

# apoA-I (12C8): sc-80551

## 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.

## REFERENCES

1. Duriez, P. and Fruchart, J.C. 1999. High-density lipoprotein subclasses and apolipoprotein A-I. Clin. Chim. Acta 286: 97-114.
2. Maezawa, I., et al. 2004. apoE isoforms and apoA-I protect from amyloid precursor protein carboxy-terminal fragment-associated cytotoxicity. J. Neurochem. 91: 1312-1321.
3. Maejima, T., et al. 2004. Effect of pitavastatin on apoA-I production in Hep G2 cell. Biochem. Biophys. Res. Commun. 324: 835-839.

## CHROMOSOMAL LOCATION

Genetic locus: APOA1 (human) mapping to 11q23.3; ApoA1 (mouse) mapping to 9 A5.2.

## SOURCE

apoA-I (12C8) is a mouse monoclonal antibody raised against full length native apoA-I of human origin.

## PRODUCT

Each vial contains 100 µg IgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

## APPLICATIONS

apoA-I (12C8) is recommended for detection of apoA-I of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)] and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000); non cross-reactive with apolipoprotein B (apoB).

Suitable for use as control antibody for apoA-I siRNA (h): sc-41177, apoA-I siRNA (m): sc-63361, apoA-I shRNA Plasmid (h): sc-41177-SH, apoA-I shRNA Plasmid (m): sc-63361-SH, apoA-I shRNA (h) Lentiviral Particles: sc-41177-V and apoA-I shRNA (m) Lentiviral Particles: sc-63361-V.

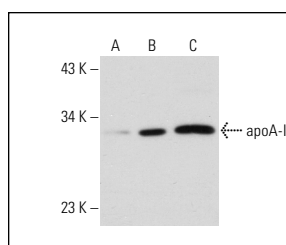
Molecular Weight of apoA-I: 28 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200, apoA-I (h): 293T Lysate: sc-111827 or apoA-I (m): 293T Lysate: sc-118477.

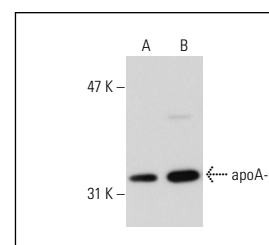
## RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).

## DATA



apoA-I (12C8): sc-80551. Western blot analysis of apoA-I expression in non-transfected 293T: sc-117752 (A), mouse apoA-I transfected 293T: sc-118477 (B) and HeLa (C) whole cell lysates.



apoA-I (12C8): sc-80551. Western blot analysis of apoA-I expression in non-transfected: sc-117752 (A) and human apoA-I transfected: sc-111827 (B) 293T whole cell lysates.

## SELECT PRODUCT CITATIONS

1. Chong, P.K., et al. 2010. Reduced plasma apoA-I level is associated with gastric tumor growth in MKN45 mouse xenograft model. J. Proteomics 73: 1632-1640.
2. Yu, Y., et al. 2016. The binding capability of plasma phospholipid transfer protein, but not HDL pool size, is critical to repress LPS induced inflammation. Sci. Rep. 6: 20845.
3. Lipps, C., et al. 2019. Non-invasive approach for evaluation of pulmonary hypertension using extracellular vesicle-associated small non-coding RNA. Biomolecules 9 pii: E666.
4. Brahmer, A., et al. 2019. Platelets, endothelial cells and leukocytes contribute to the exercise-triggered release of extracellular vesicles into the circulation. J. Extracell. Vesicles 8: 1615820.

## STORAGE

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

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.