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## For Research Use Only.

## Not for use in diagnostic procedures.



#### Smart-IP Series

# Mouse IgG2b (isotype control) -Magnetic Beads

**CODE No.** M077-11

CLONALITY Monoclonal CLONE 3D12

ISOTYPE Mouse IgG2b κ
QUANTITY 20 tests (Slurry: 1 mL)

**SOURCE** Purified IgG from hybridoma supernatant

IMMUNOGEN KLH

**REACTIVITY**No specific binding is detected on immunoprecipitation.

10 mg magnetic beads in 1 mL PBS/0.1% BSA/0.09% NaN<sub>3</sub>

\*Azide may react with copper or lead in plumbing system to form explosive metal azides. Therefore, always flush plenty of water when disposing materials containing azide into drain.

**STORAGE** This beads suspension is stable for one year from the date of purchase when stored at 4°C.

If bead agglomeration is observed, please disperse the agglomerations by careful pipetting.

\*In particular, please check the inner wall of the vial and cap.

#### **APPLICATION-CONFIRMED**

Immunoprecipitation 50 μL of beads slurry/sample

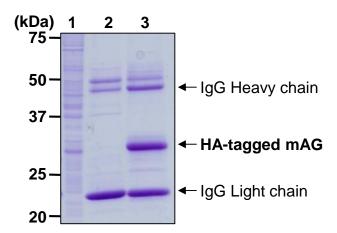
#### RELATED PRODUCTS

For more information, please visit our web site <a href="https://ruo.mbl.co.jp/">https://ruo.mbl.co.jp/</a>.

The descriptions of the following protocols are examples. Each user should determine the appropriate condition.

### **Immunoprecipitation**

- 1) Wash 2 x  $10^6$  cells 3 times with PBS and suspends them in 500  $\mu$ L of cold Extraction buffer [50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 1% NP-40].
- 2) Centrifuge the tube at 12,000 x g for 5 min. at 4°C and transfer the supernatant to another tube.
- 3) Add magnetic beads as suggested in the **APPLICATION** into 500  $\mu$ L of the cell lysate. Mix well and incubate with gentle agitation for 30 min. at 4°C.
- 4) Place the tube on the magnetic rack (MBL; code no. 3190) for a few seconds.
- 5) Remove the supernatant.
- 6) Wash the beads 4 times with 1 mL of cold Wash buffer [50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 0.05% NP-40] (place the tube on the magnetic rack for a few seconds).
- 7) Resuspend the magnetic beads in 20 µL of Laemmli's sample buffer, boil for 3 min., and place the tube on the magnetic rack for a few seconds.
- 8) Load 20 µL of the sample per lane in a 1-mm-thick SDS-polyacrylamide gel and carry out electrophoresis.
- 9) Visualize the protein bands by CBB staining.



#### Immunoprecipitation of HA-tagged protein

Sample: HEK293T cell lysate from 2 x10<sup>6</sup> cells + HA-tagged mAG 10 μg

Lane 1: Input (10 µL/lane)

Lane 2: Post-IP beads of Mouse IgG2b (isotype control)-Magnetic Beads (M077-11)

Lane 3: Post-IP beads of Anti-HA-tag mAb-Magnetic Beads (M180-11)