

Product datasheet for RC224518L1V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

PXDN (NM_012293) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: PXDN (NM_012293) Human Tagged ORF Clone Lentiviral Particle

Symbol: PXDN

Synonyms: ASGD7; COPOA; D2S448; D2S448E; MG50; PRG2; PXN; VPO

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

 Tag:
 Myc-DDK

 ACCN:
 NM_012293

ORF Size: 4437 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC224518).

Sequence:

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.

RefSeg: NM 012293.1

 RefSeq Size:
 6821 bp

 RefSeq ORF:
 4440 bp

 Locus ID:
 7837

 UniProt ID:
 Q92626

 Cytogenetics:
 2p25.3

Protein Families: Druggable Genome, Transmembrane

MW: 165.1 kDa







Gene Summary:

This gene encodes a heme-containing peroxidase that is secreted into the extracellular matrix. It is involved in extracellular matrix formation, and may function in the physiological and pathological fibrogenic response in fibrotic kidney. Mutations in this gene cause corneal opacification and other ocular anomalies, and also microphthalmia and anterior segment dysgenesis. [provided by RefSeq, Aug 2014]