BioVision

rev. 06/12

Heparin-Sepharose Column

CATALOG #:	6554-1 6554-5	1 ml 5 ml
LOT #:		
PREPARATION:	Heparin-Sepharose is heparin to epoxy-activa The coupling was opti and could be greater (such as thrombin) per	s prepared by covalently coupling ated 6% cross-linked sepharose beads. imized to give a high binding capacity than 0.4 mg of heparin-binding protein ml of wet gel.
CONTENTS:	Ready-to-use pre-pac volume in 20 % Etha Sepharose beads.	ked columns of 1 ml or 5 ml bead anol/dH ₂ O. > 2.5 mg heparin per ml
FEATURES:	Heparin-beads have be various heparin-bindi antithrombin III, lij (transcription factors, ligands. BioVison's He purpose of purification It can also be used medium. Specific prote concentrations of salt Sepharose formulation high flow rate, no sign pH stability range of 2-	een widely used in affinity purification of ing proteins or ligand, such as poprotein, DNA binding proteins virus coat proteins etc.) and other eparin-Sepharose is designed for the of these kinds of proteins and ligands. as a high capacity cation exchange eins can be separated by using different t or a salt gradient. This Heparin- n exhibits excellent binding capacity, ificant loss of the heparin ligand and a 10.
APPLICATIONS:	Purification of heparir ligands.	n-binding proteins, enzymes or other
STORAGE:	Store at 4°C. Do not fr year.	eeze. Stable, as supplied, for at least 1

PROCEDURE EXAMPLE:

- 1. Equilibrate the column to room temperature.
- 2. Remove the upper (first) then lower cap and allow the preservative to drain by gravity flow.
- 3. Equilibrate the column with 5 10 bed volumes of degassed Binding Buffer.
- 4. Add sample in Binding Buffer and recycle through column 3-5 times.
- 5. Wash with 4 5 column volumes of Binding Buffer containing 0.5 M NaCl
- 6. Wash with at least 2 column volumes of Binding Buffer and ensure the effluent reaches the same Absorbance (280 nm) as the Binding Buffer.
- 7. Elute with the elution buffer.
- 8. Combine fractions with highest absorbance.
- 9. Regenerate the column by:
 - a. Washing with ~ 5 volumes 5M NaCl.
 - b. Equilibrate with 5 volumes of Binding Buffer containing 20% Ethanol in PBS/dH₂O.
 - c. Store upright at 4 °C.

BUFFER EXAMPLES:

- (1) Binding Buffers: 1 X PBS
- (2) Elution Buffers: 3 M NaCl in PBS

RELATED PRODUCTS:

Recombinant Protein G- Sepharose Column Recombinant Protein L- Sepharose Column Recombinant Protein A/G- Sepharose Column Recombinant Protein A/G/L- Sepharose Column Recombinant Protein A Sepharose Beads Recombinant Protein G Sepharose Beads Recombinant Protein L Sepharose Beads Recombinant Protein A/G Sepharose Beads Recombinant Protein A/G Sepharose Beads Protein G Polyclonal Antibody Protein A Polyclonal Antibody Protein L Polyclonal Antibody Protein G Magnetic Beads Protein A Magnetic Beads Protein A/G Magnetic Beads Protein A/G/L Magnetic Beads

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