

Product datasheet for BM785

OriGene Technologies, Inc.

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Apolipoprotein A I (APOA1) Mouse Monoclonal Antibody [Clone ID: G2]

Product data:

Product Type: Primary Antibodies

Clone Name: G2

Applications: ELISA

Recommended Dilution: ELISA (1/2.500-1/10.000). This antibody is suitable for coating microtitre plates in a Sandwich

ELISA using catalogue number BP912HRP for detection.

Western Blot (1/250-1/1000).

Immunohistochemistry on Frozen Sections (1/20-1/80).

Reactivity: Human
Host: Mouse
Isotype: IgG1

Clonality: Monoclonal

Immunogen: Native Human Apolipoprotein A1 from Human plasma.

Specificity: Reacts with both free Apolipoprotein A1 and HDL bearing Apo-A1. Does not cross-react with

ApoE, ApoB or Albumin.

Formulation: 0.01M Sodium Phosphate, 0.01M Sodium Borate, 0.15M Sodium Chloride with 1% Mannitol

and 1% Dextran.
State: Purified

State: Lyophilized purified IgG fraction

Reconstitution Method: Restore with 1.0 ml distilled water.

Care should be taken during reconstitution as the protein may appear as a film at the bottom

of the vial. We recommend that the vial is gently mixed after reconstitution.

Purification: Affinity Chromatography on Protein A

Conjugation: Unconjugated

Storage: Store the lyophilized antibody at 2-8°C and after reconstitution at -20°C.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: apolipoprotein A1





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Database Link: Entrez Gene 335 Human

P02647

Background: Apolipoprotein A I 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. Synthesized in the liver and small intestine, it consists of two identical chains of 77 amino acids; an 18 amino acid signal peptide is removed co-translationally and a 6 amino acid propeptide is cleaved post-translationally. Apolipoprotein A I is a cofactor for lecithin cholesterolacyltransferase (LCAT) which is responsible for the formation of most plasma cholesteryl esters. Defects in the Apolipoprotein A I gene are associated with HDL deficiency

and Tangier disease.

The therapeutic potential of apoA-I has been recently assessed in patients with acute coronary syndromes, using a recombinant form of a naturally occurring variant of apoA-I. The availability of recombinant normal apoA-I should facilitate further investigation into the

potential usefulness of apoA-I in preventing atherosclerotic vascular diseases.

Synonyms: APOA1, ApoA-I, Apo-AI, ApoAI