

Synonym

LYPD1, Ly6/PLAUR domain-containing protein 1, FLJ41033, LY6/PLAUR domain containing 1, LYPDC1, PHTS, Putative HeLa tumor suppressor

Source

Human LYPD1 Protein, Mouse IgG2a Fc Tag (LY1-H5258) is expressed from human 293 cells (HEK293). It contains AA Leu 21 - Gly 115 (Accession # [Q8N2G4-1](#)).

Predicted N-terminus: Leu 21

Molecular Characterization

LYPD1(Leu 21 - Gly 115) Q8N2G4-1	mFc(Glu 98 - Lys 330) P01863
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This protein carries a mouse IgG2a Fc tag at the C-terminus

The protein has a calculated MW of 37.3 kDa. The protein migrates as 43-48 kDa under reducing (R) condition (SDS-PAGE) due to glycosylation.

Endotoxin

Less than 1.0 EU per μg by the LAL method.

Purity

>95% as determined by SDS-PAGE.

>90% as determined by SEC-MALS.

Formulation

Lyophilized from 0.22 μm filtered solution in PBS, pH7.4 with trehalose as protectant.

Contact us for customized product form or formulation.

Reconstitution

Please see Certificate of Analysis for specific instructions.

For best performance, we strongly recommend you to follow the reconstitution protocol provided in the CoA.

Storage

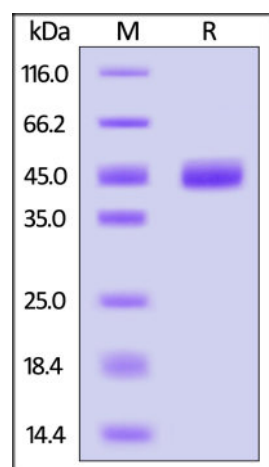
For long term storage, the product should be stored at lyophilized state at -20°C or lower.

Please avoid repeated freeze-thaw cycles.

This product is stable after storage at:

- -20°C to -70°C for 12 months in lyophilized state;
- -70°C for 3 months under sterile conditions after reconstitution.

SDS-PAGE



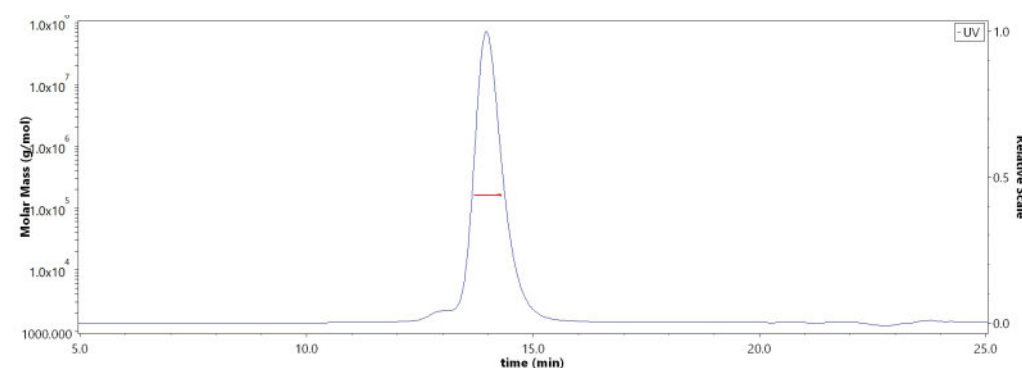
Human LYPD1 Protein, Mouse IgG2a Fc Tag on SDS-PAGE under reducing (R) condition. The gel was stained with Coomassie Blue. The purity of the protein is greater than 95%.

Bioactivity-ELISA

LYPD1 ELISA

Immobilized Human LYPD1 Protein, Mouse IgG2a Fc Tag (Cat. No. LY1-H5258) at 1 $\mu\text{g}/\text{mL}$ (100 $\mu\text{L}/\text{well}$) can bind Anti-LYPD1 antibody with a linear range of 0.02-0.625 $\mu\text{g}/\text{mL}$ (QC tested).

SEC-MALS



The purity of Human LYPD1 Protein, Mouse IgG2a Fc Tag (Cat. No. LY1-H5258) is more than 90% and the molecular weight of this protein is around 155-175 kDa verified by SEC-MALS.

[Report](#)

Background

Believed to act as a modulator of nicotinic acetylcholine receptors (nAChRs) activity. In vitro increases receptor desensitization and decreases affinity for ACh of alpha-4:beta-2-containing nAChRs. May play a role in the intracellular trafficking of alpha-4:beta-2 and alpha-7-containing nAChRs and may inhibit their expression at the cell surface. May be involved in the control of anxiety.

Clinical and Translational Updates

Please contact us via TechSupport@acrobiosystems.com if you have any question on this product.