

## **MALT1 Antibody**



Orders: 877-616-CELL (2355)

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IP	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 90	Source/Isotype: Rabbit	UniProt ID: #Q9UDY8	Entrez-Gene Id: 10892
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		MALT1 Antibody detects endogenous levels of total MALT1 protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the carboxy terminus of mouse and rat MALT1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Mucosa-associated lymphoid tissue translocation gene 1 ( <i>MALT1</i> ) is a paracaspase that is a critical mediator of T-cell receptor activation of NF-κB and may contribute to the progression of MALT lymphomas (1-4). It contains two immunoglobulin-like domains, an amino-terminal death domain and a carboxy-terminal caspase-like domain. Association of MALT1 with Bcl-10 and CARD11/Carma1 leads to activation of IKK and subsequent stimulation of NF-κB, resulting in increased proliferation and inhibition of apoptosis (5,6). A common translocation in MALT B-cell non-Hodgkin lymphomas t(11;18) (q21;q21) results in the fusion of the amino terminus of API2 (c-IAP2), a member of the inhibitor of apoptosis protein family, to the carboxy terminus of MALT1 (1,2). The API2-MALT1 fusion protein likely leads to deregulation of NF-κB, contributing to increased oncogenic potential (7).				
Background References		1. Akagi, T. et al. (1999) <i>Oncogene</i> 18, 5785-5794. 2. Uren, A.G. et al. (2000) <i>Mol. Cell</i> 6, 961-967. 3. Ruland, J. et al. (2003) <i>Immunity</i> 19, 749-758. 4. Nakagawa, M. et al. (2006) <i>Leukemia</i> 20, 929-936. 5. Che, T. et al. (2004) <i>J. Biol. Chem.</i> 279, 15870-15876. 6. Lucas, P.C. et al. (2001) <i>J. Biol. Chem.</i> 276, 19012-19019. 7. Ho, L. et al. (2005) <i>Blood</i> 105, 2891-2899.				

**Species Reactivity** Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X

TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: Human M: Mouse R: Rat

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