

## Product datasheet for **RC239243**

### PEX5 (NM\_001300789) Human Tagged ORF Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	PEX5 (NM_001300789) Human Tagged ORF Clone
Tag:	Myc-DDK
Symbol:	PEX5
Synonyms:	PBD2A; PBD2B; PTS1-BP; PTS1R; PXR1; RCDP5
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)



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ORF Nucleotide  
Sequence:

>RC239243 representing NM\_001300789  
Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
GCC**CGGATCGCC**

ATGCTCCTCTGCCGTGCTCACCGCTGCTGGGGCTGCGCGGGCTAGAGAGCTGGCGGTACCATGGCAA  
TGCGGGAGCTGGTGGAGGCCAATGCGGGGTGCCAACCCGCTCATGAAGCTCGCCGGGCACTTCACCCA  
GGACAAGGCCCTTCGGCAGGAGGATTGAGGCCTGGCCCTGGCCCCGGAGCCCCGGCCTCTGAGGCA  
GCCTCAAAGCCTTTGGGAGTAGCTTCTGAAGATGAGTTGGTGGCTGAATTCCTGCAGGACCAGAATGCAC  
CCCTTGTGTCCCCTGCCCTCAGACCTTCAAGATGGATGACCTCCTGGCTGAGATGCAGCAGATTGAGCA  
GTCAAACCTCCGCCAGGCTCCCCAGAGAGCCCCTGGTGTGGCAGACTTGGCCTTGTCTGAGAACTGGGCC  
CAGGAGTTTCTGCAGCTGGAGATGCTGTGGATGTAACCTAGGATTATAATGAGACTGACTGGTCCCAAG  
AATTCATCTCTGAAGTTACAGACCCCTTGTCTGTGTCCCCTGCCGCTGGGCTGAGGAATATTTGGAGCA  
ATCAGAGGAGAAGCTGTGGCTGGGAGAACCTGAGGGAACAGCCACCGATCGCTGGTATGATGAATATCAT  
CCTGAGGAGGATCTGCAGCACACGGCCAGTGACTTTGTGGCCAAAGTGGATGACCCCAAATTTGGCTAATT  
CTGAGTTCCTGAAATTCGTGCGGCAGATTGGCGAAGGGCAGGTGTCCCTGGAGTCCGGTGCAGGGTCCGG  
CCGAGCTCAGGCAGAACAGTGGGCAGCAGAGTTTATACAGCAGCAGGGTACATCAGATGCCTGGGTTGAC  
CAGTTCACAAGACCAGTAAACACATCTGCCCTTGATATGGAGTTTGAACGAGCCAAGTCAGCTATAGAGT  
CTGATGTCGATTTCTGGGACAAGTTGCAGGCAGAGTTGGAGGAGATGGCAAAACGGGATGCTGAGGCCCA  
CCCCTGGCTTTCTGACTATGATGACCTTACGTCAGCTACCTATGATAAGGGGTACCAGTTGAGGAGGAG  
AACCCCTTGCCTGATCACCTCAGCCTTTTGAAGAAGGGCTGCGGCGCCTTCAGGAGGGGACCTGCCAA  
ATGCTGTGCTGCTTTTTGAGGCAGCTGTGCAGCAGGATCCTAAGCACATGGAAGCTTGGCAGTATCTGGG  
TACCACCCAGGCAGAGAATGAACAAGAACTATTAGCCATCAGTGCATTGCGGAGGTGTCTGGAGCTAAAG  
CCAGATAACCAGACAGCACTGATGGCGCTGGCTGTGAGCTTACCAACGAGTCCCTGCAGCGACAGGCCT  
GTGAAACCTACGAGACTGGCTGCGGTACACACCAGCCTATGCCATCTGGTGCACCTGCTGAAGAAGG  
GGCTGGTGGGCAGGACTGGGCCCAGCAAGCGTATCCTGGGATCTCTTGTCTGACTCCCTGTTTCTT  
GAAGTGAAGAGCTTCTCCTGGCAGCTGTGCGGCTGGACCCTACCTCCATTGACCCTGATGTGCAGTGTG  
GCTTGGGAGTCTTTTCAACCTGAGTGGGAGTATGACAAGGCCGTGGACTGCTTACAGCTGCCCTCAG  
CGTTCGTCCTAATGACTATTTGCTGTGAATAAGCTAGGCGCCACCTGGCCAATGGAACCCAGAGTGAA  
GAAGCAGTAGCTGCGTACCGCCGGGCCCTCGAGCTCCAGCCTGGCTATATCCGGTCCCGCTATAACCTGG  
GCATCAGCTGCATCAACCTCGGGGCTCACCGGGAGGCTGTGGAGCACTTCTGGAGGCCCTGAACATGCA  
GAGGAAAAGCCGGGGCCCCGGGGTGAAGGAGGTGCCATGTCCGGAACATCTGGAGCACCTGCCTTTG  
GCATTGTCTATGTTAGGCCAGAGCGATGCCTATGGGGCAGCCGACGCGGGGATCTGTCCACCCTCTAA  
CTATGTTTGGCCTGCCCCAG

