

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

Catalog No. 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

AccessionP02647Calculated Molecular Weight30.7 kDaObserved molecular weight27-31 kDaTagN-His

**Bioactivity** Not validated for activity

# **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

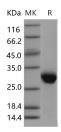
Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as

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



> 95 % as determined by reducing SDS-PAGE.

# **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.

#### For Research Use Only

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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 cholesteryl 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 anticlotting 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 premature coronary artery disease.

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