

## Caspase-4 Antibody



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

Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 45	Source/Isotype: Rabbit	UniProt ID: #P49662	Entrez-Gene Id: 837
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Caspase-4 Antibody detects endogenous levels of total caspase-4 protein. Processing intermediate forms of caspase-4 are observed at 40 kDa and 32 kDa as previously reported (7). The antibody does not cross-react with other caspases.				
Species predicted to react based on 100% sequence homology		Monkey				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ile125 within the p20 subunit of human caspase-4 protein. Antibodies were purified by protein A and peptide affinity chromatography.				
Background		Caspase-4 (TX/ICH-2/ICE <sub>rel</sub> II) is a member of the caspase family of proteases that play a key role in the execution of apoptosis and activation of inflammatory cytokines (1-3). Expression of caspase-4 has been observed in most tissues except brain, with highest levels in placenta, lung, spleen, and peripheral blood lymphocytes (PBL). Caspase-4 was originally found to contribute to Fas-mediated apoptosis (4). Several caspases (including caspase-4, caspase-5, and mouse caspase-11 and -12) are most closely related to caspase-1 and are capable of inducing apoptosis when overexpressed but are better characterized in the proteolytic activation of inflammatory cytokines (5). Caspase-4 associates with TRAF6 and is involved in the LPS inducible production of inflammatory cytokines IL-8 and MIP1 in THP-1 cells (6). While caspase-4 and mouse caspase-12 localize to the endoplasmic reticulum (ER) and may be activated by drugs that induce ER stress (7), at least one study suggests that caspase-4 and caspase-12 are not essential for ER stress-induced apoptosis (8).				
Background References		1. Faucheu, C. et al. (1995) <i>EMBO J</i> 14, 1914-22. 2. Kamens, J. et al. (1995) <i>J Biol Chem</i> 270, 15250-6. 3. Munday, N.A. et al. (1995) <i>J Biol Chem</i> 270, 15870-6. 4. Kamada, S. et al. (1997) <i>Oncogene</i> 15, 285-90. 5. Martinon, F. and Tschopp, J. (2007) <i>Cell Death Differ</i> 14, 10-22. 6. Lakshmanan, U. and Porter, A.G. (2007) <i>J Immunol</i> 179, 8480-90. 7. Hitomi, J. et al. (2004) <i>J Cell Biol</i> 165, 347-56. 8. Obeng, E.A. and Boise, L.H. (2005) <i>J Biol Chem</i> 280, 29578-87.				
Species Reactivi	ty	Species reactivity is de	etermined by testin	g in at least one approve	ed 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

Cross-Reactivity Key H: Human

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