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



Anti-Dab1 (Mouse) mAb

CODE No. D354-3

 CLONALITY
 Monoclonal

 CLONE
 4H11

 ISOTYPE
 Rat IgG1 κ

 QUANTITY
 100 μL, 1 mg/mL

SOURCE Purified IgG from hybridoma supernatant

IMMUNOGEN Mouse Dab1 (recombinant)

FORMULATION PBS containing 50% Glycerol (pH 7.2). No preservative is contained.

STORAGE This antibody solution is stable for one year from the date of purchase when stored at -20°C.

APPLICATION-CONFIRMED

Western blotting 1 μg/mL

APPLICATIONS-REPORTED

<u>Immunocytochemistry</u> Reference 1) <u>Immunohistochemistry</u> Reference 1)

SPECIES CROSS REACTIVITY on WB

Species	Human	Mouse	Rat	Hamster
Sample	Not tested	Transfectant	Not tested	Not tested
Reactivity		+		

Entrez Gene ID 13131 (Mouse)

REFERENCES 1) Onoue, A., et al., Neurosci. Res. 88, 23-27 (2014) [WB, IC, IHC-fr]

2) Uchida, T., et al., J. Neurosci. 29, 10653-10662 (2009)

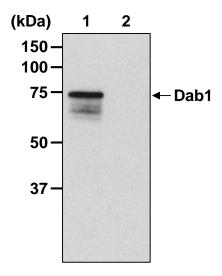
3) Morimura, T., et al., J. Biol. Chem. 280, 16901-16908 (2005)

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SDS-PAGE & Western blotting

- 1) Boil the sample for 2 min. and centrifuge. Load 20 μ L of the sample per lane in a 1-mm-thick SDS-polyacrylamide gel (7.5% acrylamide) for electrophoresis.
- 2) Blot the protein to a polyvinylidene difluoride (PVDF) membrane at 1 mA/cm² for 1 hr. in a semi-dry transfer system (Transfer Buffer: 25 mM Tris, 190 mM glycine, 20% methanol). See the manufacturer's manual for precise transfer procedure.
- 3) To reduce nonspecific binding, soak the membrane in 10% skimmed milk (in PBS, pH 7.2) overnight at 4°C.
- 4) Wash the membrane with PBS-T [0.05% Tween-20 in PBS] (5 min. x 3).
- 5) Incubate the membrane with primary antibody diluted with 1% skimmed milk (in PBS, pH 7.2) as suggested in the **APPLICATION** for 1 hr. at room temperature. (The concentration of antibody will depend on the conditions.)
- 6) Wash the membrane with PBS-T (5 min. x 3).
- 7) Incubate the membrane with HRP-conjugated anti-rat IgG antibody diluted with 1% skimmed milk (in PBS, pH 7.2) for 1 hr. at room temperature.
- 8) Wash the membrane with PBS-T (5 min. x 3).
- 9) Wipe excess buffer on the membrane, and then incubate it with appropriate chemiluminescence reagent for 1 min. Remove extra reagent from the membrane by dabbing with paper towel, and seal it in plastic wrap.
- 10) Expose to an X-ray film in a dark room for 3 min. Develop the film as usual. The condition for exposure and development may vary.

(Positive control for Western blotting; Transfectant)



Western blot analysis of mouse Dab1

Lane 1: Mouse Dab1/COS-7

Lane 2: COS-7

Immunoblotted with Anti-Dab1 (Mouse) mAb (MBL, code no. D354-3)

The sample was kindly provided by Dr. Mitsuharu Hattori. (Department of Biomedical Science, Graduate School of Pharmaceutical Sciences, Nagoya City University)