

# Anti-LAMP1 Antibody

Our Anti-LAMP1 primary antibody from PhosphoSolutions is mouse monoclonal. It detects human LAMP1 an  
Catalog # AN1434

## Product Information

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<b>Application</b>	WB, IHC, ICC
<b>Primary Accession</b>	<a href="#">P11279</a>
<b>Host</b>	Mouse
<b>Clonality</b>	Monoclonal
<b>Isotype</b>	IgG1
<b>Clone Names</b>	5H6
<b>Calculated MW</b>	44882

## Additional Information

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<b>Gene ID</b>	3916
<b>Other Names</b>	CD107 antigen like family member A antibody, CD107 antigen-like family member A antibody, CD107a antibody, CD107a antigen antibody, LAMP 1 antibody, LAMP-1 antibody, LAMP1 antibody, LAMP1_HUMAN antibody, LAMPA antibody, LGP120 antibody, IgpA antibody, Lysosomal membrane glycoprotein 120KD antibody, Lysosomal Associated Membrane Protein 1 antibody, Lysosome associated membrane glycoprotein 1 antibody, Lysosome-associated membrane glycoprotein 1 antibody, Lysosome-associated membrane protein 1 antibody, OTTHUMP00000040663 antibody
<b>Target/Specificity</b>	Lysosomal Associated Membrane Protein1 (LAMP1) is a protein that is localized primarily in lysosomes but may also be present on late endosomes and the plasma membrane. LAMP1 antibodies are therefore widely used as lysosome markers. It has recently been suggested that lysosomes are activated in microglia in the progression of multiple system atrophy (MSA) and thus play a key role in its pathology (Makioka et al., 2012).
<b>Dilution</b>	WB~~1:1000 IHC~~1:100~500 ICC~~N/A
<b>Format</b>	Protein G Purified
<b>Storage</b>	Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
<b>Precautions</b>	Anti-LAMP1 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.
<b>Shipping</b>	Blue Ice

## Background

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Lysosomal Associated Membrane Protein1 (LAMP1) is a protein that is localized primarily in lysosomes but may also be present on late endosomes and the plasma membrane. LAMP1 antibodies are therefore widely used as lysosome markers. It has recently been suggested that lysosomes are activated in microglia in the progression of multiple system atrophy (MSA) and thus play a key role in its pathology (Makioka et al., 2012).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.