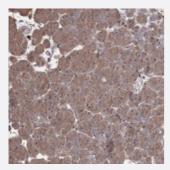


OPTC polyclonal antibody

Catalog # PAB22838 Size 100 uL

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Immunohistochemical staining of human pancreas with OPTC polyclonal antibody (Cat # PAB22838) shows moderate cytoplasmic positivity in exocrine pancreas at 1:20-1:50 dilution.

Specification	
Product Description	Rabbit polyclonal antibody raised against recombinant OPTC.
Immunogen	Recombinant protein corresponding to amino acids of human OPTC.
Sequence	RIDLSNNLISSIDNDAFRLLHALQDLILPENQLEALPVLPSGIEFLDVRLNRLQSSGIQPAAFRAMEK LQFLYLSDNLLDSIP
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Antigen affinity purification
Isotype	lgG
Recommend Usage	Immunohistochemistry (1:20-1:50) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide)



Product Information

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 shoul d be handled by trained staff only.

Applications

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Gene Info — OPTC	
Entrez GeneID	<u>26254</u>
Protein Accession#	Q9UBM4
Gene Name	OPTC
Gene Alias	OPT
Gene Description	opticin
Omim ID	605127
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Opticin belongs to class III of the small leucine-rich repeat protein (SLRP) family. Members of this family are typically associated with the extracellular matrix. Opticin is present in significant quantiti es in the vitreous of the eye and also localizes to the cornea, iris, ciliary body, optic nerve, choroid , retina, and fetal liver. Opticin may noncovalently bind collagen fibrils and regulate fibril morpholog y, spacing, and organization. The opticin gene is mapped to a region of chromosome 1 that is as sociated with the inherited eye diseases age-related macular degeneration (AMD) and posterior column ataxia with retinosa pigmentosa (AXPC1). [provided by RefSeq
Other Designations	OTTHUMP00000034332 oculoglycan

Disease

- Genetic Predisposition to Disease
- Glaucoma



- <u>Hypertension</u>
- Myopia