# abcam

## Product datasheet

## Recombinant Human Apolipoprotein A I ab50239

4 References 画像数 1

製品の詳細

製品名 Recombinant Human Apolipoprotein A I

精製度 > 95 % SDS-PAGE.

ab50239 purity is greater than 97% by SDS-PAGE gel and HPLC analyses.

**エンドトキシン・レベル** < 0.100 Ew/μg

**発現系** Escherichia coli

アクセッション番号 <u>P02647</u>

タンパク質長 Full length protein

Animal free No

**由来** Recombinant

生物種 Human

配列 MDEPPQSPWD RVKDLATVYV DVLKDSGRDY

VSQFEGSALG KQLNLKLLDN WDSVTSTFSK LREQLGPVTQ EFWDNLEKET EGLRQEMSKD LEEVKAKVQP YLDDFQKKWQ EEMELYRQKV EPLRAELQEG ARQKLHELQE KLSPLGEEMR DRARAHVDAL RTHLAPYSDE LRQRLAARLE ALKENGGARL AEYHAKATEH LSTLSEKAKP

ALEDLRQGLL PVLESFKVSF LSALEEYTKK LNTQ

**予測される分子量** 28 kDa

特性

Our **Abpromise guarantee** covers the use of **ab50239** in the following tested applications.

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

アプリケーション SDS-PAGE

Sandwich ELISA

製品の状態 Lyophilized

備考 Molecular Weight: 28.2 kDa

前処理および保存

#### 保存方法および安定性

#### 再構成

Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Initially reconstitute in water to 0.1-1.0 mg/ml. Store at 2°C to 8°C for up to 1 week or prepare for extended storage. After initial reconstitution, further dilute in a buffer containing a carrier protein or stabilizer (e.g. 0.1% BSA). Store working aliquots at -20°C to -80°C.

#### 関連情報

#### 機能

Participates in the reverse transport of cholesterol from tissues to the liver for excretion by promoting cholesterol efflux from tissues and by acting as a cofactor for the lecithin cholesterol acyltransferase (LCAT). As part of the SPAP complex, activates spermatozoa motility.

#### 組織特異性

Major protein of plasma HDL, also found in chylomicrons. Synthesized in the liver and small intestine.

#### 関連疾患

Defects in APOA1 are a cause of high density lipoprotein deficiency type 2 (HDLD2) [MIM:604091]; also known as familial hypoalphalipoproteinemia (FHA). Inheritance is autosomal dominant.

Defects in APOA1 are a cause of the low HDL levels observed in high density lipoprotein deficiency type 1 (HDLD1) [MIM:205400]; also known as analphalipoproteinemia or Tangier disease (TGD). HDLD1 is a recessive disorder characterized by the absence of plasma HDL, accumulation of cholesteryl esters, premature coronary artery disease, hepatosplenomegaly, recurrent peripheral neuropathy and progressive muscle wasting and weakness. In HDLD1 patients, ApoA-I fails to associate with HDL probably because of the faulty conversion of pro-ApoA-I molecules into mature chains, either due to a defect in the converting enzyme activity or a specific structural defect in Tangier ApoA-I.

Defects in APOA1 are the cause of amyloid polyneuropathy-nephropathy lowa type (AMYLIOWA) [MIM:107680]; also known as amyloidosis van Allen type or familial amyloid polyneuropathy type III. AMYLIOWA is a hereditary generalized amyloidosis due to deposition of amyloid mainly constituted by apolipoprotein A1. The clinical picture is dominated by neuropathy in the early stages of the disease and nephropathy late in the course. Death is due in most cases to renal amyloidosis. Severe peptic ulcer disease can occurr in some and hearing loss is frequent. Cataracts is present in several, but vitreous opacities are not observed.

Defects in APOA1 are a cause of amyloidosis type 8 (AMYL8) [MIM:105200]; also known as systemic non-neuropathic amyloidosis or Ostertag-type amyloidosis. AMYL8 is a hereditary generalized amyloidosis due to deposition of apolipoprotein A1, fibrinogen and lysozyme amyloids. Viscera are particularly affected. There is no involvement of the nervous system. Clinical features include renal amyloidosis resulting in nephrotic syndrome, arterial hypertension, hepatosplenomegaly, cholestasis, petechial skin rash.

#### 配列類似性

Belongs to the apolipoprotein A1/A4/E family.

### 翻訳後修飾

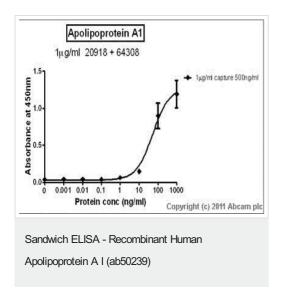
Palmitoylated.

Phosphorylation sites are present in the extracelllular medium.

#### 細胞内局在

Secreted.

#### 画像



Standard curve for Apolipoprotein A I (Analyte: ab50239); dilution range 1pg/ml to 1µg/ml using Capture Antibody **ab20918** at 1µg/ml and Detector Antibody **ab64308** at 0.5µg/ml.

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

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