Bottle 100 mL Bottle 1000 mL Bottle	'Mixed-mode' chromatography based on a combination of electrostatic and hydrophobic properties of the protein and ligands. Has an aromatic (phenylpropylamine) ligand	80-100 μm (avg)	40-60 mg/mL <sup>9</sup>	By gradient or step-elution, e.g. pH 7.0-2.6 adsorption 7.0-9.0	1-14	< 3 bar (44 psi)	Unique industry-scalable sorbents designed for protein capture and impurity removal in a biopharmaceutical environment. Provides different selectivites not accessible with traditional ion exchange or HIC.

(1) Dynamic binding capacity, 10% breakthrough, 200 cm/h; sample: 5 mg/mL BSA in 50 mM Tris-HCl buffer, pH 8.6.  $(2) \ Dynamic \ binding \ capacity, \ 10\% \ breakthrough, \ 200 \ cm/h; \ sample: 5 \ mg/mL \ lysozyme \ in \ 50 \ mM \ sodium \ acetate, \ pH \ 4.5.$  $(3) \ Dynamic \ binding \ capacity, \ 10\% \ breakthrough, \ 200 \ cm/h; \ sample: 5 \ mg/mL \ hu \ lgG \ in \ 50 \ mM \ sodium \ acetate, \ 100 \ mM \ NaCl, \ pH \ 4.7.$ 

HA Ultrogel hydroxyapatite sorbent is

composed of cross-linked agarose

60-180 μm

Cytochrome C:

> 7 mg/mL;<sup>10</sup>

BSA: < 7 mg/mL<sup>11</sup>

which is highly hydrophobic.

beads with micro-crystals of

agarose mesh.

hydroxyapatite entrapped in the

24775-075

24775-082

24775-025

24775-041

HA Ultrogel

5 mL Bottle

HCl, 2 M NaCl, pH 4.7, 10 cm bed height.

25 mL Bottle

100 mL Bottle

1000 mL Bottle

(4) Capacity determined in PBS buffer using 5 mg/mL. (5) Dynamic binding capacity, 10% breakthrough, 100 cm/h, determined using 10 mg/mL hu lgG in PBS, pH 7.4; elution in 0.1 M sodium citrate, pH 2.5; column 4.6 ID x 100 mm. (6) Dynamic binding capacity at 600 cm/h, using hu ATIII at 72.5 Ul/mL in 20 mM Tris-HCl, 0.3 M NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-HCl, 0.3 m NaCl, pH 7.4, elution with 20 mM Tris-H

(7) Dynamic binding capacity, 10% breakthrough; determined using 5 mg/mL hu lgG in PBS, flow rate: 60 cm/h (8) Dynamic binding capacity, 10% breakthrough at 300 cm/h, determined using 5 mg/mL Triton x 100 in PBS pH 7.4.

(9) Dynamic binding capacity, 10% breakthrough; 100 cm/h, 5 mg/mL BSA in PBS.  $(10)\ Determined\ using\ 5\ mg/mL\ Cytochrome\ c\ diluted\ 50/50\ in\ 10\ mM\ sodium\ phosphate\ buffer,\ pH\ 6.8,\ at\ 30\ cm/h.$ (11) Determined using 1 mg/mL BSA diluted 50/50 in 10 mM sodium phosphate buffer, pH 6.8, at 12.5 cm/h.

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Fractionation, purification of biomolecules

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