# abcam

## Product datasheet

## Heparin Sepharose® ab193268

#### 製品の概要

製品名

アッセイタイプ

検出感度

製品の概要

Heparin Sepharose®

Bead-based sandwich immunoassay (quantitative, multiplexable)

>= 0.4 mg/ml

High binding capacity (>0.4mg/mL). Minimal leaching of ligand. For column or batch purification of heparin-binding proteins & cation exchange (ab193268).

#### Contents:

Supplied as a 50% slurry in 20 % Ethanol; >2.5 mg heparin per mL Sepharose<sup>®</sup> beads.

#### Features:

Heparin beads have been widely used in affinity purification of various heparin-binding proteins or ligands, such as antithrombin III, lipoprotein, as well as DNA binding proteins (transcription factors, virus coat proteins etc). Abcam's Heparin Sepharose<sup>®</sup> is designed for purification of heparin-binding proteins and ligands. It can also be used as a high capacity cation exchange medium. Specific proteins can be separated by using different concentrations of salt or a salt gradient. This Heparin Sepharose<sup>®</sup> formulation exhibits excellent binding capacity, high flow rate, no significant loss of the heparin ligand and a pH stability range of 2-10.

These beads are for use in column purification. If used in batch purification, we recommend not exceeding  $150 \times g$  when centrifuging.

Store beads at 4°C.

The beads may be damaged above 40°C.

DO NOT FREEZE.

Wash beads 3 times with 3x bead volume of desired buffer before use.

#### **Applications:**

Purification of heparin-binding proteins, enzymes or other ligands.

Sepharose is a registered trademark of GE Healthcare

## 特記事項

This product is manufactured by BioVision, an Abcam company and was previously called 6553 Heparin Sepharose. 6553-10 is the same size as the 10 ml size of ab193268.

Heparin Sepharose<sup>®</sup> is prepared by covalently coupling heparin to epoxy-activated 6% cross-linked Sepharose<sup>®</sup> beads. The coupling was optimized to give a high binding capacity and could be greater than 0.4 mg of heparin-binding protein (such as thrombin) per ml of wet gel.

### **Suggested Protocol:**

Wash column with ddH2O to remove air bubbles.

Fill column with heparin beads.

Wash the column with 5X volume of Binding Buffer.

Dilute sample with Binding Buffer (1:1 ratio) or change the sample solution to binding buffer by means of your choice.

Add the sample solution onto the column.

Collect the solution and repeat step 5 & 6 several times if necessary.

Wash the column 5-10 times with the Binding Buffer.

Add Elution Buffer to elute bound protein.

Collect the eluent using microcentrifuge tube.

Assay protein concentration and combine the fractions containing sufficient heparin-binding protein.

Bead can be cleaned and regenerated by washing with 2-3x volume of high concentration salt solution and then the binding buffer.

## アプリケーション

#### 適用あり: Purification

## 製品の特性

### 保存方法

#### Store at +4°C. Please refer to protocols.

内容	10 ml	1 ml	50 ml
Heparin Sepharose <sup>®</sup>	1 x 10ml	1 x 1ml	1 x 50ml

#### アプリケーション

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アプリケーション	Abreviews	特記事項
Purification		Use at an assay dependent concentration. Purification of heparin-binding proteins, enzymes or other ligands.

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