



# Rabbit Anti-Human Apolipoprotein A-I/APOA1 monoclonal antibody, clone S142 (CABT-ZB605)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Specificity</b>	It reacts with Human Apolipoprotein A-I/APOA1
<b>Target</b>	APOA1
<b>Immunogen</b>	Recombinant Human Apolipoprotein A-I/ApoA1 Protein
<b>Isotype</b>	IgG2b
<b>Source/Host</b>	Rabbit
<b>Species Reactivity</b>	Human
<b>Clone</b>	S142
<b>Purification</b>	Protein A purified
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	ELISA(cap) This antibody will detect Apolipoprotein A-I/APOA1 in antibody pair set. [ABPR-ZB183]
<b>Preparation</b>	This antibody was obtained from a rabbit immunized with purified, recombinant Human Apolipoprotein A-I / ApoA1.
<b>Format</b>	Purified, Liquid
<b>Concentration</b>	Lot specific
<b>Size</b>	50 µL, 100 µL, 1 mL

Buffer	PBS
Preservative	None
Storage	This antibody can be stored at 2°C-8°C for one month without detectable loss of activity. Antibody products are stable for twelve months from date of receipt when stored at -20°C to -80°C. Preservative-Free. Avoid repeated freeze-thaw cycles.
Ship	Wet ice

## BACKGROUND

<b>Introduction</b>	Apolipoprotein A1 (APOA1) is a member of the apolipoprotein family whose members are proteins bind with lipids and form lipoproteins to translate these oil-soluble lipids such as fat and cholesterol through lymphatic and circulatory system. APOA1 is the main component of high density lipoprotein (HDL) in plasma and is involved in the esterification of cholesterol as a cofactor of lecithin-cholesterol acyltransferase (LCAT) which is responsible for the formation of most plasma cholesterol esters, and thus play a major role in cholesterol efflux from peripheral cells. As a major component of the HDL complex, APOA1 helps to clear cholesterol from arteries. APOA1 is also characterized as a prostacyclin stabilizing factor, and thus may have an anticoagulant effect. Defects in encoding gene may result in HDL deficiencies, including Tangier disease, and with systemic non-neuropathic amyloidosis. Men carrying a mutation may develop premature coronary artery disease.
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<b>Keywords</b>	APOA1; apolipoprotein A-I; apo-AI
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## GENE INFORMATION

<b>Synonyms</b>	APOA1; apolipoprotein A-I; apo-AI
<b>Entrez Gene ID</b>	<a href="#">335</a>
<b>UniProt ID</b>	<a href="#">P02647</a>