#99633

HOIP/RNF31 (E6M5B) Rabbit mAb



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Applications: Re W, IP	activity: H	Sensitivity: Endogenous	MW (kDa): 120	Source/Isotype: Rabbit IgG	UniProt ID: #Q96EP0	Entrez-Gene Id: 55072		
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 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/Sensitivi	ty	HOIP/RNF31 (E6M5B) Rabbit mAb recognizes endogenous levels of total HOIP/RNF31 protein.						
Source / Purification	n	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human HOIP/RNF31 protein.						
Background		HOIL-1-interacting protein (HOIP/RNF31), a RING-type E3 ubiquitin ligase, is the catalytic subunit of the Linear Ubiquitin Chain Assembly Complex (LUBAC) that is associated with TNF-R1 (1). Research studies have shown that the LUBAC consists of three subunits: HOIP, HOIL-1L, and Sharpin that facilitate canonical NF-kB activation in response to pro inflammatory cytokines through M1-linked linear ubiquitination of NEMO and RIP1 (2-6). As part of the LUBAC, HOIP has also been implicated in the negative regulation of interferon-mediated antiviral signaling through the suppression of RIG-I activation (7). The role of HOIP in LUBAC function and human disease is underscored by naturally occurring mutations in HOIP that impair LUBAC assembly and NF-kb activation. Patients that are homozygous for this mutation in HOIP have multi organ auto inflammation and immunodeficiency (8).						
Background Refere	nces	1. Haas, T.L. et al. (2009) <i>Mol Cell</i> 36, 831-44. 2. Smit, J.J. et al. (2012) <i>EMBO J</i> 31, 3833-44. 3. Tokunaga, F. et al. (2009) <i>Nat Cell Biol</i> 11, 123-32. 4. Tokunaga, F. et al. (2011) <i>Nature</i> 471, 633-6. 5. Blackwell, K. et al. (2013) <i>Mol Cell Biol</i> 33, 1901-15. 6. Ikeda, F. et al. (2011) <i>Nature</i> 471, 637-41. 7. Inn, K.S. et al. (2011) <i>Mol Cell</i> 41, 354-65. 8. Boisson, B. et al. (2015) <i>J Exp Med</i> 212, 939-51.						
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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