

**LAMP2 (PT0249R) PT® Rabbit mAb**

<b>Catalog No :</b>	YM8157
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	WB;IHC;IF;IP;ELISA
<b>Target :</b>	CD107b
<b>Fields :</b>	>>Autophagy - animal;>>Lysosome;>>Phagosome;>>Tuberculosis
<b>Gene Name :</b>	LAMP2
<b>Protein Name :</b>	Lysosome-associated membrane glycoprotein 2
<b>Human Gene Id :</b>	3920
<b>Human Swiss Prot No :</b>	P13473
<b>Mouse Swiss Prot No :</b>	P17047
<b>Specificity :</b>	endogenous
<b>Formulation :</b>	PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA
<b>Source :</b>	Monoclonal, rabbit, IgG, Kappa
<b>Dilution :</b>	IHC 1:200-1:1000,WB 1:1000-1:5000,IF 1:200-1:1000,ELISA 1:5000-1:20000,IP 1:50-1:200,
<b>Purification :</b>	Protein A
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)
<b>Molecularweight :</b>	45kD
<b>Observed Band :</b>	120kD

**Background :**

The protein encoded by this gene is a member of a family of membrane glycoproteins. This glycoprotein provides selectins with carbohydrate ligands. It may play a role in tumor cell metastasis. It may also function in the protection, maintenance, and adhesion of the lysosome. Alternative splicing of this gene results in multiple transcript variants encoding distinct proteins. [provided by RefSeq, Jul 2008],

**Function :**

disease:Defects in LAMP2 are the cause of Danon disease (DAND) [MIM:300257]; also known as glycogen storage disease type 2B (GSD2B). DAND is a lysosomal glycogen storage disease characterized by the clinical triad of cardiomyopathy, vacuolar myopathy and mental retardation. It is often associated with an accumulation of glycogen in muscle and lysosomes.,function:Implicated in tumor cell metastasis. May function in protection of the lysosomal membrane from autodigestion, maintenance of the acidic environment of the lysosome, adhesion when expressed on the cell surface (plasma membrane), and inter-and intracellular signal transduction.,PTM:O- and N-glycosylated; some of the 16 N-linked glycans are polylactosaminoglycans.,similarity:Belongs to the LAMP family.,subcellular location:This protein shuttles between lysosomes, endosomes, and the plasma membrane.,tissue specificity:Isoform LAMP-2A

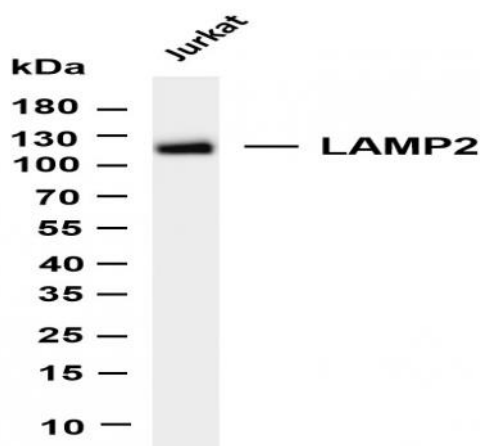
**Subcellular Location :**

Membranous

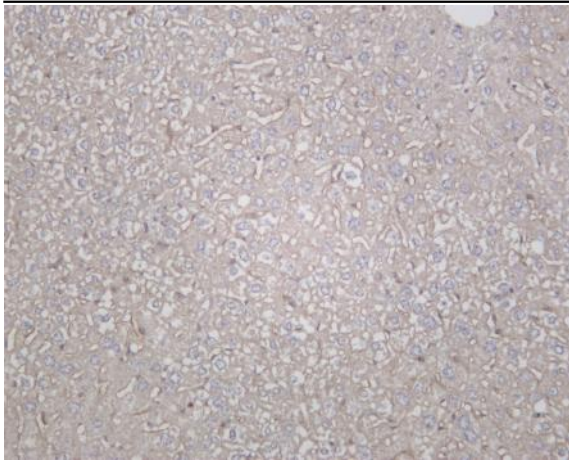
**Expression :**

Isoform LAMP-2A is highly expressed in placenta, lung and liver, less in kidney and pancreas, low in brain and skeletal muscle (PubMed:7488019, PubMed:26856698). Isoform LAMP-2B is detected in spleen, thymus, prostate, testis, small intestine, colon, skeletal muscle, brain, placenta, lung, kidney, ovary and pancreas and liver (PubMed:7488019, PubMed:26856698). Isoform LAMP-2C is detected in small intestine, colon, heart, brain, skeletal muscle, and at lower levels in kidney and placenta (PubMed:26856698).

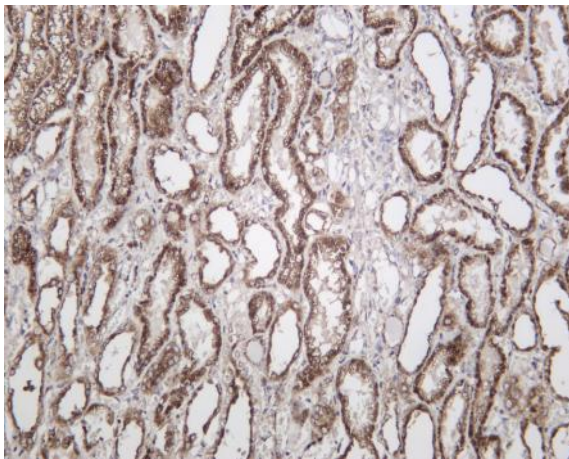
## Products Images



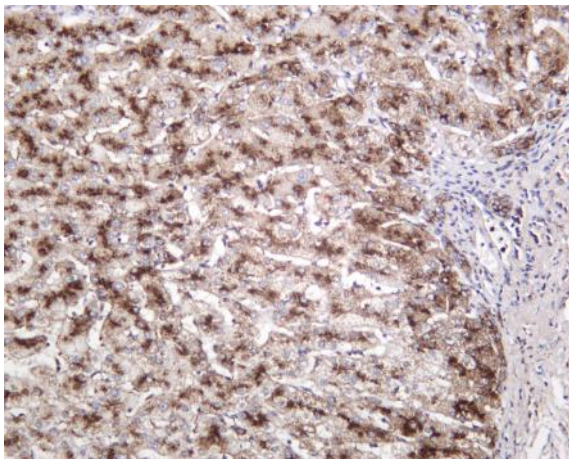
Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-LAMP2 (PT0249R) antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: Jurkat  
Predicted band size: 45kDa Observed band size: 120kDa



Mouse liver was stained with Anti-LAMP2 (PT0249R) rabbit antibody



Human kidney was stained with Anti-LAMP2 (PT0249R) rabbit antibody



Human liver was stained with Anti-LAMP2 (PT0249R) rabbit antibody