

Product datasheet for **SC206938**

PHYH (NM_001037537) Human 3' UTR Clone

Product data:

Product Type:	3' UTR Clones
Symbol:	PHYH
Synonyms:	LN1; LNAP1; PAHX; PHYH1; RD
Mammalian Cell	Neomycin
Selection:	
Vector:	pMirTarget (PSI00062)
ACCN:	NM_001037537
Insert Size:	532 bp

Insert Sequence: >SC206938 3'UTR clone of NM_001037537
The sequence shown below is from the reference sequence of NM_001037537. The complete sequence of this clone may contain minor differences, such as SNPs.
Blue=Stop Codon Red=Cloning site

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GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC
CTTGTGAAAGGAGAAAGAACCAATCTTGAATAGCCATCTGCTATAACTCTTCAACAGAAAACCAAA
ACCAACGAAATGTCTAAGGAAATGTTTTCTTAATGAGATGATGAACCTTTCTATCACTTGTTAAA
AGCAGAAAAACATGTATCAGGTACTTAATTGCATAGAGTTAGTTTTGCAGCACAATGGTGTGCTTTAAT
GGAAAAAAAACAGTAAAGTGAAATATTACTGTTTTAAGGAAAATAATTTAGGGTGCGCAGCCAATA
AAGGTGGTTGGTGTCTAATTTAAGTGTTAAATCAATTTCTTTCATTAGCTCTTTACCCAAGAAG
AAGTGAATGATTTGGAGCTTAGGGTATGTTTTGTATCCCTTTCTGATAAACCCATTCCCTACCAATTT
TATGTCATAAGAGATTTTTTCCCCAAATCTAGAACAATGTATAATACATTACATCTAGTCAAGGGC
ATAGGAACGGTGTATGGAGTCCAAATAAAGTGATATTCCTGCTCGGA
ACGCGTAAGCGGCCGCGGCATCTAGATTCAAGAAAAATGACCGACCAAGCGACGCCAACCTGCCATCA
CGAGATTTTCGATTCCACCGCCGCTTCTATGAAAGG
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Restriction Sites: SgfI-MluI

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).



Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.
Note:	Plasmids are not sterile. For experiments where strict sterility is required, filtration with 0.22um filter is required.
RefSeq:	<u>NM_001037537.2</u>
Summary:	This gene is a member of the PhyH family and encodes a peroxisomal protein that is involved in the alpha-oxidation of 3-methyl branched fatty acids. Specifically, this protein converts phytanoyl-CoA to 2-hydroxyphytanoyl-CoA. Mutations in this gene have been associated with Refsum disease (RD) and deficient protein activity has been associated with Zellweger syndrome and rhizomelic chondrodysplasia punctata. Alternate transcriptional splice variants, encoding different isoforms, have been characterized. [provided by RefSeq, Jul 2008]
Locus ID:	5264
MW:	20.5