

**APOB Rabbit Polyclonal Antibody** 

## Catalog #: EAB13953

Host/Isotype	Clonality	Applications	MW (kDa)	Reactivity
Rabbit IgG	Polyclonal	WB	516	Human, Mouse, Rat

## **Applications Dilutions**

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

WB(Western Blotting)

1:500-2000

## **Product Information**

SpecificityAPOB Rabbit Polyclonal Antibody detects endogenous levels of APOB protein.PurificationAffinity purificationConcentration1mg/mlFormatLiquidFormulationIn PBS, pH 7.4, Containing 0.02% sodium azide, 0.5% BSA and 50% Glycerol.ShippingGel PackStorageStore at -20°C least 1 year from the date of shipment. Avoid repeated freeze/thaw cycles. Aliquots may be stored at +4°C for 1-2 weeks.UniProt IDP04114			
PurificationAffinity purificationConcentration1mg/mlFormatLiquidFormulationIn PBS, pH 7.4, Containing 0.02% sodium azide, 0.5% BSA and 50% Glycerol.ShippingGel PackStorageStore at -20°C least 1 year from the date of shipment. Avoid repeated freeze/thaw cycles. Aliquots may be stored at +4°C for 1-2 weeks.UniProt IDP04114	Conjugate	Unconjugate	
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Shipping   Gel Pack     Storage   Store at -20°C least 1 year from the date of shipment. Avoid repeated freeze/thaw cycles.     UniProt ID   P04114	Format	Liquid	
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	Storage		
Entrez-Gene ID <u>338</u>	UniProt ID	<u>P04114</u>	
	Entrez-Gene ID	<u>338</u>	

## **Product Description**

APOB is the main apolipoprotein of chylomicrons and low density lipoproteins (LDL), and is the ligand for the LDL receptor. It occurs in plasma as two main isoforms, apoB-48 and apoB-100: the former is synthesized exclusively in the gut and the latter in the liver. The intestinal and the hepatic forms of apoB are encoded by a single gene from a single, very long mRNA. The two isoforms share a common N-terminal sequence. The shorter apoB-48 protein is produced after RNA editing of the apoB-100 transcript at residue 2180 (CAA->UAA), resulting in the creation of a stop codon, and early translation termination. Mutations in this gene or its regulatory region cause hypobetalipoproteinemia, normotriglyceridemic hypobetalipoproteinemia, and hypercholesterolemia due to ligand-defective apoB, diseases affecting plasma cholesterol and apoB levels.

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