ReliChrom™ BU400/SS

Lot. E905P125

Column dimensions:

Internal Diameter i.d.	**0.8 cm**
Lenght	**10 cm**
Area	**0.5 cm^2**
Resin volume	**5 ml**
Theoretical plates N	**1798 m^{-1}**
Asymmetry A_s	**0.81**

Experimental conditions

Sample	100 µl 1% Acetone (v/v)
Mobile phase	50 mM TRIS/HCl, 0.9% NaCl, pH 8.0
Flow velocity	1.25 ml/min

Instructions for use

Preliminary set up:

- Rinse the chromatographic system circuit with DI water;
- After the removal of the upper stopper of the ReliChrom™ column, connect it to the chromatographic unit;
- Remove the bottom stopper of ReliChrom™ column and connect the column outlet to the specific device of your chromatographic system (Detectors, fraction collector…).

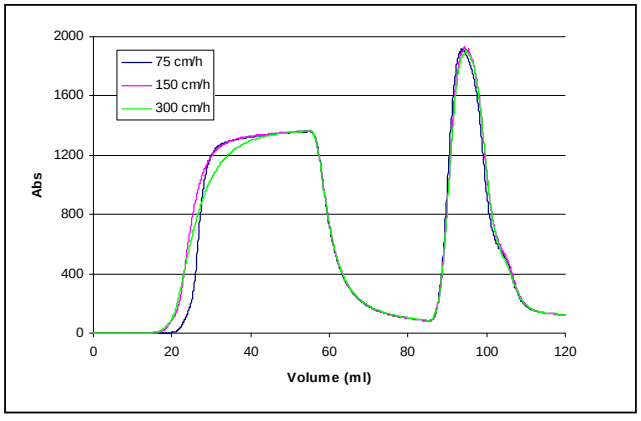
Operation mode:

- wash out the conditioning solution with 10 BV of DI water;
- start the equilibration with the desired buffer solution at an appropriate linear flow rate;
- run the chromatographic separation according to your individual protocol at the same flow rate as in the previous step;
- if necessary, perform a regeneration step following the instructions here below:

  - Wash the resin with 2 BV of DI water
  - Clean the resin with 1 BV of NaOH 0.5 M in DI water or in 10-40% alcohol solution
  - Displace NaOH solution with 2 BV of DI water
  - Rinse with 5 BV of DI water

BSA capacity vs linear velocity

Feed solution: 10 g/l BSA in 20 mM phosphate buffer pH 7 + $(NH_4)_2SO_4$ 2M
Buffer equilibration: 6 BV of 20 mM phosphate buffer pH 7 + $(NH_4)_2SO_4$ 2M
BSA loading: 10 BV
Displacement: 6 BV of 20 mM phosphate buffer pH 7 + $(NH_4)_2SO_4$ 2M
Elution: 4 BV of 20 mM phosphate buffer pH 7



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