abcam

Product datasheet

Recombinant Human Apolipoprotein A V/APOA5 ab113861

1 Image

Description

Product name Recombinant Human Apolipoprotein A V/APOA5

Purity > 70 % SDS-PAGE.

Purity determined by densitometric image analysis

Endotoxin level < 1.000 Eu/µg
Expression system HEK 293 cells

Accession Q6Q788

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence RKGFWDYFSQ TSGDKGRVEQ IHQQKMAREP

ATLKDSLEQD LNNMNKFLEK LRPLSGSEAP RLPQDPVGMR RQLQEELEEV KARLQPYMAE AHELVGWNLE GLRQQLKPYT MDLMEQVALR VQELQEQLRV VGEDTKAQLL GGVDEAWALL QGLQSRVVHH TGRFKELFHP YAESLVSGIG RHVQELHRSV APHAPASPAR LSRCVQVLSR KLTLKAKALH ARIQQNLDQL REELSRAFAG TGTEEGAGPD PQMLSEEVRQ RLQAFRQDTY LQIAAFTRAI DQETEEVQQQ LAPPPPGHSA FAPEFQQTDS GKVLSKLQAR LDDLWEDITH

SLHDQGHSHL GDPAAADYKD DDDK

Predicted molecular weight 40 kDa including tags

Amino acids 24 to 366

Tags DDDDK tag C-Terminus

Specifications

Our <u>Abpromise guarantee</u> covers the use of ab113861 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

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Applications SDS-PAGE

Western blot

ELISA

Form Lyophilized

Additional notes This product was previously labelled as Apolipoprotein A V

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot. Store at -80°C. Avoid freeze / thaw cycle.

Constituents: 0.24% Tris, 0.29% Sodium chloride

Reconstitution Add deionized water (200 μl) to prepare a working stock solution of 0.5 mg/mL

General Info

Function Minor apolipoprotein mainly associated with HDL and to a lesser extent with VLDL. May also be

associated with chylomicrons. Important determinant of plasma triglyceride (TG) levels by both being a potent stimulator of apo-CII lipoprotein lipase (LPL) TG hydrolysis and a inhibitor of the hepatic VLDL-TG production rate (without affecting the VLDL-apoB production rate) (By similarity). Activates poorly lecithin:cholesterol acyltransferase (LCAT) and does not enhance

efflux of cholesterol from macrophages.

Tissue specificity Liver and plasma.

Involvement in diseaseDefects in APOA5 are a cause of susceptibility to familial hypertriglyceridemia (FHTR)

[MIM:145750].mFamilial hypertriglyceridemia is a common inherited disorder in which the concentration of very low density lipoprotein (VLDL) is elevated in the plasma. This leads to

increased risk of heart disease, obesity, and pancreatitis.

Defects in APOA5 are a cause of hyperlipoproteinemia type 5 (HLPP5) [MIM:144650]. HLPP5 is characterized by increased amounts of chylomicrons and very low density lipoprotein (VLDL) and decreased low density lipoprotein (LDL) and high density lipoprotein (HDL) in the plasma after a fast. Numerous conditions cause this phenotype, including insulin-dependent diabetes mellitus, contraceptive steroids, alcohol abuse, and glycogen storage disease type 1A (GSD1A)

[MIM:232200].

Sequence similaritiesBelongs to the apolipoprotein A1/A4/E family.

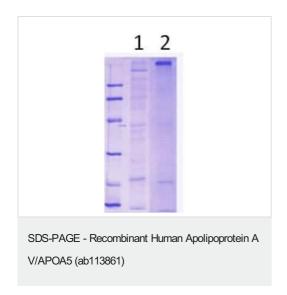
Post-translational

modifications

Phosphorylation sites are present in the extracelllular medium.

Cellular localization Secreted.

Images



12% SDS-PAGE separation of Human Apolipoprotein A V/APOA5 1.reduced and boiled sample, 5µg/lane 2.non-reduced and non-boiled sample, 5µg/lane

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