

Product datasheet for SC334829

PCMT1 (NM_001252052) Human Untagged Clone

Product data:

OriGene Technologies, Inc.

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Product Type:	Expression Plasmids
Product Name:	PCMT1 (NM_001252052) Human Untagged Clone
Tag:	Tag Free
Symbol:	PCMT1
Synonyms:	PIMT
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_001252052, the custom clone sequence may differ by one or more nucleotides
	ATGCCGGGAGCGCGCAGTGGCGGCGGCGGCGACGGCAGTAACAGCGGCAGCTACAGCGGGGACGCGA GCGGGGCGGTGACGGTGTGGGAGGTGGTCTCACTCTTGGGAAAACTGCTGGGCACCGTCGTCGCGCGCG
Restriction Sites:	Sgfl-Mlul
ACCN:	NM_001252052
OTI Disclaimer:	Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).



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ORIGENE PCMT1 (NM_001252052) Human Untagged Clone – SC334829	
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:	 Centrifuge at 5,000xg for 5min. Carefully open the tube and add 100ul of sterile water to dissolve the DNA. Close the tube and incubate for 10 minutes at room temperature. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom. 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:	<u>NM 001252052.1, NP 001238981.1</u>
RefSeq Size:	1599 bp
RefSeq ORF:	756 bp
Locus ID:	5110
Cytogenetics:	6q25.1
Protein Families:	Druggable Genome
Gene Summary:	This gene encodes a member of the type II class of protein carboxyl methyltransferase enzymes. The encoded enzyme plays a role in protein repair by recognizing and converting D-aspartyl and L-isoaspartyl residues resulting from spontaneous deamidation back to the normal L-aspartyl form. The encoded protein may play a protective role in the pathogenesis of Alzheimer's disease, and single nucleotide polymorphisms in this gene have been associated with spina bifida and premature ovarian failure. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. [provided by RefSeq, Oct 2011] Transcript Variant: This variant (5) lacks an in-frame coding exon and uses an alternate splice

Transcript Variant: This variant (5) lacks an in-frame coding exon and uses an alternate splice site in the 3' coding region, which results in a frameshift, compared to variant 1. The encoded isoform (5) is shorter and has a distinct C-terminus, compared to isoform 1.

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