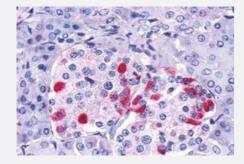


GIPR polyclonal antibody

Catalog # PAB26179 Size 50 ug

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) of human pancreas, islet of Langerhans with GIPR polyclonal antibody (Cat # PAB26179). Immunohistochemistry of formalin-fixed, paraffin-embedded tissue after heat-induced antigen retrieval.

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of GIPR.
Immunogen	A synthetic peptide corresponding to 18 amino acid at cytoplasmic domain of human GIPR.
Host	Rabbit
Reactivity	Human, Monkey
Specificity	BLAST analysis of the peptide immunogen showed no homology with other human proteins, except GHRHR (100%), GCGR (61%), GLP2R (61%).
Form	Liquid
Purification	Immunoaffinity chromatography
Recommend Usage	Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (32 ug/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.09% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -80°C. Aliquot to avoid repeated freezing and thawing.



Product Information

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

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

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Gene Info — GIPR	
Entrez GenelD	2696
Protein Accession#	P48546
Gene Name	GIPR
Gene Alias	MGC126722
Gene Description	gastric inhibitory polypeptide receptor
Omim ID	<u>137241</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Gastric inhibitory polypeptide (GIP; MIM 137240), also called glucose-dependent insulinotropic p olypeptide, is a 42-amino acid polypeptide synthesized by K cells of the duodenum and small inte stine. It was originally identified as an activity in gut extracts that inhibited gastric acid secretion a nd gastrin release, but subsequently was demonstrated to stimulate insulin release potently in the presence of elevated glucose. The insulinotropic effect on pancreatic islet beta-cells was then rec ognized to be the principal physiologic action of GIP. Together with glucagon-like peptide-1, GIP is largely responsible for the secretion of insulin after eating. It is involved in several other facets of the anabolic response.[supplied by OMIM
Other Designations	-

Pathway

Neuroactive ligand-receptor interaction



Disease

- Cardiovascular Diseases
- Diabetes Mellitus
- Genetic Predisposition to Disease
- Glucose Intolerance
- Hyperparathyroidism
- Insulin Resistance
- Metabolic Syndrome X
- Multiple Sclerosis
- Obesity