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## LAMP2 (ANT0049R) Rabbit mAb

CatalogNo: ANT8157 Recombinant R

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

Quantity: 100 ug/vial

Host Species Reactivity Applications

Rabbit
 Human, Mouse
 WB, IHC, IF, IP, ELISA

MW Isotype

• 45kD (Calculated) • IgG,Kappa

120kD (Observed)

### **Recommended Dilution Ratios**

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,

# Storage

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

#### **Basic Information**

**Clonality** Monoclonal

Clone Number ANT0049R

# **Target Information**

#### Immunogen Information Specificity

#### Endogenous

Gene name LAMP2

**Protein Name** Lysosome-associated membrane glycoprotein 2

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

Cellular Localization Membranous

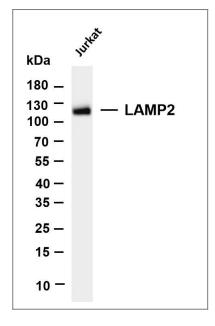
Tissue specificity Isoform LAMP-2A is highly expressed in placenta, lung and liver, less in kidney and

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., ANTM: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.,



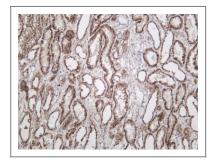
## **Validation**

#### **Data**

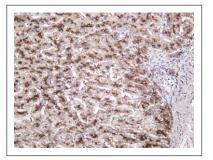
Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-LAMP2 (ANT0049R) antibody. The HRPconjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: Jurkat Predicted band size: 45kDa Observed band size: 120kDa



Mouse liver was stained with Anti-LAMP2 (ANT0049R) rabbit antibody



Human kidney was stained with Anti-LAMP2 (ANT0049R) rabbit antibody



Human liver was stained with Anti-LAMP2 (ANT0049R) rabbit antibody

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Contact Antagene Inc Tel 1-866-964-2589 Email: info@antageneinc.com