

**Malonyl-Lysine [Mal-K] MultiMab® Rabbit mAb mix**

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

<b>Applications:</b> W	<b>Reactivity:</b> All	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG
<b>Product Usage Information</b>	<b>Application</b> Western Blotting	<b>Dilution</b> 1:1000	
<b>Storage</b>	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.		
<b>Specificity/Sensitivity</b>	Malonyl-Lysine [Mal-K] MultiMab® Rabbit mAb mix recognizes endogenous levels of proteins only when malonylated at a lysine residue. This antibody does not cross-react with other lysine modifications.		
<b>Source / Purification</b>	MultiMab® Rabbit mAb mixes are prepared by combining individual rabbit monoclonal clones in optimized ratios for the approved applications. Each antibody in the mix is carefully selected based on motif recognition and performance in multiple assays. Each mix is engineered to yield the broadest possible coverage of the modification being studied while ensuring a high degree of specificity for the modification or motif.		
<b>Background</b>	<p>Lysine is subject to a wide array of regulatory post-translational modifications due to its positively charged ε-amino group side chain. The most prevalent of these are ubiquitination and acetylation, which are highly conserved among prokaryotes and eukaryotes (1,2). Acyl group transfer from the metabolic intermediates acetyl-, succinyl-, malonyl-, glutaryl-, butyryl-, propionyl-, and crotonyl-CoA all neutralize lysine's positive charge and confer structural alterations affecting substrate protein function. Lysine acetylation is catalyzed by histone acetyltransferases, HATs, using acetyl-CoA as a cofactor (3,4). Deacetylation is mediated by histone deacetylases, HDACs 1-11, and NAD-dependent Sirtuins 1-7. Some sirtuins have little to no deacetylase activity, suggesting that they are better suited for other acyl lysine substrates (5).</p> <p>SIRT5 is a predominantly mitochondrial desuccinylase and demalonylase (5,6). In the absence of a known malonyltransferase, nonenzymatic protein malonylation is likely driven by the concentration of malonyl-CoA and intracellular pH and is subject to metabolic fluctuations (7). Malonylation is especially prevalent among mitochondrial metabolic proteins. In type II diabetes mouse models, notably elevated malonylation can be detected mainly, but not exclusively, on proteins of glucose and fatty acid metabolism (8). Yeast histone H3K56 malonylation suggests poor DNA binding efficiency and may lead to reduced cell viability (9).</p>		
<b>Background References</b>	<div>1. Liu, Z. et al. (2014) <i>Nucleic Acids Res</i> 42, D531-6.</div> <div>2. Lee, S. (2013) <i>Toxicol Res</i> 29, 81-6.</div> <div>3. Lin, H. et al. (2012) <i>ACS Chem Biol</i> 7, 947-60.</div> <div>4. Zhang, Z. et al. (2011) <i>Nat Chem Biol</i> 7, 58-63.</div> <div>5. Du, J. et al. (2011) <i>Science</i> 334, 806-9.</div> <div>6. Peng, C. et al. (2011) <i>Mol Cell Proteomics</i> 10, M111.012658.</div> <div>7. Newman, J.C. et al. (2012) <i>J Biol Chem</i> 287, 42436-43.</div> <div>8. Du, Y. et al. (2015) <i>Mol Cell Proteomics</i> 14, 227-36.</div> <div>9. Xie, Z. et al. (2012) <i>Mol Cell Proteomics</i> 11, 100-7.</div>		
<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).		
<b>Western Blot Buffer</b>	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.		
<b>Applications Key</b>	<b>W:</b> Western Blotting		
<b>Cross-Reactivity Key</b>	<b>All:</b> All Species Expected		
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