

# PARD6A antibody, C-term

## Cat. No. GTX26022

Host	Goat
Clonality	Polyclonal
Isotype	IgG
Applications	IHC-P
Reactivity	Human

Package  $100 \, \mu g$ 

## **Applications**

## **Application Note**

\*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution	
IHC-P	5-10μg/ml	
Note: Human Pancreas shows strong cytoplasmic staining in rare cells of intralobular ducts.		

Not tested in other applications.

This antibody is expected to recognise both reported isoforms (NP\_058644.1; NP\_001032358.1). **Product Note** 

Properties	
Form	Liquid
Buffer	TBS, 0.5% BSA
Preservative	0.02% Sodium azide
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	0.50 mg/ml (Please refer to the vial label for the specific concentration.)
Immunogen	Peptide with sequence C-GSRIRGDGSGFSL, from the C Terminus of the protein sequence according to NP_058644.1; NP_001032358.1.
Purification	Purified by ammonium sulphate precipitation followed by antigen affinity chromatography
Conjugation	Unconjugated
Note	For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.
	Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.



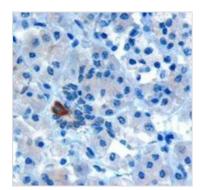
For full product information, images and publications, please visit our website.

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## DATA IMAGES

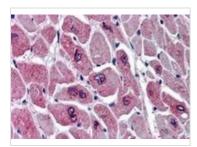


#### GTX26022 IHC-P Image

IHC-P analysis of human pancreas using GTX26022 PARD6A antibody, C-term.

Antigen retrieval: Tris/EDTA buffer pH 9

Dilution: 10µg/ml



#### GTX26022 IHC-P Image

IHC-P analysis of human heart using GTX26022 PARD6A antibody, C-term.

Antigen retrieval: citrate buffer pH 6

Dilution: 5µg/ml



For full product information, images and publications, please visit our <u>website</u>.

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