

Recombinant Human Apolipoprotein A-I/ApoA1 Protein (His Tag)



Catalog Number:PKSH031486

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

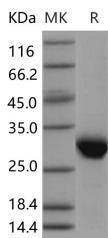
Description

Synonyms	Apolipoprotein A-I;Apo-AI;ApoA-I;Apolipoprotein A1;APOA1
Species	Human
Expression Host	E.coli
Sequence	Asp25-Gln267
Accession	P02647
Calculated Molecular Weight	30.7 kDa
Observed molecular weight	27-31 kDa
Tag	N-His

Properties

Purity	> 95 % as determined by reducing SDS-PAGE.
Endotoxin	Please contact us for more information.
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 sterile PBS, 10% Glycerol, pH 7.4 Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as protectants before lyophilization. Please refer to the specific buffer information in the printed manual.
Reconstitution	Please refer to the printed manual for detailed information.

Data



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Background

Apolipoprotein A1 (APOA1) is a member of the apolipoprotein family whose members are proteins bind with lipids and form lipoproteins to translate these oil-soluble lipids such as fat and cholesterol through lymphatic and circulatory system. APOA1 is the main component of high density lipoprotein (HDL) in plasma and is involved in the esterification of cholesterol as a cofactor of lecithin-cholesterol acyltransferase (LCAT) which is responsible for the formation of most plasma cholesterol esters; and thus play a major role in cholesterol efflux from peripheral cells. As a major component of the HDL complex; APOA1 helps to clear cholesterol from arteries. APOA1 is also characterized as a prostacyclin stabilizing factor; and thus may have an anticoagulant effect. Defects in encoding gene may result in HDL deficiencies; including Tangier disease; and with systemic non-neuropathic amyloidosis. Men carrying a mutation may develop

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premature coronary artery disease.

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