## Fas (C18C12) Rabbit mAb





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Applications: W, IHC-P	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 40-50	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P25445	Entrez-Gene Id: 355	
Product Usage Information		<b>Application</b> Western Blotting Immunohistochemistry (Paraffin)			<b>Dilution</b> 1:1000 1:250 - 1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.					
		For a carrier free (BSA and azide free) version of this product see product #49414.					
Specificity/Sen	<b>nsitivity</b> Fas (C18C12) Rabbit mAb detects endogenous levels of total human Fas protein.						
				uced by immunizing animals with a synthetic peptide corresponding to within the intracellular region of human Fas protein.			
Background		Association of the receptor Fas with its ligand FasL triggers an apoptotic pathway that plays an important role in immune regulation, development, and progression of cancers (1,2). Loss of function mutation in either Fas (lpr mice) or FasL (gld mice) leads to lymphadenopathy and splenomegaly as a result of decreased apoptosis in CD4-CD8- T lymphocytes (3,4). FasL (CD95L, Apo-1L) is a type II transmembrane protein of 280 amino acids (runs at approximately 40 kDa upon glycosylation) that belongs to the TNF family, which also includes TNF-α, TRAIL, and TWEAK. Binding of FasL to its receptor triggers the formation of a death-inducing signaling complex (DISC) involving the recruitment of the adaptor protein FADD and caspase-8 (5). Activation of caspase-8 from this complex initiates a caspase cascade resulting in the activation of caspase-3 and subsequent cleavage of proteins leading to apoptosis. Unlike Fas, which is constitutively expressed by various cell types, FasL is predominantly expressed on activated T lymphocytes, NK cells, and at immune privileged sites (6). FasL is also expressed in several tumor types as a mechanism to evade immune surveillance (7). Similar to other members of the TNF family, FasL can be cleaved by metalloproteinases producing a 26 kDa trimeric soluble form (8,9).					
Background Re	eferences	1. Suda, T. et al. (1993) <i>Cell</i> 75, 1169-78. 2. Lee, H.O. and Ferguson, T.A. (2003) <i>Cytokine Growth Factor Rev</i> 14, 325-35. 3. Watanabe-Fukunaga, R. et al. (1992) <i>Nature</i> 356, 314-7. 4. Hahne, M. et al. (1995) <i>Int Immunol</i> 7, 1381-6. 5. Nagata, S. (1997) <i>Cell</i> 88, 355-65. 6. Green, D.R. and Ferguson, T.A. (2001) <i>Nat Rev Mol Cell Biol</i> 2, 917-24. 7. Walker, P.R. et al. (1997) <i>J Immunol</i> 158, 4521-4. 8. Kayagaki, N. et al. (1995) <i>J Exp Med</i> 182, 1777-83. 9. Tanaka, M. et al. (1995) <i>EMBO J</i> 14, 1129-35.					
		Cranica vanstivitaria da			d angligation (a.g.		
Species Reactiv	vity	Species reactivity is determined by testing in at least one approved application (e.g., western blo		western blot).			
Western Blot B	Suffer	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 K	ey	W: Western Blotting IHC-P: Immunohistochemistry (Paraffin)					
Cross-Reactivit	y Key	H: Human					
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		U.S. Patent No. 7,429,	487, foreign equival	ents, and child patents	deriving therefrom.		

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