## LAMP1 (D2D11) XP<sup>®</sup> Rabbit mAb





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Applications: W, IP, IHC-P, IF-IC, FC-FP	Reactivity: H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 42 (non- glycosylated), 90- 120 (glycosylated)	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P11279	Entrez-Gene Id: 3916
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation Immunohistochemistry (Paraffin) Immunofluorescence (Immunocytochemistry) Flow Cytometry (Fixed/Permeabilized)			<b>Dilution</b> 1:1000 1:100 1:100 - 1:400 1:100 - 1:400 1:100 - 1:400	
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. For a carrier-free (BSA and azide free) version of this product see product #52252.				
Specificity/Sensitivity		LAMP1 (D2D11) XP <sup>®</sup> Rabbit mAb recognizes endogenous levels of total LAMP1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a recombinant protein fragment of human LAMP1 protein.				
Background		Lysosome-associated membrane protein 1 and 2 (LAMP1 and LAMP2) are two abundant lysosomal membrane proteins (1,2). Both are transmembrane proteins and are heavily glycosylated at the amino-terminal luminal side of the lysosomal inner leaflet, which protects the proteins from proteolysis (3). The carboxy terminus of LAMP1 is exposed to the cytoplasm and contains a tyrosine sorting motif that targets LAMP to lysosomal membranes (4). LAMP1 and LAMP2 are 37% homologous in their protein sequences. Both LAMP1 and LAMP2 are involved in regulating lysosomal motility during lysosome-phagosome fusion and cholesterol trafficking (5,6).				
Background Refe	rences	1. Eskelinen, E.L. et al. (2003) <i>Trends Cell Biol</i> 13, 137-45. 2. Fukuda, M. (1991) <i>J Biol Chem</i> 266, 21327-30. 3. Kundra, R. and Kornfeld, S. (1999) <i>J Biol Chem</i> 274, 31039-46. 4. Rohrer, J. et al. (1996) <i>J Cell Biol</i> 132, 565-76. 5. Huynh, K.K. et al. (2007) <i>EMBO J</i> 26, 313-24. 6. Eskelinen, E.L. et al. (2004) <i>Mol Biol Cell</i> 15, 3132-45.				
Species Reactivity		Species reactivity is determined by testing in at least one approve			d application (e.g., western blot).	
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.			primary antibody in 5% w/v BSA, 1X	
Applications Key		W: Western Blotting IP: Immunoprecipitation IHC-P: Immunohistochemistry (Paraffin) IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)				,
Cross-Reactivity Key		H: Human Mk: Monkey				
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