



# **Immunoprecipitation/IP Kit- Anti-Influenza M1 Immunomagnetic Beads**

**Catalog Number: MB40010-T60**

Please read this instruction manual carefully before using the product

Product Contents

| Contents                                              | Package 1                                    | Package 2                                  | Storage            |
|-------------------------------------------------------|----------------------------------------------|--------------------------------------------|--------------------|
| Anti-Influenza M1 Immunomagnetic Beads <sup>1 3</sup> | 1 mL                                         | 5 mL                                       | 2-8℃ for 12 months |
| NP40 Cell Lysis Buffer                                | 4 mL                                         | 22 mL                                      | -20℃ for 12 months |
| 5×TBST (pH7.4)                                        | Required but not supplied                    |                                            |                    |
| 1×TBST (pH7.4)                                        | Required but not supplied                    |                                            |                    |
| ddH <sub>2</sub> O                                    | Required but not supplied                    |                                            |                    |
| Alkaline Elution Buffer                               | 3 mL                                         | 15 mL                                      | 2-8℃ for 12 months |
| Acidity Elution Buffer                                | 3 mL                                         | 15 mL                                      | 2-8℃ for 12 months |
| Neutralization Buffer                                 | 2 mL                                         | 8 mL                                       | 2-8℃ for 12 months |
| Magnetic Separator                                    | Not included (refer related product MAGS001) | One MAGS001 included in China <sup>2</sup> |                    |

- [1] The IP KIT contains anti-Influenza M1 Immunomagnetic Beads(2 mg/mL) in phosphate buffered saline (PBS, pH 7.4) with sodium azide (0.1%).
- [2] The Magnetic Separator cannot be included for oversea customers due to shipment prohibition.
- [3] Immunomagnetic Beads kits are shipped at ambient temperature in which immunomagnetic beads are provided in liquid buffer.

Product Description

The Anti-Influenza M1 Immunomagnetic Beads, conjugated with Anti-Influenza M1 antibody, are used for immunoprecipitation (IP) of Influenza M1 proteins which expressed in vitro expression systems and bacterial and mammalian cell lysates.

For IP, the beads are added to a sample containing Influenza M1 proteins to form a bead-protein complex. The complex is removed from the solution manually using a Magnetic Separator. The bound Influenza M1 proteins are dissociated from the Immunomagnetic Beads using an elution buffer.

Antibody Information

**Antibody:** Influenza A H1N1 (A/Puerto Rico/8/34/Mount Sinai) Matrix protein 1 / M1 Antibody, Rabbit PAb, Antigen Affinity Purified(40010-T60)

**Immunogen:** Recombinant Influenza A H1N1 (A/Puerto Rico/8/34/Mount Sinai) Matrix protein 1 / M1 Protein (Catalog#40010-V07E)

**Clone ID:**

**Isotype:** Rabbit IgG

**Specificity:** Influenza A H1N1 (A/Puerto Rico/8/34/Mount Sinai) Matrix protein 1 / M1

**Guaranteed Applications:** IP, Minimum Protein Purification

**Preparation:** Produced in rabbits immunized with purified, recombinant Influenza A H1N1 (A/Puerto Rico/8/34/Mount Sinai) Matrix protein 1 / M1 (Catalog#40010-V07E; NP\_040978.1; Ser2-Lys252). Influenza A H1N1 (A/Puerto Rico/8/34/Mount Sinai) Matrix protein 1 / M1 specific IgG was purified by Influenza A H1N1 (A/Puerto Rico/8/34/Mount Sinai) Matrix protein 1 / M1 affinity chromatography.

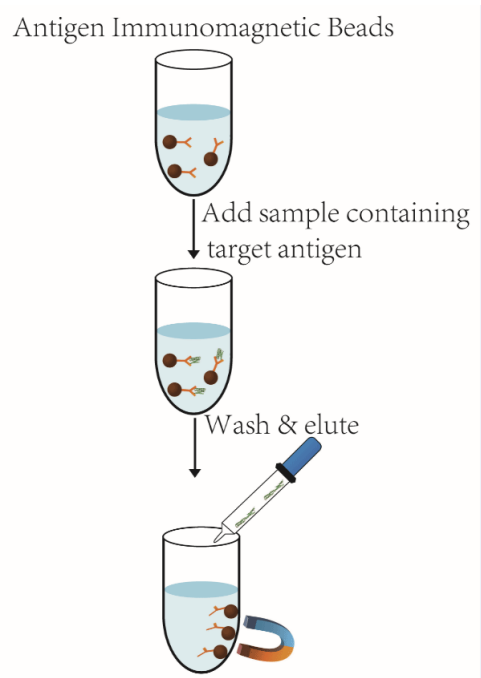


Fig. 1 Immunoprecipitation (IP) Protocol

## Protocol

The protocol (Fig. 1) uses 50  $\mu\text{L}$  Anti-Influenza M1Immunomagnetic Beads, but this can be scaled up or down as required.

### Cell Lysis

Cells may be lysed using any standard cell lysis protocol in accordance with your starting materials. We suggest using NP40 Cell Lysis Buffer (supplied with kit).

