SANTA CRUZ BIOTECHNOLOGY, INC.

GIPR (H-70): sc-98795



BACKGROUND

GIPR (gastric inhibitory polypeptide receptor) is a 466 amino acid protein belonging to the G protein-coupled receptor 2 family. The activity of GIPR is mediated by G proteins, which activate adenylyl cyclase. Expressed as two isoforms produced by alternative splicing, GIPR is a multi-pass cell membrane protein that acts as a receptor for the glucose-dependent Insulinotropic polypeptide (GIP). GIP is a major physiologic factor in the augmentation of the Insulin response to oral glucose. GIP is a peptide hormone that is released postprandially from the small intestine and acts in concert with glucagon-like peptide GLP1 to potentiate glucose-induced Insulin secretion from the pancreatic β cell. GIP has been shown to increase adenylyl cyclase activity, elevate intracellular calcium levels, and stimulate a mitogen-activated protein kinase pathway in the pancreatic β cell. GIP release is demonstrated predominantly after ingestion of carbohydrate and fat and the effects of acid on GIP are consistent with a role for GIP as an enterogastrone.

REFERENCES

- 1. Yamada, Y. and Seino, Y. 2004. Physiology of GIP—a lesson from GIP receptor knockout mice. Horm. Metab. Res. 36: 771-774.
- Boylan, M.O., et al. 2006. Sp1/Sp3 binding is associated with cell-specific expression of the glucose-dependent Insulinotropic polypeptide receptor gene. Am. J. Physiol. Endocrinol. Metab. 290: E1287-E1295.
- Marenah, L., et al. 2006. A stable analogue of glucose-dependent insulinotropic polypeptide, GIP(LysPAL16), enhances functional differentiation of mouse embryonic stem cells into cells expressing islet-specific genes and hormones. Biol. Chem. 387: 941-947.
- Lampron, A., et al. 2006. Whole genome expression profiling of glucosedependent Insulinotropic peptide (GIP)- and adrenocorticotropin-dependent adrenal hyperplasias reveals novel targets for the study of GIP-dependent Cushing's syndrome. J. Clin. Endocrinol. Metab. 91: 3611-3618.
- Tsukiyama, K., et al. 2006. Gastric inhibitory polypeptide as an endogenous factor promoting new bone formation after food ingestion. Mol. Endocrinol. 20: 1644-1651.
- Irwin, N., et al. 2006. Biological activity and antidiabetic potential of synthetic fragment peptides of glucose-dependent Insulinotropic polypeptide, GIP(1-16) and (Pro3)GIP(1-16). Regul. Pept. 135: 45-53.

CHROMOSOMAL LOCATION

Genetic locus: GIPR (human) mapping to 19q13.32; Gipr (mouse) mapping to 7 A3.

SOURCE

GIPR (H-70) is a rabbit polyclonal antibody raised against amino acids 268-337 mapping within an internal region of GIPR of human origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

GIPR (H-70) is recommended for detection of GIPR of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

GIPR (H-70) is also recommended for detection of GIPR in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for GIPR siRNA (h): sc-75134, GIPR siRNA (m): sc-75135, GIPR shRNA Plasmid (h): sc-75134-SH, GIPR shRNA Plasmid (m): sc-75135-SH, GIPR shRNA (h) Lentiviral Particles: sc-75134-V and GIPR shRNA (m) Lentiviral Particles: sc-75135-V.

Molecular Weight of GIPR: 53 kDa.

Positive Controls: rat pituitary gland extract: sc-364807.

DATA

132 K – 90 K –			
55 K – 43 K –	-	< mm GIPR	
34 K –			

GIPR (H-70): sc-98795. Western blot analysis of GIPR expression in rat pituitary tissue extract.

SELECT PRODUCT CITATIONS

- 1. Puddu, A., et al. 2015. Effects of high glucose levels and glycated serum on GIP responsiveness in the pancreatic β cell line HIT-T15. J. Diabetes Res. 2015: 326359.
- Berglund, L.M., et al. 2015. Glucose-dependent insulinotropic polypeptide (GIP) stimulates osteopontin expression in the vasculature via endothelin-1 and CREB. Diabetes. E-published.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.