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For Research Use Only. Not for use in diagnostic procedures.



Smart-IP Series

Anti-V5-tag mAb-Magnetic Beads

CODE No.	M167-11
CLONALITY CLONE ISOTYPE	Monoclonal 1H6 Mouse IgG2a κ
QUANTITY	20 tests (Slurry: 1 mL)
SOURCE IMMUNOGEN FORMULATION	Purified IgG from hybridoma supernatant Carrier protein conjugated synthetic peptide, GKPIPNPLLGLDST (V5-tag) 10 mg magnetic beads in 1 mL PBS/0.1% BSA/0.1% ProClin 150 *Azide may react with copper or lead in plumbing system to form explosive metal azides. Therefore,
STORAGE	always flush plenty of water when disposing materials containing azide into drain. This gel slurry is stable for one year from the date of purchase when stored at 4°C.

APPLICATION-CONFIRMED

Immunoprecipitation

50 µL of beads slurry/sample

*The purification capacity of Anti-V5-tag mAb-Magnetic Beads varies depending upon the characteristics of a V5-tagged protein. For example, 50 µL of beads slurry bounds 3.2 µg of a V5-tagged protein (32 kDa).

APPLICATION-REPORTED

Co-Immunoprecipitation

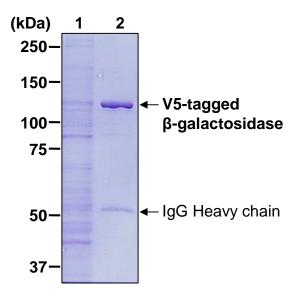
50 μ L of beads slurry/sample

For more information, please visit our website https://ruo.mbl.co.jp/.

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Immunoprecipitation

- 1) Wash 5 x 10⁶ cells 3 times with PBS and suspend with 1 mL of cold Lysis buffer [50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 1% NP-40] containing appropriate protease inhibitors.
- 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 400 μL of the supernatant prepared in step 2). Mix well and incubate with gentle agitation for 1 hr. 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 μ L of Laemmli's sample buffer, boil for 2 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 V5-tagged protein

Sample: V5-tagged β -galactosidase/HEK293T whole cell lysate Lane 1: Input (5 μ L/lane) Lane 2: Post-IP beads of Anti-V5-tag mAb (MBL; code no. M167-11)