

#### OriGene Technologies, Inc.

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# Product datasheet for TA501390S

## PEX5 Mouse Monoclonal Antibody [Clone ID: OTI6C4]

## **Product data:**

Product Type:	Primary Antibodies
Clone Name:	OTI6C4
Applications:	IF, WB
Recommended Dilution:	WB 1:2000, IF 1:100
Reactivity:	Human, Mouse, Rat
Host:	Mouse
lsotype:	lgG2b
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human PEX5 (NP_000310) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.91 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	69.7 kDa
Gene Name:	peroxisomal biogenesis factor 5
Database Link:	<u>NP 000310</u>
	<u>Entrez Gene 19305 MouseEntrez Gene 312703 RatEntrez Gene 5830 Human</u> <u>P50542</u>



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#### Serigene PEX5 Mouse Monoclonal Antibody [Clone ID: OTI6C4] – TA501390S

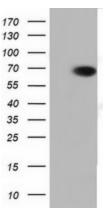
Background: The product of this gene binds to the C-terminal PTS1-type tripeptide peroxisomal targeting signal (SKL-type) and plays an essential role in peroxisomal protein import. Peroxins (PEXs) are proteins that are essential for the assembly of functional peroxisomes. The peroxisome biogenesis disorders (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 of neonatal adrenoleukodystrophy (NALD), a cause of Zellweger syndrome (ZWS) as well as may be a cause of infantile Refsum disease (IRD). Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq]

Synonyms: PBD2A; PBD2B; PTS1-BP; PTS1R; PXR1; RCDP5

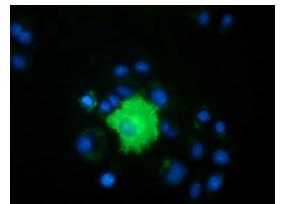
**Protein Families:** 

Druggable Genome

### **Product images:**



HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY PEX5 ([RC202062], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-PEX5. Positive lysates [LY424800] (100ug) and [LC424800] (20ug) can be purchased separately from OriGene.



Anti-PEX5 mouse monoclonal antibody ([TA501390]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY PEX5 ([RC202062]).

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