

Product datasheet for SC330392

OriGene Technologies, Inc.

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Prostaglandin dehydrogenase 1 (HPGD) (NM_001256305) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Prostaglandin dehydrogenase 1 (HPGD) (NM_001256305) Human Untagged Clone

Tag: Tag Free
Symbol: HPGD

Synonyms: 15-PGDH; PGDH; PGDH1; PHOAR1; SDR36C1

Vector: pCMV6-Entry (PS100001)

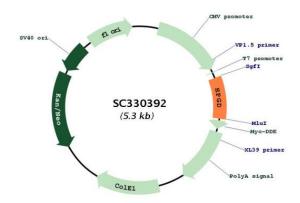
Fully Sequenced ORF: >SC330392 representing NM_001256305.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

TTAGCAGCCCACCATTGA

Restriction Sites: Sgfl-Mlul

Plasmid Map:





ORÏGENE

ACCN: NM_001256305

Insert Size: 432 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).

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 001256305.1</u>

 RefSeq Size:
 2803 bp

 RefSeq ORF:
 432 bp

 Locus ID:
 3248

 UniProt ID:
 P15428

 Cytogenetics:
 4q34.1

Protein Families: Druggable Genome

MW: 15.5 kDa

Gene Summary: This gene encodes a member of the short-chain nonmetalloenzyme alcohol dehydrogenase

protein family. The encoded enzyme is responsible for the metabolism of prostaglandins, which function in a variety of physiologic and cellular processes such as inflammation. Mutations in this gene result in primary autosomal recessive hypertrophic osteoarthropathy

and cranioosteoarthropathy. Multiple transcript variants encoding different isoforms have

been found for this gene. [provided by RefSeq, Mar 2009]

Transcript Variant: This variant (4) lacks two alternate exons in the 3' coding region, which results in a frameshift and early stop codon, compared to variant 1. It encodes isoform 4,

which is shorter and contains a distinct C-terminus, compared to isoform 1.