

Smart-IP Series

# Anti-Myc-tag mAb-Magnetic Beads

<b>CODE No.</b>	M047-11
<b>CLONALITY</b>	Monoclonal
<b>CLONE</b>	PL14
<b>ISOTYPE</b>	Mouse IgG1
<b>QUANTITY</b>	20 tests (Slurry: 1 mL)
<b>SOURCE</b>	Purified IgG from mouse ascites fluid
<b>IMMUNOGEN</b>	6myc-tagged fusion protein
<b>FORMULATION</b>	15 mg magnetic beads in 1 mL PBS/0.1% BSA/0.1% ProClin 150
<b>STORAGE</b>	This beads suspension 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-Myc-tag mAb-Magnetic Beads varies depending upon the characteristics of a Myc-tagged protein.

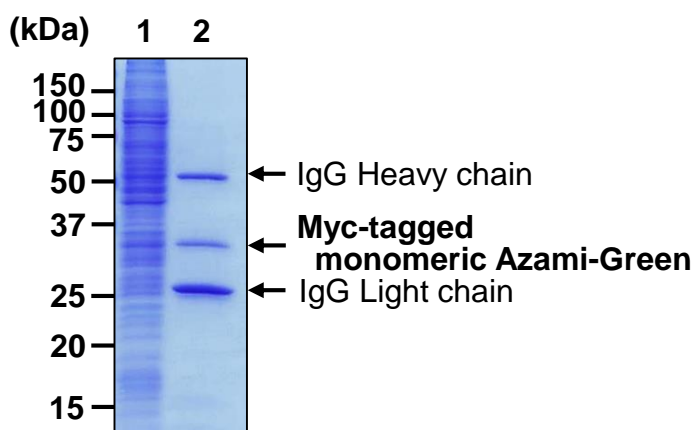
For example, 50 µL of beads slurry bounds 1.5 µg of a Myc-tagged protein (35 kDa).

**REFERENCE** 1) Jain, P., *et al.*, *Oncogene* **36**, 6348-6358 (2017) [Co-IP]

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

### **Immunoprecipitation**

- 1) Wash  $1 \times 10^7$  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  $12,000 \times g$  for 5 min. at  $4^\circ\text{C}$  and transfer the supernatant to another tube.
- 3) Add magnetic beads as suggested in the **APPLICATION** and purified Myc-tagged protein into 300  $\mu\text{L}$  of the supernatant prepared in step 2). Mix well and incubate with gentle agitation for 1 hr. at  $4^\circ\text{C}$ .
- 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  $\mu\text{L}$  of Laemmli's sample buffer, boil for 2 min., and place the tube on the magnetic rack for a few seconds.
- 8) Load 20  $\mu\text{L}$  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 Myc-tagged protein***

Sample: 293T cell lysate from  $3 \times 10^6$  cells + Myc-tagged monomeric Azami-Green 10  $\mu\text{g}$

Lane 1: Input (5  $\mu\text{L}/\text{lane}$ )

Lane 2: Post-IP beads of Anti-Myc-tag mAb (MBL, code no. M047-11)