abcam

Product datasheet

Protein A Magnetic Beads ab214286

5 References

概述

产品名称 蛋白A Magnetic Beads

样**品**类型 Serum, Cell Lysate

种属反应性 与反应: Mouse, Rabbit, Horse, Guinea pig, Hamster, Cow, Human, Pig

不与反应: Rat, Sheep, Goat, Chicken

产品概述 Features:

Easy to use, high-binding capacity, non-adherent beads.

Support Characteristics: Paramagnetic, spherical, 6 % cross-linked agarose.

Ligand: Recombinant Protein A.

Particle Size: 75 – 150 µm.

Binding Capacity: Generally >25 mg human lgG/ml wet beads.

Working Temperature: Room temperature.

Storage Solution: PBS with 0.02% Sodium Azide.

Storage Temperature: 4 – 8 °C.

Applications:

Useful for immunoprecipitation and enrichment of IgG antibodies.

High affinity for Fc region of IgG antibodies from a variety of species.

Protein A binds to most human and mouse IgG subclasses (e.g., human IgG1, IgG2, IgG4; mouse IgG1, IgG2a, IgG2b, IgG3).

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It also binds to total IgG from cow, guinea pig, hamster, horse, pig, and rabbit. Protein A has little affinity to chicken, goat, rat and sheep.

说明

This product is manufactured by BioVision, an Abcam company and was previously called 6507 Protein A Magnetic Beads. 6507-1 is the same size as the 1 ml size of ab214286.

Description:

Protein A Magnetic Beads are prepared by covalently coupling Recombinant Protein A to 6% crosslinked magnetically beaded agarose. The coupling technique is optimized to give a high binding capacity for lgG. The capacity of lgG binding is generally greater than 25 mg of human lgG per ml of wet gel.

SUGGESTED PROTOCOL:

Prepare the antibody solution by diluting the required amount of antibody in binding buffer before running the protocol.

- 1. Magnetic Bead Preparation (perform three times)
- a. Dispense the required amount of magnetic beads into a 1.5 ml microfuge tube.
- b. Place the tube in the magnetic rack and remove the storage solution.
- c. Add 500 µl binding buffer.
- d. Resuspend the beads.
- e. Remove the liquid
- 2. Antibody Capture
- a. Immediately add the antibody solution.
- b. Resuspend and mix (slow end-over-end) for at least 15 minutes.
- c. Remove the liquid.
- 3. Washing
- a. Add 500 µl Binding Buffer containing 0.5 M NaCl; Remove the liquid.
- b. Add 500 µl Binding Buffer; Remove the liquid.
- 4. Target Binding
- a. Add sample diluted in binding buffer.
- b. Incubate with slow end-over-end mixing for up to 60 minutes.
- c. Remove and collect unbound fraction.
- 5. Washing (perform three times)
- a. Add 500 µl wash buffer
- b. Remove liquid (save washes to troubleshoot)
- 6. Elution (perform three times)

- a. Add 2 volumes elution buffer (vs. bead volume).
- b. Completely resuspend beads and incubate at least 2 minutes.
- c. Remove and collect elution fraction.

RECOMMENDED BUFFER EXAMPLES:

Binding buffer: 50 mM Tris, 150 mM NaCl, pH 7.5

Wash buffer: 50 mM Tris, 150 mM NaCl, pH 7.5 (or add 1% Octylglucoside to this buffer) (Could

also try 1X PBS as both binding and wash buffer)

Elution buffer: 0.1 M -0.2 M Glycine pH 2.5-3.1 (or 0.1 M citric acid, pH 2.5-3.1 or 2.5 % Acetic

Acid)

经测试应用 适用于: Purification, IP

性能

存放说明

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

组 件	1 ml
Protein A magnetic beads	1 x 1ml

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab214286于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Purification		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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