### Immunoprecipitate Target Antigen

1. Add 50  $\mu\text{L}$  of Immunomagnetic Beads into a 1.5 mL microcentrifuge tube.
2. Add 150  $\mu\text{L}$  of  $1\times$  TBST buffer to the Immunomagnetic Beads and gently vortex to mix.
3. Place the tube into a Magnetic Separator to collect the beads against the wall side of the tube. Remove and discard the supernatant.
4. Add 1 mL of  $1\times$  TBST buffer to the tube. Invert the tube several times or gently vortex to mix for 1 min. Collect Immunomagnetic Beads with a Magnetic Separator. Remove and discard the supernatant.
5. Add the sample containing target protein ( $\sim 100\text{ }\mu\text{g}$  of protein in 100  $\mu\text{L}$ ) to the pre-washed Immunomagnetic Beads, add 400  $\mu\text{L}$  of  $1\times$  TBST buffer and incubate at room temperature for 30 min with mixing.
6. Collect the Immunomagnetic Beads with a Magnetic Separator, remove the unbounded sample and save for analysis.
7. Add 300  $\mu\text{L}$  of  $5\times$  TBST buffer to the tube and gently mix. Collect the Immunomagnetic Beads and discard the supernatant. Repeat this wash twice.
8. Add 300  $\mu\text{L}$  of ddH<sub>2</sub>O to the tube and gently mix. Collect the Immunomagnetic Beads on a Magnetic Separator and discard the supernatant.

### Elute Target Antigen.

#### A. Alkaline Elution Protocols

1. Add 100  $\mu\text{L}$  of Alkaline Elution buffer to the tube.
2. Gently vortex to mix and incubate the sample at room temperature on a rotator for 5 min.
3. Magnetically separate the Immunomagnetic Beads and save the supernatant containing the target antigen.
4. To neutralize the sample, add 50  $\mu\text{L}$  of Neutralization Buffer for each 100  $\mu\text{L}$  of eluate.

#### B. Acidity Elution

1. Add 100  $\mu\text{L}$  Acidity Elution Buffer.
2. Gently vortex to mix and incubate the sample at room temperature on a rotator for 5-10 min.
3. Magnetically separate the Immunomagnetic Beads and save the supernatant containing the target antigen.

4. To neutralize the low pH, add 15  $\mu\text{L}$  of Neutralization Buffer for each 100  $\mu\text{L}$  of eluate.

#### C. Elution Using Sample Buffer

1. Add 100  $\mu\text{L}$  of SDS-PAGE sample buffer to the tube.
2. Gently vortex to mix and incubate the sample at 95-100°C for 5-10 min.
3. Magnetically separate the Immunomagnetic Beads and save the supernatant containing the antigen.

## Reference Information

### Related Products

| Products                                                                         | Cat No.  |
|----------------------------------------------------------------------------------|----------|
| Magnetic Separator-1.5 (2 tubes)                                                 | MAGS001  |
| Immunoprecipitation Kit<br>-Immunomagnetic Beads Protein A Kit                   | BA10600  |
| Immunoprecipitation Kit<br>-Immunomagnetic Beads Protein G Kit                   | BG13103  |
| Immunoprecipitation Kit<br>-Immunomagnetic Beads Protein L Kit                   | BL11044  |
| Immunoprecipitation Kit<br>-Anti-DYKDDDDK(Flag®) Tag<br>Immunomagnetic Beads Kit | TB101274 |
| Immunoprecipitation Kit<br>-Anti-GFP Tag Immunomagnetic Beads Kit                | TB13105  |
| Immunoprecipitation Kit<br>-Anti-Myc Tag Immunomagnetic Beads Kit                | TB100029 |
| Immunoprecipitation Kit<br>-Anti-HA Tag Immunomagnetic Beads Kit                 | TB100028 |
| Immunoprecipitation Kit<br>-Anti-V5 Tag Immunomagnetic Beads Kit                 | TB100378 |

### Trouble Shooting

| Problem                          | Possible Cause                      | Solution                                                        |
|----------------------------------|-------------------------------------|-----------------------------------------------------------------|
| Little or no protein is detected | Protein degraded                    | Include protease inhibitors (PMSF) in the lysis buffer          |
|                                  |                                     | Use new lysate or lysate stored at -80° C                       |
|                                  | No or minimal protein was expressed | Verify protein expression by SDS-PAGE or Western blot           |
|                                  |                                     | Analysis of the lysate using an positive control as a reference |
|                                  |                                     | Increase the amount of lysate used for IP/Co-IP                 |
|                                  |                                     | Use a more sensitive detection system                           |
|                                  |                                     |                                                                 |
|                                  |                                     |                                                                 |

| Problem                              | Possible Cause                                                           | Solution                                                                               |
|--------------------------------------|--------------------------------------------------------------------------|----------------------------------------------------------------------------------------|
| Magnetic Beads aggregated            | Magnetic Beads were frozen or centrifuged                                | Handle the Beads as directed in the instructions                                       |
|                                      | Buffer was incompatible with magnetic beads                              |                                                                                        |
|                                      | Detergent was not added to the wash and bind solutions                   |                                                                                        |
| Failure to co-IP interacting protein | Wash conditions were too stringent for the weak or transient interaction | Reduce the number of washes                                                            |
|                                      |                                                                          | Lower the ionic strength of the wash buffer                                            |
|                                      | Interacting protein was expressed at a low level                         | Apply additional protein sample                                                        |
|                                      |                                                                          | Use a more sensitive detection system                                                  |
|                                      | Buffer system was not optimal for the protein: protein interaction       | Optimize the co-IP buffer                                                              |
|                                      | Insufficient sample was loaded on the gel for Western blot detection     | Elute sample in 30% acetonitrile 0.5% formic acid, then                                |
|                                      |                                                                          | Bring the sample back up in SDS-PAGE Sample Buffer and load entire elution fraction on |