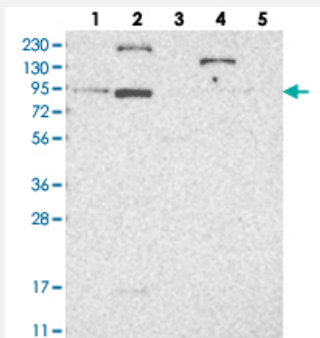


LCA5 polyclonal antibody

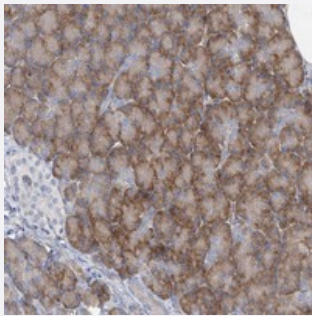
Catalog # PAB22243 Size 100 uL

Applications



Western Blot

Western blot analysis of Lane 1: RT-4, Lane 2: U-251 MG, Lane 3: Human Plasma, Lane 4: Liver, Lane 5: Tonsil with LCA5 polyclonal antibody (Cat # PAB22243).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human pancreas with LCA5 polyclonal antibody (Cat # PAB22243) shows moderate cytoplasmic positivity in exocrine glandular cells.

Specification

Product Description	Rabbit polyclonal antibody raised against recombinant LCA5.
Immunogen	Recombinant protein corresponding to amino acids of human LCA5.
Sequence	RKSQEKERATEKRVKDTESSELFRTKFSLQKLKEISEARHLPERDDLAKKLVSaelKLDDTERRIKE LSKNLELSTNSFQRQLLAERKRA
Host	Rabbit
Reactivity	Human, Mouse, Rat
Form	Liquid

Purification	Antigen affinity purification
Isotype	IgG
Recommend Usage	Immunohistochemistry (1:50-1:200) Western Blot (1:250-1:500) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Western Blot

Western blot analysis of Lane 1: RT-4, Lane 2: U-251 MG, Lane 3: Human Plasma, Lane 4: Liver, Lane 5: Tonsil with LCA5 polyclonal antibody (Cat # PAB22243).

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human pancreas with LCA5 polyclonal antibody (Cat # PAB22243) shows moderate cytoplasmic positivity in exocrine glandular cells.

Gene Info — LCA5

Entrez GeneID	167691
Protein Accession#	Q86VQ0
Gene Name	LCA5
Gene Alias	C6orf152
Gene Description	Leber congenital amaurosis 5
Omim ID	604537 611408
Gene Ontology	Hyperlink
Gene Summary	This gene encodes a protein that is thought to be involved in centrosomal or ciliary functions. Mutations in this gene cause Leber congenital amaurosis type V. Alternative splicing results in two transcript variants. [provided by RefSeq]

Other Designations

OTTHUMP00000016774|OTTHUMP00000165989|lebercilin

Disease

- [Blindness](#)
- [Retinal Degeneration](#)