## TRADD (7G8) Rabbit mAb



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

<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 32	Source/Isotype: Rabbit	<b>UniProt ID:</b> #Q15628	Entrez-Gene Id: 8717
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 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		TRADD (7G8) Rabbit mAb detects endogenous levels of total TRADD protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding glycine 227 of human TRADD.				
Background		Apoptosis mediated by death factors like FasL and TNF-α involves the formation of a death-inducing signaling complex (DISC) to their respective receptors (1). Upon ligand activation to their receptors, Fas and TNF-R1 associate with death domain (DD) containing adaptor proteins FADD (Fas associated death domain) (2,3) and TRADD (TNF-R1 associated death domain) (4). In addition to its carboxy-terminal DD, FADD contains an amino-terminal death effector domain (DED) that binds to DEDs found on caspase-8 which leads to activation of this initiator caspase (5,6). Caspase-8 subsequently activates downstream effector caspases, like caspase-3, resulting in the cleavage of proteins involved in the execution of apoptosis. Unlike FADD, TRADD does not contain a DED (4). Apoptosis driven by TNF-R1 binding to TRADD involves association of TRADD and FADD which then leads to activation of caspase-8 (7).				
Background Re	ferences	1. Nagata, S. (1997) <i>Cell</i> 88, 355-65. 2. Chinnaiyan, A.M. et al. (1995) <i>Cell</i> 81, 505-12. 3. Boldin, M.P. et al. (1995) <i>J. Biol. Chem.</i> 270, 7795-8. 4. Hsu, H. et al. (1995) <i>Cell</i> 81, 495-504. 5. Muzio, M. et al. (1996) <i>Cell</i> 85, 817-27. 6. Boldin, M.P. et al. (1996) <i>Cell</i> 85, 803-15. 7. Hsu, H. et al. (1996) <i>Cell</i> 84, 299-308.				

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

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