Revision 1

Store at -20C

#68405

FasL (D1N5E) Rabbit mAb

Cell Signaling TECHNOLOGY®

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Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W	H	Endogenous	40	Rabbit IgG	#P48023	356

Product Usage Information

Application	Dilution
Western Blotting	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.

Specificity/Sensitivity

FasL (D1N5E) Rabbit mAb recognizes endogenous levels of total FasL protein.

Source / Purification

Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the carboxy terminus of human FasL protein. The antigen is within the extracellular domain of FasL.

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

1. Suda, T. et al. (1993) *Cell* 75, 1169-78.
2. Lee, H.O. and Ferguson, T.A. (2003) *Cytokine Growth Factor Rev* 14, 325-35.
3. Watanabe-Fukunaga, R. et al. (1992) *Nature* 356, 314-7.
4. Hahne, M. et al. (1995) *Int Immunol* 7, 1381-6.
5. Nagata, S. (1997) *Cell* 88, 355-65.
6. Green, D.R. and Ferguson, T.A. (2001) *Nat Rev Mol Cell Biol* 2, 917-24.
7. Walker, P.R. et al. (1997) *J Immunol* 158, 4521-4.
8. Kayagaki, N. et al. (1995) *J Exp Med* 182, 1777-83.
9. 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 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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Limited Uses

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