

## Product datasheet for **RG226067**

### PEX5 (NM\_001131023) Human Tagged ORF Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	PEX5 (NM_001131023) Human Tagged ORF Clone
Tag:	TurboGFP
Symbol:	PEX5
Synonyms:	PBD2A; PBD2B; PTS1-BP; PTS1R; PXR1; RCDP5
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-AC-GFP (PS100010)
E. coli Selection:	Ampicillin (100 ug/mL)



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ORF Nucleotide  
Sequence:

>RG226067 representing NM\_001131023  
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
 GCC**CGGATCGCC**

ATGGCAATGCGGGAGCTGGTGGAGGCCGAATGCGGGGTGCCAACCCGCTCATGAAGCTCGCCGGGCACT  
 TCACCCAGGACAAGGCCCTTCGGCAGGAGGGATTGAGGCCTGGCCCTGGCCCCGGAGCCCCGGCCTC  
 TGAGGCAGTGAGTGTTCTTGAGGTGAAAAGCCAGGTGCAGCCTCTGAGGCAGCCTCCAAGCCTTTGGGA  
 GTAGTCTCTGAAGATGAGTTGGTGGCTGAATTCCTGCAGGACCAGAATGCACCCCTTGTGTCCCCTGCC  
 CTCAGACCTTCAAGATGGATGACCTCCTGGCTGAGATGCAGCAGATTGAGCAGTCAAACCTCCGCCAGGC  
 TCCCCAGAGAGCCCTGGTGTGGCAGACTTGGCCTTGTCTGAGAAGTGGGCCAGGAGTTTCTTGCAGCT  
 GGAGATGCTGTGGATGTAACCTCAGGATTATAATGAGACTGACTGGTCCCAAGAATTCATCTCTGAAGTTA  
 CAGACCCCTTGTCTGTGTCCCCTGCCCGTGGGCTGAGGAATATTTGGAGCAATCAGAGGAGAAGCTGTG  
 GCTGGGAGAACCAGGGAAACAGCCACCGATCGCTGGTATGATGAATATCATCTGAGGAGGATCTGCAG  
 CACACGGCCAGTGACTTTGTGGCCAAAGTGGATGACCCCAAATGGCTAATTCAGTTCCTGAAATTCG  
 TGGCGCAGATTGGCGAAGGGCAGGTGTCCCTGGAGTCCGGTGCAGGGTCCGGCCGAGCTCAGGCAGAACA  
 GTGGGCAGCAGAGTTTATACAGCAGCAGGGTACATCAGATGCCTGGGTTGACCAGTTCACAAGACCAGTA  
 AACACATCTGCCCTTGATATGGAGTTTGAACGAGCCAAGTCAGCTATAGAGTCTGATGTCGATTTCTGGG  
 ACAAGTTGCAGGCAGAGTTGGAGGAGATGGCAAAACGGGATGCTGAGGCCACCCCTGGCTTTCTGACTA  
 TGATGACCTTACGTCAGTACCTATGATAAGGGGTACCAGTTTGGAGGAGAGAACCCTTGGCGTATCAC  
 CCTCAGCCTTTTGAAGAAGGGCTGCGGCCCTTCAGGAGGGGGACCTGCCAAATGCTGTGCTGTTTTTG  
 AGGCAGCTGTGCAGCAGGATCCTAAGCACATGGAAGCTTGGCAGTATCTGGGTACCACCCAGGCAGAGAA  
 TGAACAAGAACTATTAGCCATCAGTGCATTGCGGAGGTGTCTGGAGCTAAAGCCAGATAACCAGACAGCA  
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 GGCTGCGGTACACACCAGCCTATGCCCATCTGGTACACCTGCTGAAGAAGGGGCTGGTGGGGCAGGACT  
 GGGCCCCAGCAAGCGTATCCTGGGATCTCTTGTCTGACTCCCTGTTTCTTGAAGTGAAGAGCTCTTC  
 CTGGCAGCTGTGCGGCTGGACCCTACCTCCATTGACCCTGATGTGCAGTGTGGCTTGGGAGTCTTTTCA  
 ACCTGAGTGGGGAGTATGACAAGGCCGTGGACTGCTTACAGCTGCCCTCAGCGTTCGTCCCAATGACTA  
 TTTGCTGTGGAATAAGCTAGGCGCCACCCTGGCCAATGGAACCAGAGTGAAGAAGCAGTAGCTGCGTAC  
 CGCCGGGCCCTCGAGCTCCAGCCTGGCTATATCCGGTCCCCTATAACCTGGGCATCAGCTGCATCAACC  
 TCGGGGCTCACCGGAGGCTGTGGAGCACTTTCTGGAGGCCCTGAACATGCAGAGGAAAAGCCGGGGCCC  
 CCGGGGTGAAGGAGGTGCCATGTGCGGAGAACATCTGGAGCACCCCTGCGTTTGGCATTGTCTATGTTAGGC  
 CAGAGCGATGCCTATGGGGCAGCCGACGCGGGGATCTGTCCACCCTCTAACTATGTTTGGCCTGCCCC  
 AG

