

## **MLKL Antibody (Mouse Specific)**



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

<b>Applications:</b> W, IP	<b>Reactivity:</b> M	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 54	Source/Isotype: Rabbit	UniProt ID: #Q9D2Y4	Entrez-Gene Id: 74568
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:100	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		MLKL Antibody (Mouse Specific) recognizes endogenous levels of total MLKL protein from mouse.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of mouse MLKL protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Necroptosis, a regulated pathway for necrotic cell death, is triggered by a number of inflammatory signals including cytokines in the tumor necrosis factor (TNF) family, pathogen sensors such as toll-like receptors (TLRs), and ischemic injury (1,2). The process is negatively regulated by caspases and is initiated through a complex containing the RIP1 and RIP3 kinases, typically referred to as the necrosome. Mixed lineage kinase domain-like protein (MLKL) is a pseudokinase that was identified as a downstream target of RIP3 in the necroptosis pathway (3,4). During necroptosis RIP3 is phosphorylated at Ser227, which recruits MLKL and leads to its phosphorylation at Thr357 and Ser358 (3). Knockdown of MLKL through multiple mechanisms results in inhibition of necroptosis (3-5). While the precise mechanism for MLKL-induced necroptosis is unclear, some studies have shown that necroptosis leads to oligomerization of MLKL and translocation to the plasma membrane, where it affects membrane integrity (6-9).				
Background References		<ol> <li>Christofferson, D.E. and Yuan, J. (2010) <i>Curr Opin Cell Biol</i> 22, 263-8.</li> <li>Kaczmarek, A. et al. (2013) <i>Immunity</i> 38, 209-23.</li> <li>Sun, L. et al. (2012) <i>Cell</i> 148, 213-27.</li> <li>Wang, Z. et al. (2012) <i>Cell</i> 148, 228-43.</li> <li>Wu, J. et al. (2013) <i>Cell Res</i> 23, 994-1006.</li> <li>Cai, Z. et al. (2014) <i>Nat Cell Biol</i> 16, 55-65.</li> <li>Chen, X. et al. (2014) <i>Cell Res</i> 24, 105-21.</li> <li>Wang, H. et al. (2014) <i>Mol Cell</i> 54, 133-46.</li> <li>Dondelinger, Y. et al. (2014) <i>Cell Rep</i> 7, 971-81.</li> </ol>				
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 nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				

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**Applications Key** 

W: Western Blotting IP: Immunoprecipitation

**Cross-Reactivity Key** 

M: Mouse

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