abcam

Product datasheet

Anti-Apolipoprotein A I antibody ab64308

★★★★★ 3 Abreviews 9 References 画像数 3

製品の概要

免疫原

製品名 Anti-Apolipoprotein A I antibody

製品の詳細 Rabbit polyclonal to Apolipoprotein A I

由来種 Rabbit

アプリケーション 適用あり: Sandwich ELISA, WB, ICC/IF

種交差性 交差種: Human

交差が予測される動物種: Baboon 4

Synthetic peptide conjugated to KLH derived from within residues 200 to the C-terminus of

Human Apolipoprotein A I.Immunogen の所有権に関して(Peptide available as ab66674.)

ポジティブ・コントロール Recombinant Human Apolipoprotein A I (ab50239) can be used as a positive control in WB. This

antibody gave a positive signal in the following Human Tissue Lysates: Testis, Ovary, Lung,

Thymus ICC/IF: HepG2 cells

特記事項
The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide Constituents: PBS, 1% BSA

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

1

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab64308の使用に適用されますアプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
Sandwich ELISA		Use a concentration of 0.5 μ g/ml. For sandwich ELISA, use this antibody as Detection at 0.5 μ g/ml with <u>ab20918</u> as Capture.
WB	★★★★☆ (2)	Use a concentration of 1 µg/ml. Detects a band of approximately 27 kDa (predicted molecular weight: 31 kDa).
ICC/IF		Use a concentration of 5 µg/ml.

ターゲット情報

機能

組織特異性

関連疾患

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.

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

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.

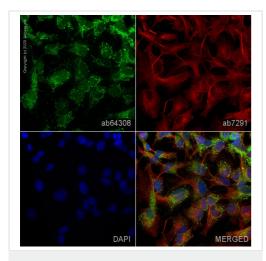
配列類似性 Belongs to the apolipoprotein A1/A4/E family.

翻訳後修飾 Palmitoylated.

Phosphorylation sites are present in the extracelllular medium.

細胞内局在 Secreted.

画像



Immunocytochemistry/ Immunofluorescence - Anti-Apolipoprotein A I antibody (ab64308)

ab64308 staining Apolipoprotein A I in HepG2 cells. The cells were fixed with 4% paraformaldehyde (10 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab64308 at 5µg/ml and ab7291, Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control. Cells were then incubated with ab150081, Goat polyclonal Secondary Antibody to Rabbit lgG - H&L (Alexa Fluor® 488), pre-adsorbed at 1/1000 dilution (shown in green) and ab150120, Goat polyclonal Secondary Antibody to Mouse lgG - H&L (Alexa Fluor® 594), pre-adsorbed at 1/1000 dilution (shown in pseudocolour red). Nuclear DNA was labelled with DAPI (shown in



Western blot - Anti-Apolipoprotein A I antibody (ab64308)

All lanes: Anti-Apolipoprotein A lantibody (ab64308) at 1 µg/ml

Lane 1 : Human testis tissue lysate - total protein (<u>ab30257</u>)

Lane 2 : Human ovary tissue lysate - total protein (<u>ab30222</u>)

Lane 4: Human thymus tissue lysate - total protein (ab30146)

Lane 3: Lung (Human) Tissue Lysate - adult normal tissue

Lysates/proteins at 10 µg per lane.

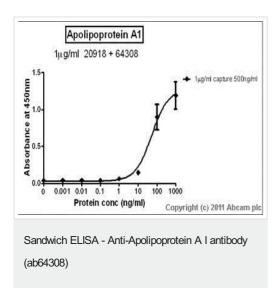
Secondary

blue).

 $\begin{tabular}{ll} \textbf{All lanes:} Goat polyclonal to Rabbit lgG - H\&L - Pre-Adsorbed (HRP) at 1/3000 dilution \end{tabular}$

Performed under reducing conditions.

Predicted band size: 31 kDa **Observed band size:** 27 kDa



Standard Curve for Apolipoprotein A I (Analyte: <u>ab50239</u>) dilution range 1pg/ml to 1ug/ml using Capture Antibody Mouse monoclonal [1409] to Apolipoprotein A I (<u>ab20918</u>) at 1ug/ml and Detector Antibody Rabbit polyclonal to Apolipoprotein A I (ab64308) at 0.5ug/ml

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

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