

HiScreen Capto Blue

Product Information

Cat#No# Hi-065P

Maximum operating pressure

3 bar [0.3 MPa] (44 psi)

Matrix

Cross-linked agarose, 6%, spherical

Average particle size

~ 75 µm

Ligand density

11 to 16 µmol/ml Cibacron Blue/mL resin

Dynamic binding capacity

Approx. 25 mg human serum albumin/ml medium

Recommended flow rate

1.2 ml/min - 2.3 ml/min

Recommended column height

100 mm

Chemical stability

Stable to commonly used aqueous buffers, 0.01 M NaOH 8 M urea, 6 M guanidine hydrochloride, 30% isopropanol, and 70% ethanol.

pH working range

3 to 13

CIP stability

2 to 13.5

Storage

HiScreen Capto Blue

2°C to 8°C

Elution buffer

50 mM KH₂PO₄ , 1.5 M KCl, pH 7.0 or 20 mM sodium phosphate, 2 M NaCl, pH 7.0.

Cleaning-in-place

Precipitated proteins: 1. Wash the column with 4 column volumes (CV) of either 0.5 M (HiScreen Capto Blue) or 0.1 M NaOH (HiScreen Blue FF) at 40 cm/h. 2. Wash with 3 to 4 CV of 70% ethanol or 2 M potassium thiocyanate. 3. Wash immediately with at least 5 CV filtered start buffer, pH 8.0. or 1. Wash the column with 2 CV of 6 M guanidine hydrochloride. 2. Wash immediately with at least 5 CV filtered start buffer, pH 8.0.

Strongly bound hydrophobic proteins, lipoproteins, and lipids: 1. Wash the column with 3 to 4 CV of up to 70% ethanol or 30% isopropanol. 2. Wash immediately with at least 5 CV filtered start buffer, pH 8.0. or 1. Wash with 2 CV detergent in a basic or acidic solution, e.g., 0.1% non-ionic detergent in 1 M acetic acid. Wash at a flow rate of 40 cm/h. 2. Remove residual detergent by washing with 5 CV of 70% ethanol. 3. Wash immediately with at least 5 CV filtered start buffer, pH 8.0.

Purification procedures

1. Remove the stoppers and connect the column to the system. Avoid introducing air into the column.
 2. Equilibrate with at least 5 column volumes (CV) start buffer.
 3. Adjust the sample to the chosen starting conditions and load on the column.
 4. Wash with 5 to 10 CV start buffer until the UV trace of the effluent returns to near baseline.
 5. Elute either by linear gradient elution or a step elution at recommended flow rates.
 6. Re-equilibrate the column with 5 to 10 CV start buffer or until the UV baseline, eluent pH, and conductivity reach the required values.
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Maximum flow velocity

4.7 mL/min

Dimensions

7.7 × 100 mm

Column volume

4.7 ml



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Column i.d.

7.7 mm

Column hardware pressure limit

0.8 MPa (8 bar, 116 psi)
