

Product datasheet for TP720715L

OriGene Technologies, Inc.

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Glycerol 3 Phosphate Dehydrogenase (GPD1) (NM 005276) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Human glycerol-3-phosphate dehydrogenase 1 (soluble)

(GPD1)

Species: Human Expression Host: HEK293

Expression cDNA Clone

or AA Sequence:

Met1-Met349

Tag: C-His

Predicted MW: 38.6 kDa

Purity: >95% as determined by SDS-PAGE and Coomassie blue staining

Buffer: Provided lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl

Endotoxin: Endotoxin level is < 0.1 ng/µg of protein (< 1 EU/µg)

Storage: Store at -80°C.

Stability: Stable for at least 3 months from date of receipt under proper storage and handling

conditions.

RefSeq: NP 005267

Locus ID: 2819

UniProt ID: <u>P21695</u>, <u>A0A024R138</u>

RefSeq Size: 3083

Cytogenetics: 12q13.12

RefSeq ORF: 1047

Synonyms: GPD-C; GPDH-C; HTGTI





Glycerol 3 Phosphate Dehydrogenase (GPD1) (NM_005276) Human Recombinant Protein – TP720715L

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]

Protein Pathways:

Glycerophospholipid metabolism