## SUMO-2/3 (18H8) Rabbit mAb



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

<b>Applications:</b> W, IF-IC	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P55854, #P61956	<b>Entrez-Gene Id:</b> 6612, 6613
Product Usage Information		<b>Application</b> Western Blotting Immunofluorescence (Ir	nmunocytochemistry)		<b>Dilution</b> 1:1000 1:50 - 1:200
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		SUMO-2/3 (18H8) Rabbit mAb detects endogenous levels of SUMO-2/3. It does not cross-react with recombinant SUMO-1.			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide from the amino terminus of human SUMO-3.			
Background		Small ubiquitin-related modifier 1, 2 and 3 (SUMO-1, -2 and -3) are members of the ubiquitin-like protein family (1). The covalent attachment of the SUMO-1, -2 or -3 (SUMOylation) to target proteins is analogous to ubiquitination. This post-translational modification is a reversible, multi-step process that is initiated by cleaving a precursor protein to a mature protein. Mature SUMO-1, -2 or -3 is then linked to the activating enzyme E1, conjugated to E2 and in conjunction with E3, SUMO-1, -2 or -3 is ligated to the target protein (2). Ubiquitin and the individual SUMO family members are all targeted to different proteins with diverse biological functions. Ubiquitin predominantly regulates degradation of its target (1). In contrast, SUMO-1 is conjugated to RanGAP, PML, p53 and IkB-a to regulate nuclear trafficking, formation of subnuclear structures, regulation of transcriptional activity and protein stability (3-7). SUMO-2/-3 forms poly-(SUMO) chains, is conjugated to topoisomerase II and APP, regulates chromosomal segregation and cellular responses to environmental stress, and plays a role in the progression of Alzheimer disease (8-11).			
Background References		<ol> <li>Schwartz, D.C. and Hochstrasser, M. (2003) <i>Trends Biochem. Sci.</i> 28, 321-8.</li> <li>Kim, K.I. et al. (2002) <i>J. Cell Physiol.</i> 191, 257-68.</li> <li>Matunis, M.J. et al. (1996) <i>J. Cell Biol.</i> 135, 1457-70.</li> <li>Duprez, E. et al. (1999) <i>J. Cell Sci.</i> 112, 381-93.</li> <li>Gostissa, M. et al. (1999) <i>EMBO J.</i> 18, 6462-74.</li> <li>Rodriguez, M.S. et al. (1999) <i>EMBO J.</i> 18, 6455-61.</li> <li>Desterro, J.M. et al. (1998) <i>Mol. Cell</i> 2, 233-9.</li> <li>Tatham, M.H. et al. (2001) <i>J. Biol. Chem.</i> 276, 35368-74.</li> <li>Azuma, Y. et al. (2003) <i>Proc. Natl. Acad. Sci. USA</i> 100, 259-64.</li> <li>Saitoh, H. and Hinchey, J. (2000) <i>J. Biol. Chem.</i> 275, 6252-8.</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 BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** 

W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key** 

H: Human M: Mouse R: Rat

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