abcam

Product datasheet

Recombinant Human Apolipoprotein L1/APOL1 ab152886

1 Image

Description

Product name Recombinant Human Apolipoprotein L1/APOL1

Purity > 80 % Affinity purified.

Glutathione Sepharose 4 Fast Flow

Expression system Wheat germ
Accession <u>014791</u>

Protein length Protein fragment

Animal free No.

Nature Recombinant

Species Human

Sequence MEGAALLRVSVLCIWMSALFLGVGVRAEEAGARVQQNVP

SGTDTGDPQSK

PLGDWAAGTMDPESSIFIEDAIKYFKEKVSTQNLLLLLTDN

EAWNGFVAA

AELPRNEADELRKALDNLARQMIMKDKNWHDKGQQYRN

WFLKEFPRLKSE

LEDNIRRLRALADGVQKVHKGTTIANVVSGSLSISSGILTLV

GMGLAPFT

EGGSLVLLEPGMELGITAALTGITSSTMDYGKKWWTQA

Predicted molecular weight 52 kDa including tags

Amino acids 1 to 238

Specifications

Our **Abpromise guarantee** covers the use of **ab152886** in the following tested applications.

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

Applications ELISA

SDS-PAGE

Western blot

Form Liquid

Additional notes Protein concentration is above or equal to 0.05 μg/μl

1

Preparation and Storage

Stability and Storage Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.31% Glutathione, 0.79% Tris HCI

General Info

Function May play a role in lipid exchange and transport throughout the body. May participate in reverse

cholesterol transport from peripheral cells to the liver.

Tissue specificity Plasma. Found on APOA-I-containing high density lipoprotein (HDL3). Expressed in pancreas,

lung, prostate, liver, placenta and spleen.

Involvement in diseaseDefects in APOL1 are the cause of focal segmental glomerulosclerosis type 4 (FSGS4)

[MIM:612551]. It is a renal pathology defined by the presence of segmental sclerosis in glomeruli and resulting in proteinuria, reduced glomerular filtration rate and edema. Renal insufficiency often progresses to end-stage renal disease, a highly morbid state requiring either dialysis therapy or

kidney transplantation.

Sequence similarities Belongs to the apolipoprotein L family.

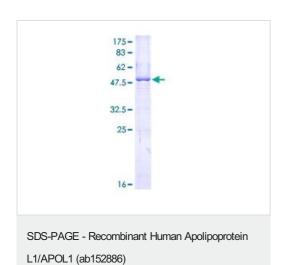
Post-translational

modifications

Phosphorylation sites are present in the extracelllular medium.

Cellular localization Secreted.

Images



12.5% SDS-PAGE analysis of ab152886 stained with Coomassie Blue

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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