Ver. HB230112

Anti-His Affinity Gel

Product description

Anti-His Affinity Gel is prepared by conjugating high quality murine IgG1 monoclonal antibody with Sepharose 4B gel. This product has a high His label fusion protein loading capacity (at least 1.1 mg protein/mL gel) and less non-specific binding of miscellaneous proteins. It can be used for immunoprecipitation (IP) and purification of fusion proteins with His labels.

The Anit-His affinity purification gel provided by YEASEN can be used to purify N-terminal, C-terminal, $4 \times \text{His}$, $5 \times \text{His}$, $6 \times \text{His}$ fusion proteins.

Components

Components No.	Name	20589ES03	20589ES08	20589ES25	20589ES60
20589	Anti-His Affinity Gel	1 mL	5 mL	25 mL	100 mL

Specifications

Isotype	Mouse IgG1
Purity	Protein A
Antibody Concentration	7.5 g antibody/L Gel
Application	Protein purification、IP
Protein binding capacity	≥1.1 mg protein/mL Gel
Storage buffer TBS, 50% glycerin, pH7.4, contains 0.02% (w/v) sodium azide	

Storage

The unopened products should be stably stored at -25°C ~-15°C for 1 year. Do not freeze in solution without glycerin!

Instructions

- 1. Fully re-suspension of Anti-His affinity purification gel to form a uniform solution as far as possible. Transfer 40-100 μ L mixture (20-50 μ L gel) to a new centrifuge tube.
- $2. \text{Add } 500 \,\mu\text{L TBS}$, gently reinsert the Anti-His Affinity Gel, centrifuge at $10,\!000 \,\text{rpm}$ for $60 \,\text{s}$, discard the supernatant, and repeat the above steps $3-4 \,\text{times}$.

Note: Remove supernatant as much as possible, but do not inhale gel.

- 3. $100 \,\mu\text{L}$ cell lysate was added to the above precipitation. Incubate at room temperature for at least 1 hour, shaking gently while incubating.
- 4. Centrifuge at 10,000 rpm for 60 s and transfer the supernatant to a new centrifuge tube for storage for later detection.
- 5. The precipitates were washed with 0.5mL PBST for 3-4 times and centrifuged at 10,000 rpm for 60 s to remove the supernatant.





6. Elution: 40 μ L 2 \times protein loading buffer was added and boiled for 10 min. Centrifuge at 10,000 rpm for 60 s and transfer supernatant carefully for SDS-PAGE detection.

Notes

- 1. All centrifugal operations are performed at 4°C.
- 2. For your safety and health, please wear lab coats and disposable gloves for operation.
- 3. For research use only.