

Product datasheet for SC306272

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

Persephin receptor (GFRA4) (NM_145762) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Persephin receptor (GFRA4) (NM_145762) Human Untagged Clone

Tag: Tag Free

Symbol: Persephin receptor

Mammalian Cell

Selection:

None

Vector: pCMV6-XL5

E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene ORF sequence for NM_145762 edited

Restriction Sites: Please inquire **ACCN:** NM_145762

Insert Size: 900 bp

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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OTI Annotation: The open reading frame of this TrueClone was fully sequenced and found to be a perfect

match to the protein associated to this reference.

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: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. 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 145762.1</u>, <u>NP 665705.1</u>

 RefSeq Size:
 1517 bp

 RefSeq ORF:
 900 bp

 Locus ID:
 64096

 UniProt ID:
 Q9GZZ7

 Cytogenetics:
 20p13

Protein Families: Druggable Genome

Gene Summary: The protein encoded by this gene is a member of the GDNF receptor family. It is a

glycosylphosphatidylinositol(GPI)-linked cell surface receptor for persephin, and mediates

activation of the RET tyrosine kinase receptor. This gene is a candidate gene for RET-associated diseases. Alternatively spliced transcript variants encoding different isoforms have

been described for this gene. [provided by RefSeq, Jul 2008]

Transcript Variant: This variant (2) includes additional segments in the coding region compared to variant 1, that cause internal frameshift. The resulting isoform (b) contains additional aa in the middle of the protein, but has the same N- and C-terminal ends as

isoform a.