

PDX1 (PT1159R) PT™ Rabbit mAb

CatalogNo: YM9008 Recombinant R

Key Features

Host Species

Rabbit

MW
• 31kD (Calculated)
42kD (Observed)

Reactivity

Human, Mouse, Rat

Isotype

• IgG,Kappa

Applications

• WB,IHC,IF,IP,ELISA

Recommended Dilution Ratios

IHC 1:100-1:500 WB 1:500-1:20000 IF 1:200-1:1000

ELISA 1:5000-1:20000

IP 1:50-1:200

Storage

Storage* -15°C to -25°C/1 year(Do not lower than -25°C)

Formulation PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA

Basic Information

Clonality Monoclonal

Clone Number PT1159R

Immunogen Information

Specificity Endogenous

Target Information

Gene name

PDX1

Protein Name

Pancreas/duodenum homeobox protein 1

Organism	Gene ID	UniProt ID
Human	<u>3651</u> ;	<u>P52945;</u>
Mouse	<u>18609;</u>	<u>P52946;</u>
Rat	<u>29535;</u>	<u>P52947;</u>

Cellular Localization

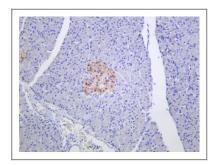
Nucleus. Cytoplasm, cytosol.

Tissue specificity Duodenum and pancreas (Langerhans islet beta cells and small subsets of endocrine nonbeta-cells, at low levels in acinar cells).

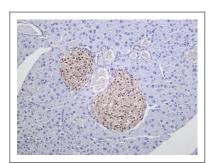
Function

Disease: Defects in PDX1 are a cause of pancreatic agenesis [MIM:260370]. This autosomal recessive disorder is characterized by absence or hypoplasia of pancreas, leading to earlyonset insulin-dependent diabetes mellitus. This was found in a frameshift mutation that produces a truncated protein and results in a second initiation that produces a second protein that act as a dominant negative mutant., Disease: Defects in PDX1 are the cause of maturity onset diabetes noninsulin-dependent diabetes mellitus (NIDDM) [MIM:125853]: also known as diabetes mellitus type II., Disease: Defects in PDX1 are the cause of maturity onset diabetes of the young type 4 (MODY4) [MIM:606392]; also symbolized MODY-4. MODY [MIM:606391] is a form of diabetes mellitus characterized by an autosomal dominant mode of inheritance, age of onset of 25 years or younger and a primary defect in insulin secretion., Domain: The Antp-type hexapeptide mediates heterodimerization with PBX on a regulatory element of the somatostatin promoter., Domain: The homeodomain, which contains the nuclear localization signal, not only mediates DNA-binding, but also acts as a protein-protein interaction domain for TCF3(E47), NEUROD1 and HMG-I(Y)...Function: Activates insulin, somatostatin, glucokinase, islet amyloid polypeptide and glucose transporter type 2 gene transcription. Particularly involved in glucose-dependent regulation of insulin gene transcription. Binds preferentially the DNA motif 5'-[CT]TAAT[TG]-3'. During development, specifies the early pancreatic epithelium, permitting its proliferation, branching and subsequent differentiation. At adult stage, required for maintaining the hormone-producing phenotype of the beta-cell., miscellaneous: According to PubMed:16141209, it may be methylated by SETD7 in vitro. However, the relevance of methylation is unsure in vivo., PTM: Phosphorylated by the SAPK2 pathway at high intracellular glucose concentration., similarity: Belongs to the Antp homeobox family. IPF1/XIHbox-8 subfamily., similarity: Contains 1 homeobox DNA-binding domain., subunit: Interacts with the basic helix-loop-helix domains of TCF3(E47) and NEUROD1 and with HMG-I(Y). Interacts with SPOP (By similarity). Interacts with the methyltransferase SETD7.,tissue specificity:Duodenum and pancreas (Langerhans islet beta cells and small subsets of endocrine non-beta-cells, at low levels in acinar cells).,

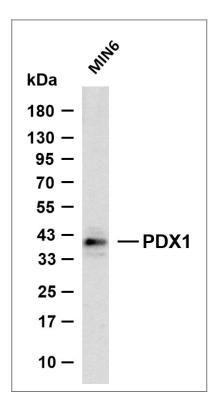
Validation Data



Rat pancreas was stained with anti-PDX1 (PT1159R) Rabbit antibody



Mouse pancreas was stained with anti-PDX1 (PT1159R) Rabbit antibody



Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-PDX1 (PT1159R) antibody. The HRP-conjugated Goat anti-Rabbit IgG (H + L) antibody was used to detect the antibody. Lane 1: MIN6 Predicted band size: 31kDa Observed band size: 42kDa

| Contact information

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Please scan the QR code to access additional product information:
PDX1 (PT1159R)
PT™ Rabbit mAb

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