

Product Data Sheet

Purified anti-Microtubule Associated Protein 2ab (MAP2ab)

Catalog # / Size: 631101 / 25 µg

631102 / 100 µg

Clone: MT-08

Isotype: Mouse IgG1, κ

Immunogen: Microtubule protein enriched for kinesin

Reactivity: Human, Porcine, Mouse

Preparation: The antibody was purified by affinity chromatography.

Formulation: This antibody is provided in phosphate-buffered solution, pH 7.2, containing

0.09% sodium azide. Final antibody concentration is 0.5 mg/ml.

Concentration: 0.5 mg/ml

Storage: Upon receipt, store undiluted at 4°C.

Applications:

Applications: WB - Quality tested IP, IHC, ICC, ELISA - Reported in the literature

Recommended Usage: Each lot of this antibody is quality control tested by Western blotting. Western

blotting, suggested working dilution(s): Use 5 µg antibody per 5 ml antibody dilution buffer for each mini-gel. It is recommended that the reagent be

titrated for optimal performance for each application.

Description: Microtubule associated protein 2 also know as MAP2, has three isoforms that include two high molecular weight forms MAP2A and MAP2B (~280KD), and

a low molecular weight form MAB2C (~70KD). MAP2 belongs to the

microtubule-associated protein family and expressed in the nervous system. The exact function of MAP2 is unknown but MAPs may stabilize the microtubules against depolymerization. They also seem to have a stiffening effect on microtubules. MAP2 has been shown to interact with Fyn, Grb2, PRKACA, and CPEB1. The MT-08 antibody has been shown to be useful for Western blotting, Immunoprecipitation, Immunohistochemistry (frozen

sections), Immunocytochemistry, and ELISA of the human, mouse and porcine MAP2A and MAP2B isoforms.

Antigen References: 1. Garner CC, et al. 1988. Nature 336:674.

2. Kindler S, et al. 1994. Molec. Brain Res. 26:218.

3. Chang L, et al. 2003. Dev. Cell 4:521.

Related Products: Product

Clone Purified Mouse IgG1, κ Isotype Ctrl MOPC-21 HRP Goat anti-mouse IgG (minimal x-reactivity) Poly4053

MAP2ab 220 97. 46.

Mouse brain tissue extract was resolved by electrophoresis, transferred nitrocellulose, and probed with monoclonal anti-MAP2ab (clone MT-08) antibody. Proteins were visualized using HRP goat anti-mouse IgĞ and a chemiluminescence detection system.

FC, ICFC, ICC, IF, IHC, IP, WB

ELISA, IHC, WB



