

OriGene Technologies, Inc.

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Product datasheet for RC220492L2V

PEAMT (PEMT) (NM_148172) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type:	Lentiviral Particles
Product Name:	PEAMT (PEMT) (NM_148172) Human Tagged ORF Clone Lentiviral Particle
Symbol:	PEAMT
Synonyms:	PEAMT; PEMPT; PEMT2; PLMT; PNMT
Mammalian Cell Selection:	None
Vector:	pLenti-C-mGFP (PS100071)
Tag:	mGFP
ACCN:	NM_148172
ORF Size:	708 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC220492).
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. <u>More info</u>
OTI Annotation:	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.
RefSeq:	<u>NM 148172.1</u>
RefSeq Size:	1026 bp
RefSeq ORF:	711 bp
Locus ID:	10400
UniProt ID:	<u>Q9UBM1</u>
Cytogenetics:	17p11.2
Domains:	PEMT
Protein Families:	Transmembrane



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GRIGENE PEAMT (PEMT) (NM_148172) Human Tagged ORF Clone Lentiviral Particle – RC220492L2V	
Protein Pathways:	Glycerophospholipid metabolism, Metabolic pathways
MW:	25.9 kDa
Gene Summary:	Phosphatidylcholine (PC) is the most abundant mammalian phospholipid. This gene encodes an enzyme which converts phosphatidylethanolamine to phosphatidylcholine by sequential methylation in the liver. Another distinct synthetic pathway in nucleated cells converts intracellular choline to phosphatidylcholine by a three-step process. The protein isoforms encoded by this gene localize to the endoplasmic reticulum and mitochondria-associated membranes. Alternate splicing of this gene results in multiple transcript variants encoding different isoforms. [provided by RefSeq, May 2012]

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