**ACGCGT**ACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAATGATATCCTGGATT  
ACAAGGATGACGACGATAAGGTTTAA

**Protein Sequence:** >RC239243 representing NM\_001300789  
Red=Cloning site Green=Tags(s)

MLLCRAHRVGLRGARELAVTMAMRELVEAECGGANPLMKLAGHFTQDKALRQEGLRPGPWPAPASEA  
 ASKPLGVASEDELVAEFLQDQNAFLVSRAPQTFKMDLLAEMQQIEQSNFRQAPQRAPGVADLALSENWA  
 QEFLAAGDAVDVTQDYNEDWSQEFISEVTDPLSVSPARWAEYELEQSEEKWLWGEPEGTATDRWYDEYH  
 PEEDLQHTASDFVAKVDDPKLANSEFLKFRVQIGEGQVLSLESGAGSGRAQAEQWAAEFIQQGTSDAWVD  
 QFTRPVNTSALDMEFERAKSAIESDVDFWDLQAELEEMAKRDAEAHPWLSYDDLTSATYDKGYQFEEE  
 NPLRDHPQPFEEGLRRLQEGDL PNAVLLFEAAVQQDPKHMEAWQYL GTTQAEENEQELLAISALRRCLELK  
 PDNQ TALMALAVSFTNESLQRQACETLRDWLRYTPAYAHLVTPAEEGAGGAGLGPSKRILGSLSDSLFL  
 EVKELFLAAVRLDPTSIDPDVQCGLGVL FNL SGEYDKAVDCFTAALSVRPNDYLLWNKLGATLANGNQSE  
 EAVAAYRRALELQPGYIRSRYNLGISINLGAHREAVEHFLEALNMQRKSRGPRGEGGAMSENIWSTLRL  
 ALSMLGQSDAYGAADARDLSTLLTMFGLPQ

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

**Restriction Sites:**

Sgfl-MluI

**Cloning Scheme:**



**ACCN:** NM\_001300789

**ORF Size:** 1980 bp

**OTI Disclaimer:** 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\\_001300789.1](#), [NP\\_001287718.1](#)

**RefSeq Size:** 3350 bp

**RefSeq ORF:** 1920 bp

**Locus ID:** 5830

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

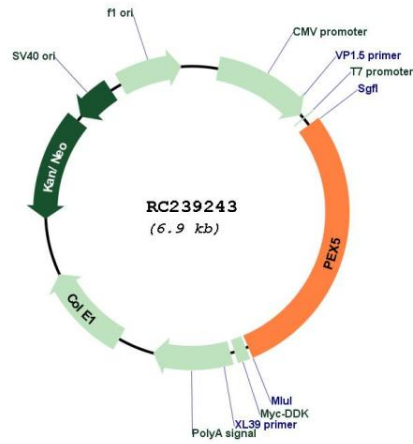
**Cytogenetics:** 12p13.31

**Protein Families:** Druggable Genome

**MW:** 73.6 kDa

**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 RC239243