

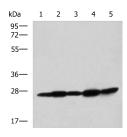
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

# **QDPR Polyclonal Antibody**

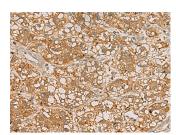
Catalog No.E-AB-52713ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IHC,ELISAIsotypeIgG

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

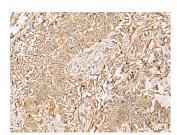
## **Images**



Western blot analysis of Mouse liver tissue Mouse brain tissue Rat brain tissue Rat liver tissue and Human fetal liver tissue lysates using QDPR Polyclonal Antibody at dilution of 1:800



Immunohistochemistry of paraffinembedded Human liver cancer tissue using QDPR Polyclonal Antibody at dilution of 1:70(×200)



Immunohistochemistry of paraffinembedded Human ovarian cancer tissue using QDPR Polyclonal Antibody at dilution of 1:70(×200)

## **Immunogen Information**

Immunogen Fusion protein of human QDPR

**Gene Accession** BC000576 **Swissprot** P09417

**Synonyms** 6,7 dihydropteridine reductase,DHPR,DHPR,HDHPR

,HPR,PKU2,Qdpr,member 1

#### **Product Information**

Calculated MW 26 kDa

**Observed MW** Refer to figures

**Buffer** PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

**Purify** Antigen affinity purification

**Dilution** WB 1:500-1:2000, IHC 1:50-1:200, ELISA

1:5000-1:10000

#### **Background**

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 phenylketonuria due to insufficient production of biopterin, a cofactor for phenylalanine hydroxylase.

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