**ACGCGT**ACGCGGGCCGCTCGAG – GFP Tag – GTTTAA

Protein Sequence: >RG226067 representing NM\_001131023  
 Red=Cloning site Green=Tags(s)

MAMRELVEAECGGANPLMKLAGHFTQDKALRQEGLRPGWPPGAPASEAVSVLEVESPGAASEAASKPLG  
 VASEDELVAEFLQDQNAFLVSRAPQTFKMDDLAEQQIEQSNFRQAPQAPGVADLALSENWAQEFLLAA  
 GDAVDVTQDYNETDWSQEFISEVTDPLSVSPARWAEYLEQSEEKWLGEPEGTADRWDYDEYHPEEDLQ  
 HTASDFVAKVDDPKLANSEFLKFVRQIGEGQVLSLESGAGSGRAQAEQWAAEFIQQQGTSDAWVDQFTRPV  
 NTSALDMEFERAKSAIESDVDFWDLQAELEEMAKRDAEAHPWLSDYDDLTSATYDKGYQFEEENPLRDH  
 PQPFEEGLRRLQEGDLPNAVLLFEAAVQQDPKHMEAWQYLGTQAENEQELLAISALRRCLELKPNDQTA  
 LMAVLSFTNESLQRQACETLRDRLRYTPAYAHLVTPAEEGAGGAGLGPSKRILGSLLSDSLFLVKELF  
 LAAVRLDPTSIDPDVQCGLGVLFNLSGEYDKAVDCFTAALSVRPNDYLLWNKLGATLANGQSEEVAAY  
 RRALQLPGYIRSRYNLGISCNLGAHREAVEHFLEALNMQRKSRGPRGEGGAMSENIWSTLRLALSMLG  
 QSDAYGAADARDLSTLLTMFGLPQ

TRTRPLE - GFP Tag - V

Restriction Sites: SgfI-MluI

Cloning Scheme:



ACCN: NM\_001131023

ORF Size: 1962 bp

**OTI Disclaimer:** Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at [custsupport@origene.com](mailto:custsupport@origene.com) or by calling 301.340.3188 option 3 for pricing and delivery.

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

**Components:** The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

**Reconstitution Method:**

1. Centrifuge at 5,000xg for 5min.
2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

**RefSeq:** [NM\\_001131023.2](#)

**RefSeq Size:** 3324 bp

**RefSeq ORF:** 1965 bp

**Locus ID:** 5830

**UniProt ID:** [P50542](#)

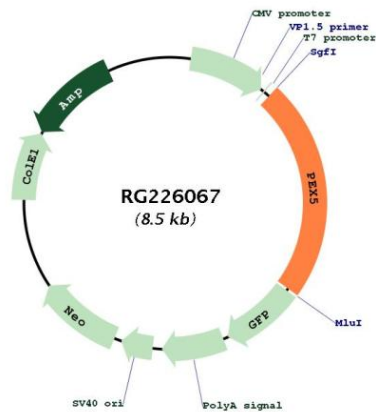
**Cytogenetics:** 12p13.31

**Protein Families:** Druggable Genome

**Gene Summary:**

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, Oct 2008]

**Product images:**



Circular map for RG226067