

Product datasheet for TP720718L

OriGene Technologies, Inc.

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QDPR (NM_000320) Human Recombinant Protein

Ala2-Phe244

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Human quinoid dihydropteridine reductase (QDPR)

Species: Human Expression Host: HEK293

Expression cDNA Clone

or AA Sequence:

: C-His

Tag: C-His
Predicted MW: 26.8 kDa
Concentration: lot specific

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)

Reconstitution Method: Always centrifuge tubes before opening. Do not mix by vortex or pipetting. Dissolve the

lyophilized protein in ddH2O. It is not recommended to reconstitute a concentration less than 100 µg/ml. Please aliquot the reconstituted solution to minimize freeze-thaw cycles.

Storage: Store at -80°C.

Stability: Stable for at least 6 months from date of receipt under proper storage and handling

conditions.

RefSeq: NP 000311

Locus ID: 5860

UniProt ID: <u>P09417</u>, <u>A0A140VKA9</u>

RefSeq Size: 1550

Cytogenetics: 4p15.32

RefSeq ORF: 732

Synonyms: DHPR; HDHPR; PKU2; SDR33C1





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Summary: This gene encodes the enzyme dihydropteridine reductase, which catalyzes the NADH-

mediated reduction of quinonoid dihydrobiopterin. This enzyme is an essential component of the pterin-dependent aromatic amino acid hydroxylating systems. Mutations in this gene resulting in QDPR deficiency include aberrant splicing, amino acid substitutions, insertions, or premature terminations. Dihydropteridine reductase deficiency presents as atypical

premature terminations. Dihydropteridine reductase deficiency presents as atypical phenylketonuria due to insufficient production of biopterin, a cofactor for phenylalanine

hydroxylase. [provided by RefSeq, Jul 2008]

Protein Families: Druggable Genome

Protein Pathways: Folate biosynthesis, Metabolic pathways