

Rabbit PEX3 Polyclonal Antibody | anti-PEX3 antibody

PEX3 Polyclonal Antibody

CATALOG # AAA311511	HOST Rabbit	REACTIVITY Human, Mouse, Rat	APPLICATIONS Immunohistochemistry, Western Blot
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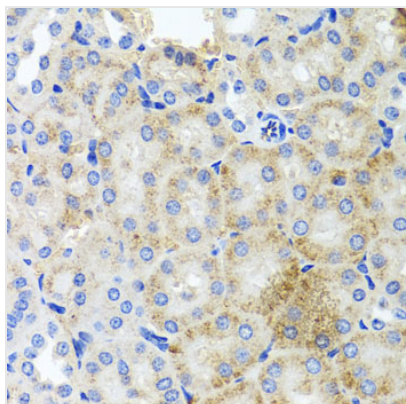
PRODUCT SYNONYMS

PEX3; Polyclonal Antibody; PEX3 Polyclonal Antibody; PBD10A; PBD10B; TRG18; peroxisomal biogenesis factor 3; anti-PEX3 antibody

PRODUCT SPECIFICATIONS

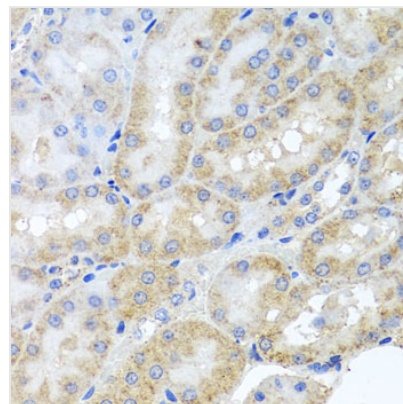
Host	Rabbit
Reactivity	Human, Mouse, Rat
Clonality	Polyclonal
Isotype	IgG
Purity/Purification	Affinity purification
Form/Format	PBS with 50% glycerol, pH7.4
Applicable Applications for anti-PEX3 antibody	IHC (Immunohistochemistry), WB (Western Blot)
Immunogen	Recombinant fusion protein of human PEX3 (NP_003621.1).
Conjugation	Unconjugated
Preparation and Storage	Store at -20 degree C. Avoid freeze/thaw cycles.

IMAGES



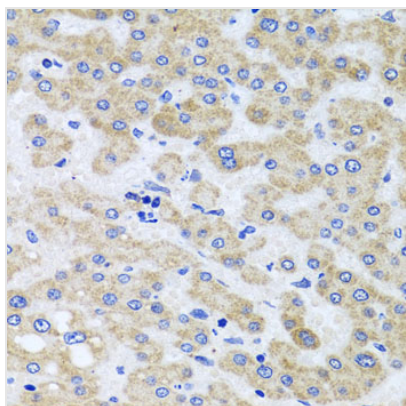
IHC (Immunohistochemistry)

Immunohistochemistry of paraffin-embedded mouse kidney using PEX3 antibody.

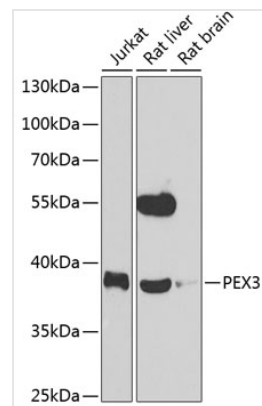


IHC (Immunohistochemistry)

Immunohistochemistry of paraffin-embedded rat kidney using PEX3 antibody.



IHC (Immunohistochemistry)
Immunohistochemistry of paraffin-embedded human liver damage using PEX3 antibody.



WB (Western Blot)
Western blot analysis of extracts of various cell lines, using PEX3 antibody.

ADDITIONAL INFORMATION

Related Product Information for anti-PEX3 antibody	The product of this gene is involved in peroxisome biosynthesis and integrity. It assembles membrane vesicles before the matrix proteins are translocated. Peroxins (PEXs) are proteins that are essential for the assembly of functional peroxisomes. The peroxisome biogenesis disorders (PBDs) are a group of genetically heterogeneous autosomal recessive, lethal diseases characterized 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 features 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 Zellweger syndrome (ZWS).
Product Categories/Family for anti-PEX3 antibody	Total protein Ab

NCBI AND UNIPROT INFORMATION

NCBI GI #	4505727
NCBI GeneID	8504
NCBI Accession #	NP_003621.1
NCBI GenBank Nucleotide #	NM_003630.2
UniProt Accession #	P56589; Q6FGP5
Molecular Weight	42,140 Da
NCBI Official Full Name	peroxisomal biogenesis factor 3
NCBI Official Synonym Full Names	peroxisomal biogenesis factor 3
NCBI Official Symbol	PEX3
NCBI Official Synonym Symbols	TRG18; PBD10A
NCBI Protein Information	peroxisomal biogenesis factor 3; peroxin-3; peroxisomal assembly protein PEX3; transformation-related protein 18
UniProt Protein Name	Peroxisomal biogenesis factor 3
UniProt Gene Name	PEX3
UniProt Entry Name	PEX3_HUMAN

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Small volumes of product may become entrapped in the seal of the vial during shipment and storage; if necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the cap.