

# TRAF4 (D1N3A) Rabbit mAb



Orders: 877-616-CELL (2355)

orders@cellsignal.com

**Support:** 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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

Applications: W	Reactivity: H M	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 50	Source/Isotype: Rabbit IgG	UniProt ID: #Q9BUZ4	Entrez-Gene Id: 9618
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 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		TRAF4 (D1N3A) Rabbit mAb recognizes endogenous levels of total TRAF4 protein. An unknown background band is detected in some cell lines at 80kDa.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Arg124 of human TRAF4 protein.				
Background		TRAFs (TNF receptor-associated factors) are a family of multifunctional adaptor proteins that bind to surface receptors and recruit additional proteins to form multiprotein signaling complexes capable of promoting cellular responses (1-3). Members of the TRAF family share a common carboxy-terminal "TRAF domain", which mediates interactions with associated proteins; many also contain aminoterminal Zinc/RING finger motifs. The first TRAFs identified, TRAF1 and TRAF2, were found by virtue of their interactions with the cytoplasmic domain of TNF-receptor 2 (TNFRII) (4). The six known TRAFs (TRAF1-6) act as adaptor proteins for a wide range of cell surface receptors and participate in the regulation of cell survival, proliferation, differentiation, and stress responses.  TRAF4, also referred to as CART1 and MLN62, is a divergent member of the TRAF family with relatively weak binding to TNFR family members (5-7). Interactions have been observed between TRAF4 and the neurotrophin receptor p75-NGFR, lymphotoxin-β receptor, and GITR (8-10). While originally identified in metastatic breast carcinoma, TRAF4 has been shown to contribute to tumor growth and invasion in various cancers including breast, lung and colon (11-13). Expression of Traf4 is induced by the tumor suppressor p53 in response to DNA damage, and can promote apoptosis (14).TRAF4 has also been shown to play a critical role in TGF-β signaling, where it has been found to antagonize the E3 ligase Smurf, resulting in enhanced receptor stabilization driving breast cancer metastasis (15).				
Background References		1. Arch, R.H. et al. (1998) <i>Genes Dev</i> 12, 2821-30. 2. Chung, J.Y. et al. (2002) <i>J Cell Sci</i> 115, 679-88. 3. Bradley, J.R. and Pober, J.S. (2001) <i>Oncogene</i> 20, 6482-91. 4. Rothe, M. et al. (1994) <i>Cell</i> 78, 681-92. 5. Kawamata, S. et al. (1998) <i>J Biol Chem</i> 273, 5808-14. 6. Régnier, C.H. et al. (1995) <i>J Biol Chem</i> 270, 25715-21. 7. Bièche, I. et al. (1996) <i>Cancer Res</i> 56, 3886-90. 8. Yang, K. et al. (2015) <i>Int J Clin Exp Pathol</i> 8, 1419-26. 9. Camilleri-Broët, S. et al. (2007) <i>Oncogene</i> 26, 142-7. 10. Li, W. et al. (2013) <i>Cancer Res</i> 73, 6938-50. 11. Ye, X. et al. (1999) <i>J Biol Chem</i> 274, 30202-8. 12. Esparza, E.M. and Arch, R.H. (2004) <i>Cell Mol Life Sci</i> 61, 3087-92. 13. Krajewska, M. et al. (1998) <i>Am J Pathol</i> 152, 1549-61. 14. Sax, J.K. and El-Deiry, W.S. (2003) <i>J Biol Chem</i> 278, 36435-44. 15. Zhang, L. et al. (2013) <i>Mol Cell</i> 51, 559-72.				

**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 M: Mouse

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