## AssayGenie

## Product Data:

Product SKU: RPES2352
Species: Human

Size: $10 \mu \mathrm{~g}$
Expression host: Human Cells

Uniprot: P09417

## Protein Information:

Molecular Mass: $\quad 26.8$ kDa
AP Molecular Mass: 29 kDa
Tag: C-6His
Bio-activity:
Purity: $\quad>95 \%$ as determined by reducing SDS-PAGE.
Endotoxin: $\quad<1.0 \mathrm{EU}$ per $\mu \mathrm{g}$ as determined by the LAL method.
Storage: Lyophilized proteins are stable for up to 12 months when stored at -20 to $-80^{\circ} \mathrm{C}$. Reconstituted protein solution can be stored at $4-8^{\circ} \mathrm{C}$ for 2-7 days. Aliquots of reconstituted samples are stable at $<-20^{\circ} \mathrm{C}$ for 3 months.

Shipping: This product is provided as lyophilized powder which is shipped with ice packs.
Formulation: Lyophilized from a $0.2 \mu \mathrm{~m}$ filtered solution of 20 mM TrisHCl, pH8.0.
Reconstitution: Please refer to the printed manual for detailed information.

## Application:

Synonyms: Dihydropteridine Reductase; HDHPR; Quinoid Dihydropteridine Reductase; QDPR; DHPR

Immunogen information:
Sequence: Ala2-Phe244

## Background:

Dihydropteridine reductase, also known as HDHPR and Quinoid dihydropteridine reductase, QDPR and DHPR, belongs to the short-chain dehydrogenases/reductases (SDR) family. QDPR exists as a homodimer. QDPR is part of the pathway that recycles a substance called tetrahydrobiopterin, also known as BH4 and tryptophan hydroxylases. The regeneration of this substance is critical for the proper processing of several other amino acids in the body. Tetrahydrobiopterin also helps produce certain chemicals in the brain called neurotransmitters, which transmit signals between nerve cells. Defects in QDPR are the cause of BH4deficient hyperphenylalaninemia type $C(H P A B H 4 C)$ which is a rare autosomal recessive disorder and is lethal.

