

Product datasheet for RC206538L3V

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

Glycerol 3 Phosphate Dehydrogenase (GPD1) (NM_005276) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Glycerol 3 Phosphate Dehydrogenase (GPD1) (NM_005276) Human Tagged ORF Clone

Lentiviral Particle

Symbol: Glycerol 3 Phosphate Dehydrogenase

Synonyms: GPD-C; GPDH-C; HTGTI

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM_005276

 ORF Size:
 1047 bp

ORF Nucleotide

Sequence:

The ORF insert of this clone is exactly the same as(RC206538).

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.

RefSeq: <u>NM 005276.2</u>

 RefSeq Size:
 3083 bp

 RefSeq ORF:
 1050 bp

 Locus ID:
 2819

 UniProt ID:
 P21695

Cytogenetics: 12q13.12

Protein Pathways: Glycerophospholipid metabolism





MW: 37.6 kDa

Gene Summary:

This gene encodes a member of the NAD-dependent glycerol-3-phosphate dehydrogenase family. The encoded protein plays a critical role in carbohydrate and lipid metabolism by catalyzing the reversible conversion of dihydroxyacetone phosphate (DHAP) and reduced nicotine adenine dinucleotide (NADH) to glycerol-3-phosphate (G3P) and NAD+. The encoded cytosolic protein and mitochondrial glycerol-3-phosphate dehydrogenase also form a glycerol phosphate shuttle that facilitates the transfer of reducing equivalents from the cytosol to mitochondria. Mutations in this gene are a cause of transient infantile hypertriglyceridemia. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. [provided by RefSeq, Mar 2012]