

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

*Smart-IP Series*

# Anti-V5-tag mAb-Magnetic Beads

<b>CODE No.</b>	M215-11
<b>CLONALITY</b>	Monoclonal
<b>CLONE</b>	OZA3
<b>ISOTYPE</b>	Mouse IgG2b $\kappa$
<b>QUANTITY</b>	20 tests (Slurry: 1 mL)
<b>SOURCE</b>	Purified IgG from hybridoma supernatant
<b>IMMUNOGEN</b>	Carrier protein conjugated synthetic peptide, GKPIPPLLGLDST (V5-tag)
<b>FORMULATION</b>	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.

## **APPLICATION-CONFIRMED**

Immunoprecipitation 50  $\mu$ L of beads slurry/sample

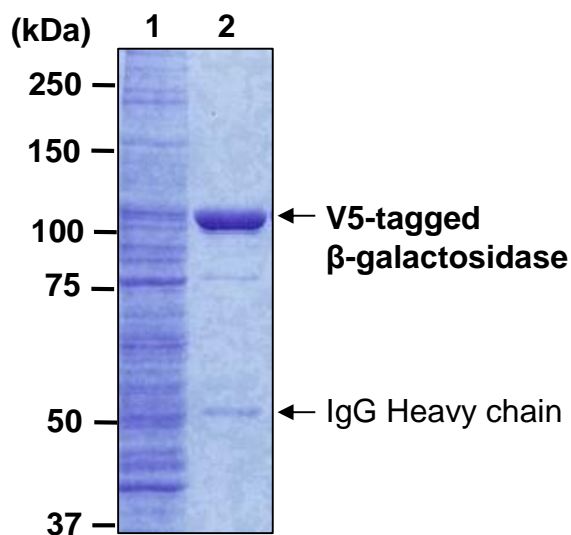
\*The purification capacity of Anti-V5-tag mAb-Magnetic Beads varies depending upon the characteristics of a V5-tagged protein. For example, 50  $\mu$ L of beads slurry bounds 3.2  $\mu$ g of a V5-tagged protein (35 kDa).

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

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

### **Immunoprecipitation**

- 1) Wash  $5 \times 10^6$  cells 3 times with PBS and suspend with 1 mL of 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 400  $\mu$ L 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 bead pellet 4 times with 1 mL of 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 bead pellet in 20  $\mu$ 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  $\mu$ 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 V5-tagged protein***

Sample: V5-tagged  $\beta$ -galactosidase/HEK293T whole cell lysate  
Lane 1: Input (10  $\mu$ L/lane)  
Lane 2: Post-IP beads of Anti-V5-tag mAb (M215-11)