

Recombinant Human LAMP1/CD107a Protein (His Tag)



Catalog Number:PKSH032682

Note: Centrifuge before opening to ensure complete recovery of vial contents.

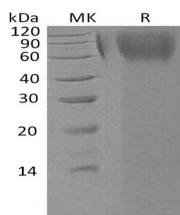
Description

Synonyms	Lysosome-Associated Membrane Glycoprotein 1;LAMP-1;Lysosome-Associated Membrane Protein 1;CD107 Antigen-Like Family Member A;CD107a;LAMP1;CD107a;LAMPA;LGP120
Species	Human
Expression Host	HEK293 Cells
Sequence	Ala29-Met382
Accession	P11279
Calculated Molecular Weight	39.4 kDa
Observed molecular weight	55-120 kDa
Tag	C-His

Properties

Purity	> 95 % as determined by reducing SDS-PAGE.
Endotoxin	< 1.0 EU per µg of the protein as determined by the LAL method.
Storage	Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.
Shipping	This product is provided as lyophilized powder which is shipped with ice packs.
Formulation	Lyophilized from a 0.2 µm filtered solution of 20mM PB, 150mM NaCl, pH 7.2. Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as protectants before lyophilization.
Reconstitution	Please refer to the specific buffer information in the printed manual for detailed information.

Data



> 95 % as determined by reducing SDS-PAGE.

Background

Lysosome-Associated Membrane Glycoprotein 1 (LAMP1) is a single-pass type I membrane protein belonging to the LAMP family. LAMP1 is expressed largely in the endosome-lysosome membranes of cells. It shuttles between lysosomes; endosomes; and the plasma membrane. LAMP1 functions to present carbohydrate ligands to selectins and it has also been implicated in tumor cell metastasis. It has been proposed LAMP1 can be used as a therapeutic agent for certain cancers; as well as a marker for lysosomal storage disorders and degranulation on lymphocytes such as CD8+ and NK cells. Cell surface LAMP1 and LAMP2 have been shown to promote adhesion of human peripheral blood mononuclear

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cells(PBMC) to vascular endothelium; therefore they are possibly involved in the adhesion of PBMCs to the site of inflammation.

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