

Lrp2 / Megalin Rabbit mAb

货号: B28465

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Monoclonal
预测反应	
应用	WB
推荐浓度	WB: 1:500 - 1:2000
理论分子量	521kDa
实测分子量	280kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	Mouse lung
细胞定位	Membrane,Single-pass type I membrane protein,coated pit
纯化	Affinity purification

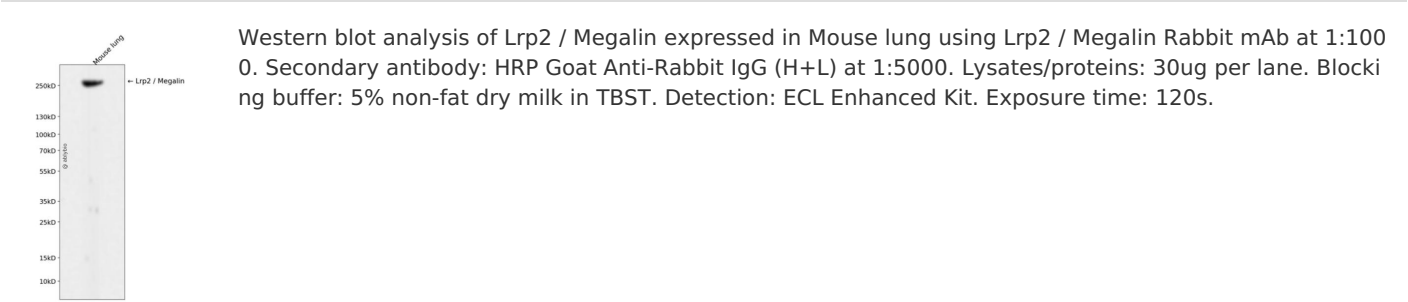
抗原信息

抗原信息	Recombinant fusion protein corresponding to Human Lrp2 / Megalin.
序列	FHYRRTGSLLPALPKLPSSLVKPSENGVTFRSGADLNMDIGVSGFGPETAI DRSMAMSEDFVMEMGKQPIIFENPM YSARDSAVKVVQPIQVTVSENV DNKNYGSPINPSEIVPETNPTSPAADGTQVTKWNLFKRKSKQTTNFENPIYAQMENEQK ESVAATPPPSPLPAKPKPSRRDPTPTYSATEDTFKDTANLVKEDSEV

靶点信息

研究背景	The protein encoded by this gene, low density lipoprotein-related protein 2 (LRP2) or megalin, is a multi-ligand endocytic receptor that is expressed in many different tissues but primarily in absorptive epithelial tissues such as the kidney. This glycoprotein has a large amino-terminal extracellular domain, a single transmembrane domain, and a short carboxy-terminal cytoplasmic tail. The extracellular ligand-binding domains bind diverse macromolecules including albumin, apolipoproteins B and E, and lipoprotein lipase. The LRP2 protein is critical for the reuptake of numerous ligands, including lipoproteins, sterols, vitamin-binding proteins, and hormones. This protein also has a role in cell-signaling; extracellular ligands include parathyroid hormones and the morphogen sonic hedgehog while cytosolic ligands include MAP kinase scaffold proteins and JNK interacting proteins. Recycling of this membrane receptor is regulated by phosphorylation of its cytoplasmic domain. Mutations in this gene cause Donnai-Barrow syndrome (DBS) and facio-otico-acoustico-renal syndrome (FOAR).
基因ID	4036
基因名	LRP2
Swiss	P98164
别名	LRP2;DBS;GP330

产品验证



实验步骤

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