

Product datasheet for SC208004

PEX16 (NM 004813) Human 3' UTR Clone

Product data:

Insert Size:

Product Type: 3' UTR Clones

Product Name: PEX16 (NM_004813) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

PEX16 Symbol:

Synonyms: PBD8A; PBD8B

ACCN: NM 004813

621 bp >SC208004 3'UTR clone of NM_004813 **Insert Sequence:**

The sequence shown below is from the reference sequence of NM_004813. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CAGAAAATCTACTTCTACAGTTGGGGCTGACAGACCTCCCGGAAGGAGGGTGTGGGGAGGGGTGGGGCA GGGAGCCCCTCTTCCCTAATAAAACTGACTCCGGCAGCGTCCCAGCGTGCGGCCTCTCCGTGCCTACCG GCCAGGCCCACACACACACCTGGTCGCCACCAGCGTTCCTCCCAGGACACCCTTGACTGCGCTCTCCT GTGACGATGCCACTGCAGCCCGCACCTTGTCACTGCTGGGCCAAGAAGCCTTCACTAGGAGTGGGATCC AGGCTCCTCTCCCACAGAAAGCGGTGACTTCACCTCATGGAGCCCGGGAAGCTGCTCGCCTCGGCAGCC ATAGGAGCGAACACTGCTGCTCTCTCGCTGGCCCCTGGTGAGGACAGGAAGCCTGAACCCGGGTGATGG CTGAACGCTGCCCAGCGTGTCTTCTGGCTGGGGCCCTCCGTCTGCCCCTTCTCCGCAGGGCCCTGTGGC TCTGGCAGCCCCAGGCCATGGCGTTGCCAGCCTCCCTGTGACAGAGCCTGGTGAACAGTGAGCCTGGCT CCCACGCAAGTGGCACTTTAAGCCCTGCATCCTCGGTTGAGAGTAAAAGGCTTTTCTCCCTTAGAAAAA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

OTI Disclaimer: Our molecular clone sequence data has been matched to the sequence identifier above as a

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The Components:

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.



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PEX16 (NM_004813) Human 3' UTR Clone - SC208004

RefSeq: <u>NM 004813.4</u>

Summary: The protein encoded by this gene is an integral peroxisomal membrane protein. An

inactivating nonsense mutation localized to this gene was observed in a patient with Zellweger syndrome of the complementation group CGD/CG9. Expression of this gene product morphologically and biochemically restores the formation of new peroxisomes, suggesting a role in peroxisome organization and biogenesis. Alternative splicing has been observed for this gene and two variants have been described. [provided by RefSeq, Jul 2008]

Locus ID: 9409 **MW:** 22.2