

## **Product datasheet for TA354264**

## OriGene Technologies, Inc.

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## **Amylin (IAPP) Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type:** Primary Antibodies

Applications: IHC

Recommended Dilution: WB 0.1-1 μg/ml ELISA 0.01-0.1 μg/ml IP 2-5 μg/ml IHC 2-10 μg/ml FC 5-10 μg/ml

Reactivity: Human
Host: Rabbit
Isotype: IgG

Clonality: Polyclonal

**Immunogen:** A synthetic peptide derived from N-term of human Amylin peptide.

**Formulation:** This affinity purified antibody is supplied in sterile Phosphate buffered saline (pH7.2)

containing antibody stabilizer.

**Purification:** The Rabbit IgG is purified by Epitope Affinity Purification

**Conjugation:** Unconjugated

Storage: Store at -20°C as received.

**Stability:** Stable for 12 months from date of receipt.

**Predicted Protein Size:** 10 kDa

**Gene Name:** islet amyloid polypeptide

Database Link: NP 000406

Entrez Gene 3375 Human

P10997

**Background:** Amylin is a 37 amino acid peptide co-secreted with insulin from pancreatic islet beta-cells.

Amylin shares structural similarities with calcitonin gene-related peptide (CGRP) and calcitonin (CT), including an N-terminal ring structure linked by a disulfide bridge, and an amidated C-terminus. The function of Amylin is to maintain the gluocose homoeostasis, include lowered insulin secretion, raised hepatic glucose output, stimulation of glycogenolysis and inhibition of glycogen synthesis in liver cells and in skeletal muscle. Amylin peptide is

predominantly found in the beta cells of the pancreas and to a lesser extent in the

gastrointestinal tract and nervous system. This antibody can be used for staining of amylin in

normal and diabetic pancreas and pancreatic carcinoma

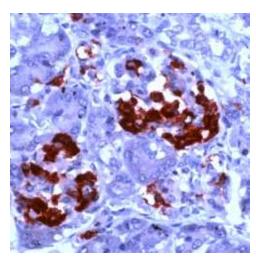




Synonyms: DAP; IAP

Protein Families: Druggable Genome, Secreted Protein
Protein Pathways: Maturity onset diabetes of the young

## **Product images:**



IHC: Human pancreatic tissue stained with Anti-Amylin antibody, at 1:100 for 10 min at RT. Staining of formalinfixed tissue requires boiling tissue sections in 10 mM Citrate Buffer, pH 6.0 for 10 min followed by cooling at RT for 20 min.