Fas (4C3) Mouse mAb



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Applications: W, IP, IF-IC, FC-FP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 40-50	Source/Isotype: Mouse IgG1	UniProt ID: #P25445	Entrez-Gene Id: 355
Product Usage Information		Application Western Blotting Immunoprecipitation Immunofluorescence Flow Cytometry (Fixed	•	istry)		Dilution 1:1000 1:50 1:50 1:100
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Fas (4C3) Mouse mAb recognizes endogenous levels of total Fas protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a recombinant protein specific to the carboxy terminus 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 References		 Suda, T. et al. (1993) Cell 75, 1169-78. Lee, H.O. and Ferguson, T.A. (2003) Cytokine Growth Factor Rev 14, 325-35. Watanabe-Fukunaga, R. et al. (1992) Nature 356, 314-7. Hahne, M. et al. (1995) Int Immunol 7, 1381-6. Nagata, S. (1997) Cell 88, 355-65. Green, D.R. and Ferguson, T.A. (2001) Nat Rev Mol Cell Biol 2, 917-24. Walker, P.R. et al. (1997) J Immunol 158, 4521-4. Kayagaki, N. et al. (1995) J Exp Med 182, 1777-83. Tanaka, M. et al. (1995) EMBO J 14, 1129-35. 				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
Western Blot Bu	ıffer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfadry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				

Applications Key

W: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry) FC-

FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key H: Human

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