

DNAxPAb



## PEX5 DNAxPab

Catalog # H00005830-W01P Size 200 ug

Specification	
Product Description	Rabbit polyclonal antibody raised against a partial-length human PEX5 DNA using DNAx™ Immune t echnology.
Technology	DNAx <sup>™</sup> Immune
Immunogen	Extracellular membrane domain (ECD) human DNA
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## Applications

- Western Blot (Transfected lysate)
  <u>Protocol Download</u>
- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

## Gene Info — PEX5

😵 Abnova	Product Information
Entrez GenelD	5830
GeneBank Accession#	<u>NM_000319.3</u>
Protein Accession#	<u>NP_000310.2</u>
Gene Name	PEX5
Gene Alias	FLJ50634, FLJ50721, FLJ51948, PTS1-BP, PTS1R, PXR1
Gene Description	peroxisomal biogenesis factor 5
Omim ID	<u>202370 214100 600414</u>
Gene Ontology	Hyperlink
Gene Summary	The product of this gene binds to the C-terminal PTS1-type tripeptide peroxisomal targeting signa I (SKL-type) and plays an essential role in peroxisomal protein import. Peroxins (PEXs) are protein ns that are essential for the assembly of functional peroxisomes. The peroxisome biogenesis disor rders (PBDs) are a group of genetically heterogeneous autosomal recessive, lethal diseases cha racterized by multiple defects in peroxisome function. The peroxisomal biogenesis disorders are a heterogeneous group with at least 14 complementation groups and with more than 1 phenotype being observed in cases falling into particular complementation groups. Although the clinical featu res of PBD patients vary, cells from all PBD patients exhibit a defect in the import of one or more classes of peroxisomal matrix proteins into the organelle. Defects in this gene are a cause of neo natal adrenoleukodystrophy (NALD), a cause of Zellweger syndrome (ZWS) as well as may be a c ause of infantile Refsum disease (IRD). Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq
Other Designations	peroxin-5 peroxisomal C-terminal targeting signal import receptor peroxisomal targeting signal 1 ( SKL type) receptor peroxisomal targeting signal import receptor peroxisomal targeting signal rec eptor 1 peroxisome receptor 1