

APOO Antibody (N-term)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP13324a

Specification

APOO Antibody (N-term) - Product Information

| | |
|-------------------|-----------------------------|
| Application | IHC-P, WB,E |
| Primary Accession | O9BUR5 |
| Other Accession | NP_077027.1 |
| Reactivity | Human |
| Host | Rabbit |
| Clonality | Polyclonal |
| Isotype | Rabbit IgG |
| Calculated MW | 22285 |
| Antigen Region | 9-38 |

APOO Antibody (N-term) - Additional Information

Gene ID 79135

Other Names

Apolipoprotein O, Protein FAM121B, APOO, FAM121B

Target/Specificity

This APOO antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 9-38 amino acids from the N-terminal region of human APOO.

Dilution

IHC-P~~1:10~50

WB~~1:1000

E~~Use at an assay dependent concentration.

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

APOO Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

APOO Antibody (N-term) - Protein Information

Name APOO

Function Component of the MICOS complex, a large protein complex of the mitochondrial inner membrane that plays crucial roles in the maintenance of crista junctions, inner membrane architecture, and formation of contact sites to the outer membrane. Plays a crucial role in crista junction formation and mitochondrial function (PubMed:[25764979](#)). Can promote cardiac lipotoxicity by enhancing mitochondrial respiration and fatty acid metabolism in cardiac myoblasts (PubMed:[24743151](#)). Promotes cholesterol efflux from macrophage cells. Detected in HDL, LDL and VLDL. Secreted by a microsomal triglyceride transfer protein (MTTP)-dependent mechanism, probably as a VLDL-associated protein that is subsequently transferred to HDL (PubMed:[16956892](#)).

Cellular Location

Mitochondrion inner membrane; Single-pass membrane protein. Secreted. Mitochondrion. Golgi apparatus membrane. Endoplasmic reticulum membrane. Note=Exists in three distinct forms: a glycosylated and secreted form, an ER/Golgi-resident form and a non- glycosylated mitochondrial form.

Tissue Location

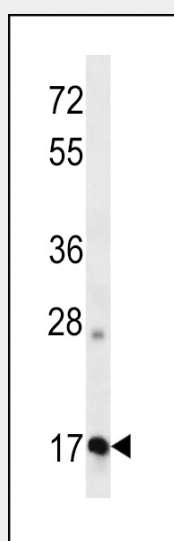
Expressed in all tissues examined. Up-regulated in diabetic heart.

APOO Antibody (N-term) - Protocols

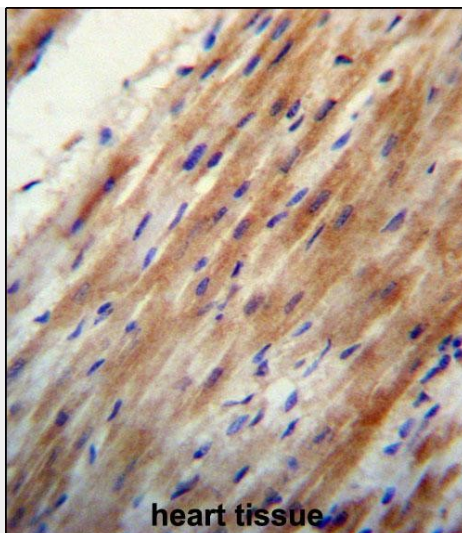
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

APOO Antibody (N-term) - Images



APOO Antibody (N-term) (Cat. #AP13324a) western blot analysis in K562 cell line lysates (35ug/lane). This demonstrates the APOO antibody detected the APOO protein (arrow).



APOO Antibody (N-term) (Cat. #AP13324a) immunohistochemistry analysis in formalin fixed and paraffin embedded human heart tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of APOO Antibody (N-term) for immunohistochemistry. Clinical relevance has not been evaluated.

APOO Antibody (N-term) - Background

This gene is a member of the apolipoprotein family. Members of this protein family are involved in the transport and metabolism of lipids. The encoded protein associates with HDL, LDL and VLDL lipoproteins and is characterized by chondroitin-sulfate glycosylation. This protein may be involved in preventing lipid accumulation in the myocardium in obese and diabetic patients. Alternative splicing results in multiple transcript variants. Pseudogenes of this gene are found on chromosomes 3, 4, 5, 12 and 16.

APOO Antibody (N-term) - References

Bailey, S.D., et al. Diabetes Care (2010) In press :
Talmud, P.J., et al. Am. J. Hum. Genet. 85(5):628-642(2009)
Chapuis, J., et al. Mol. Psychiatry 14(11):1004-1016(2009)
Lamant, M., et al. J. Biol. Chem. 281(47):36289-36302(2006)