

# **LAMP2 (PT0249R) PT™ Rabbit mAb**

CatalogNo: YM8157 Recombinant R

## **Key Features**

**Host Species** 

Rabbit

MW
• 45kD (Calculated)

120kD (Observed)

Reactivity

Human, Mouse

Isotype

IgG,Kappa

**Applications** 

• WB,IHC,IF,ELISA

#### **Recommended Dilution Ratios**

IHC 1:200-1:1000 WB 1:1000-1:5000 IF 1:200-1:1000

ELISA 1:5000-1:20000

#### Storage

Storage\* -15°C to -25°C/1 year(Do not lower than -25°C)

**Formulation** PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA

#### **Basic Information**

**Clonality** Monoclonal

Clone Number PT0249R

## Immunogen Information

**Specificity** Endogenous

## **Target Information**

Gene name

LAMP2

**Protein Name** 

Lysosome-associated membrane glycoprotein 2

Organism	Gene ID	UniProt ID	
Human	<u>3920;</u>	<u>P13473;</u>	
Mouse		<u>P17047</u> ;	

#### Cellular Localization

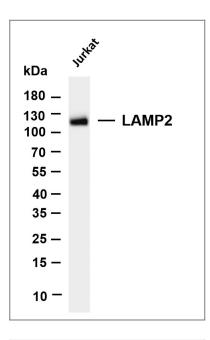
Membranous

**Tissue specificity** 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).

**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 is highly expressed in placenta, lung and liver, less in kidney and pancreas, low in brain and skeletal muscle. Isoform LAMP-2B is highly expressed in skeletal muscle, less in brain, placenta, lung, kidney and pancreas, very low in liver.,

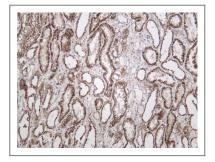
#### **I** Validation Data



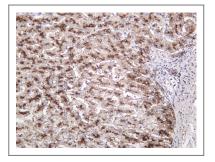
Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-LAMP2 antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: Jurkat Predicted band size: 45kDa Observed band size: 120kDa



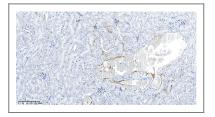
Mouse liver was stained with Anti-LAMP2 rabbit antibody



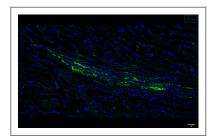
Human kidney was stained with Anti-LAMP2 rabbit antibody



Human liver was stained with Anti-LAMP2 rabbit antibody



Rat kidney was stained with anti-LAMP2 Rabbit antibody



Rat kidney was stained with anti-LAMP2 Rabbit antibody

#### | Contact information

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Please scan the QR code to access additional product information:

LAMP2 (PT0249R)

PT™ Rabbit mAb

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

Antibody | ELISA Kits | Protein | Reagents