

Monoclonal Antibody to MAP-2 - FITC

Alternate names:	MAP2, Microtubule-associated protein 2, Neuronal Marker
Catalog No.:	BM2472F
Quantity:	0.25 mg
Concentration:	0.1 mg/ml (OD280)
Background:	Microtubule Associated Protein 2 exists in two high molecular weight forms (2a and 2b) and a low molecular weight form (2c). The expression of MAP2 is developmentally regulated and its multiple forms arise by alternative splicing of a single gene.
Uniprot ID:	P11137
NCBI:	NP_002365
GeneID:	4133
Host / Isotype:	Mouse / IgG1
Clone:	AP20
Format:	State: Liquid Ig fraction Purification: Protein G affinity chromatography Buffer System: 0.01 M PBS, pH 7.2, containing 2 mM EDTA, 1% BSA and 0.09% sodium azide Label: FITC – Fluorescein has been covalently attached to anti-MAP-2 and purified to assure optimal fluorochrome/protein molar ratio <i>Absorption / Emission:</i> 492 nm / 518 nm <i>Molar Ratio:</i> F / P = 3.75
Applications:	This antibody can be used in immunohistochemistry on tissues or cell using immunofluorescence techniques. The suggested working dilution for immunohistochemical staining is approximately 1:500. Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.
Specificity:	Anti-MAP-2 recognizes (Mr 300 kDa) MAP-2 protein from bovine brain and dendrites and cell bodies of neurons. Anti-MAP-2 (Clone AP20) reacts with the high molecular weight forms (2a & 2b) of MAP-2 but not with the low molecular weight form (2c). Species: Bovine. The antibody strongly cross-reacts with MAP-2 from human, rat, mouse, Xenopus, quail and chicken brains. Other species not tested.
Storage:	Store the antibody at 2-8°C. DO NOT FREEZE. Protect from light. Shelf life: one year from despatch.

For research and in vitro use only. Not for diagnostic or therapeutic work.

Material Safety Datasheets are available at www.acris-antibodies.com or on request.

Antibody Hotline - Technical Questions - Antibody Location Service
Free Call: 0800-2274746 (Germany only) - www.acris-antibodies.com

- General References:** 1. Kalcheva N; et al. Journal of Neurochemistry, 1994 Dec, 63(6):2336-41.
2. Binder LI; et al. Annals of the New York Academy of Sciences, 1986, 466:145-66

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