SANTA CRUZ BIOTECHNOLOGY, INC.

apoA-IV (G-8): sc-374543



BACKGROUND

Apolipoproteins are protein components of plasma lipoproteins. The human apoA-I gene encodes a single chain, 243 amino acid protein which promotes cholesterol efflux from tissues to the liver for excretion. Apolipoprotein A-I is the major protein component of high density lipoprotein (HDL) in the plasma. It can function as a cofactor for lecithin cholesterolacyltransferase (LCAT), which is responsible for the formation of most plasma cholesteryl esters. The human apoA-II gene encodes the second most abundant protein of HDL particles, where it influences plasma levels of free fatty acids (FFA). The human apoA-IV gene encodes a 396 amino acid preprotein, which after proteolytic processing is secreted from the intestine in association with chylomicron particles. ApoA-IV is a potent activator of LCAT *in vitro*. The human apoA-V gene encodes a 366 amino acid protein that is believed to be an important determinant of plasma triglyceride levels.

CHROMOSOMAL LOCATION

Genetic locus: APOA4 (human) mapping to 11q23.3.

SOURCE

apoA-IV (G-8) is a mouse monoclonal antibody raised against a peptide mapping near the C-terminus of apoA-IV of human origin.

PRODUCT

Each vial contains 200 μg IgG_1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

apoA-IV (G-8) is available conjugated to agarose (sc-374543 AC), 500 μ g/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-374543 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-374543 PE), fluorescein (sc-374543 FITC), Alexa Fluor[®] 488 (sc-374543 AF488), Alexa Fluor[®] 546 (sc-374543 AF546), Alexa Fluor[®] 594 (sc-374543 AF594) or Alexa Fluor[®] 647 (sc-374543 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-374543 AF680) or Alexa Fluor[®] 790 (sc-374543 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

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APPLICATIONS

apoA-IV (G-8) is recommended for detection of apoA-IV of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for apoA-IV siRNA (h): sc-41178, apoA-IV shRNA Plasmid (h): sc-41178-SH and apoA-IV shRNA (h) Lentiviral Particles: sc-41178-V.

Molecular Weight of apoA-IV: 46 kDa.

Positive Controls: apoA-IV (h): 293T Lysate: sc-373359, Hep G2 cell lysate: sc-2227 or HeLa whole cell lysate: sc-2200.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA



apoA-IV (G-8): sc-374543. Western blot analysis of apoA-IV expression in non-transfected: sc-117752 (A) and human apoA-IV transfected: sc-373359 (B) 293T whole cell lysates.



apoA-IV (G-8): sc-374543. Immunoperoxidase staining of formalin fixed, paraffin-embedded human duodenum tissue showing cytoplasmic staining of glandular cells (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human kidney tissue showing apical membrane staining of cells in tubules (B).

SELECT PRODUCT CITATIONS

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- 3. Xu, X.R., et al. 2018. Apolipoprotein A-IV binds α IIb β 3 Integrin and inhibits thrombosis. Nat. Commun. 9: 3608.
- Choi, J.W., et al. 2020. Proteome analysis of human natural killer cell derived extracellular vesicles for identification of anticancer effectors. Molecules 25: 5216.
- 5. Park, S.R., et al. 2021. Holistic characterization of single-hepatocyte transcriptome responses to high-fat diet. Am. J. Physiol. Endocrinol. Metab. 320: E244-E258.
- Zhang, X., et al. 2022. Identification of serum biomarkers in patients with Alzheimer's disease by 2D-DIGE proteomics. Gerontology 68: 686-698.
- Liu, X.H., et al. 2022. Apolipoprotein A-IV reduced metabolic inflammation in white adipose tissue by inhibiting IKK and JNK signaling in adipocytes. Mol. Cell. Endocrinol. 559: 111813.
- Gruver, A.M., et al. 2023. Pathologist-trained machine learning classifiers developed to quantitate celiac disease features differentiate endoscopic biopsies according to modified marsh score and dietary intervention response. Diagn. Pathol. 18: 122.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.