

## Product datasheet for **SC303226**

### PDGF AA (PDGFA) (NM\_002607) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	PDGF AA (PDGFA) (NM_002607) Human Untagged Clone
Tag:	Tag Free
Symbol:	PDGF AA
Synonyms:	PDGF-A; PDGF1
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>OriGene sequence for NM_002607 edited GCGCCTCGGGACGCGATGAGGACCTTGGCTTGCCTGCTGCTCCTCGGCTGCGGATACCTC GCCCATGTTCTGGCCGAGGAAGCCGAGATCCCCCGGAGGTGATCGAGAGGCTGGCCCCG AGTCAGATCCACAGCATCCGGGACCTCCAGCGACTCCTGGAGATAGACTCCGTAGGGAGT GAGGATTCTTTGGACACCAGCCTGAGAGCTCACGGGGTCCATGCCACTAAGCATGTGCC GAGAAGCGGCCCTGCCATTTCGGAGGAAGAGAAGCATCGAGGAAGTGTCCCCGCTGTC TGCAAGACCAGGACGGTCATTTACGAGATTCTCGGAGTCAGGTCGACCCACGTCCGCC AACTTCTGATCTGGCCCCGTGCGTGGAGGTGAAACGCTGCACCGGCTGCTGCAACAG AGCAGTGTCAAGTGCCAGCCCTCCCGCTCCACCACCGCAGCGTCAAGGTGGCCAAGGTG GAATACGTCAGGAAGAAGCCAAAATTAAGAAGTCCAGGTGAGGTTAGAGGAGCATTTG GAGTGCCTGCGCGACCACAAGCCTGAATCCGGATTATCGGGAAGAGGACCGGAAGG CCTAGGGAGTCAGGTAACGAAAGAAAAGGTTAAACCCACCTAA
Restriction Sites:	Please inquire
ACCN:	NM_002607
Insert Size:	600 bp



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**OTI Disclaimer:** Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at [custsupport@origene.com](mailto:custsupport@origene.com) or by calling 301.340.3188 option 3 for pricing and delivery.

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:** no

**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:** [NM\\_002607.4](#), [NP\\_002598.4](#)

**RefSeq Size:** 2818 bp

**RefSeq ORF:** 636 bp

**Locus ID:** 5154

**UniProt ID:** [P04085](#)

**Cytogenetics:** 7p22.3

**Protein Families:** Druggable Genome

**Protein Pathways:** Cytokine-cytokine receptor interaction, Focal adhesion, Gap junction, Glioma, MAPK signaling pathway, Melanoma, Pathways in cancer, Prostate cancer, Regulation of actin cytoskeleton

**Gene Summary:**

This gene encodes a member of the protein family comprised of both platelet-derived growth factors (PDGF) and vascular endothelial growth factors (VEGF). The encoded preproprotein is proteolytically processed to generate platelet-derived growth factor subunit A, which can homodimerize, or alternatively, heterodimerize with the related platelet-derived growth factor subunit B. These proteins bind and activate PDGF receptor tyrosine kinases, which play a role in a wide range of developmental processes. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2015]

Transcript Variant: This variant (1) represents the longer transcript and encodes the longer isoform (1). The protein contains a C-terminal basic motif encoded by exon 6, the exon missing in variant 2. Sequence Note: This RefSeq record was created from transcript and genomic sequence data because no single transcript was available for the full length of the gene. The extent of this transcript is supported by transcript alignments and experimental evidence.