

Rabbit Anti-APOL5 antibody

SL12500R

Product Name:	APOL5
Chinese Name:	载LipoproteinL5抗体
Alias:	APOL 5; APOL V; APOL5; Apolipoprotein L5; Apolipoprotein L; Apolipoprotein L V; apoL-V; Apolipoprotein L 5; APOLV; OTTHUMP00000028773; APOL5 HUMAN.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800ICC=1:100-500IF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	47kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human APOL5/Apolipoprotein L5:301-400/433
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20 °C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	<u>PubMed</u>
Product Detail:	The apolipoprotein L gene family maps to a region on chromosome 22 and encodes six highly homologous proteins designated apoL-I, apoL-II, apoL-III, apoL-IV, apoL-V and apoL-VI, all of which function as components of plasma lipoproteins. ApoL-V (apolipoprotein L-V), also known as APOL5, is a 433 amino acid protein that localizes

to the cytoplasm and belongs to the apolipoprotein L family. Expressed in a variety of tissues including testis, stomach, uterus and skeletal muscle, apoL-V is thought to affect the movement of lipids in the cytoplasm and may allow the binding of lipids to organelles. Like other members of the apolipoprotein L family, apoL-V is thought to be involved in the development of schizophrenia.

Function:

Apolipoprotein L 5 is a member of the apolipoprotein L gene family. The encoded protein is found in the cytoplasm, where it may affect the movement of lipids or allow the binding of lipids to organelles.

Subcellular Location:

Cytoplasm (Probable).

Tissue Specificity:

Low level of expression; detected in uterus, testis, skeletal muscle and stomach.

Similarity:

Belongs to the apolipoprotein L family.

SWISS:

O9BWW9

Gene ID:

80831

Database links:

Entrez Gene: 80831Human

Omim: 607255Human

SwissProt: Q9BWW9Human

Important Note:

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.