

## MAP2 Ab

[References\(2\)](#) [Images\(17\)](#)

Cat.#: AF4081	Concn.: ~1mg/ml	Mol.Wt.: 55-75(2c/2d), 280(2a/2b)kd
Size: 100ul,200ul,50ul	Source: Rabbit	Clonality: Polyclonal

**Application:** WB 1:500-1:1000, IF/ICC 1:200, IHC 1:50-1:200, ELISA(peptide)  
1:20000-1:40000

\*The optimal dilutions should be determined by the end user.

**Reactivity:** Human, Mouse, Rat

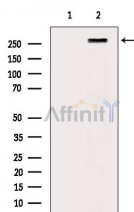
**Purification:** The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).

**Immunogen:** A synthesized peptide derived from human MAP2, corresponding to a region within N-terminal amino acids.

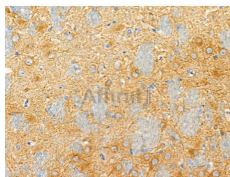
**Uniprot:** P11137

**Description:** 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.

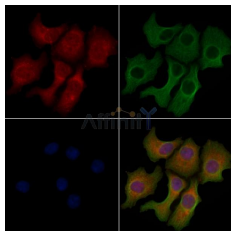
**Storage:** Supplied at 1.0mg/mL in phosphate buffered saline (without Mg<sup>2+</sup> and Ca<sup>2+</sup>), pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20 °C. Stable for 12 months from date of receipt.



Western blot analysis of extracts from Mouse muscle, using MAP2 Ab. The lane on the left was treated with blocking peptide.



AF4081 at 1/100 staining Rat brain tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the primary Ab at 4°C overnight. An HRP conjugated anti-Rabbit Ab was used as the secondary Ab.



AF4081 staining HeLa cells by IF/ICC. The samples were fixed with PFA and permeabilized in 0.1% Triton X-100, then blocked in 10% serum for 45 minutes at 25°C. Samples were then incubated with primary Ab(AF4081) and mouse anti-beta tubulin Ab(T0023) for 1 hour at 37°C. An AlexaFluor594 conjugated goat anti-rabbit IgG(H+L) Ab(Red) and an AlexaFluor488 conjugated goat anti-mouse IgG(H+L) Ab(Green) were used as the secondary Ab.

The nuclear counter stain is DAPI (blue).

**IMPORTANT:** For western blot, incubate membrane with diluted primary Ab in 5% w/v milk, 1X TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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