

## Product datasheet for **TP304694L**

### PAH (NM\_000277) Human Recombinant Protein

#### Product data:

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human phenylalanine hydroxylase (PAH), 1 mg

**Species:** Human

**Expression Host:** HEK293T

**Expression cDNA Clone or AA Sequence:** >RC204694 protein sequence  
**Red**=Cloning site **Green**=Tags(s)

MSTAVLENPGLGRKLSDFGQETSYIEDNCNQNQNGAISLIFSLKEEVGALAKVLRLEFENDVNLTHIESRPS  
RLKKDEYEFFTHLDKRSLPALTNIILRHLDIGATVHELSDKDDKDTVPWFPRTIQELDRFANQILSYGA  
ELDADHPGFKDPVYRARRKQFADIAYNYRHGQPIPRVEYMEEGKKTWGTVEFKTLKSLYKTHACYEYNHIF  
PLLEKYCGFHEDNIPQLEDVVSQFLQTCTGFRLRPVAGLLSSRDFLGGLAFRVFHTQYIRHGSKPMYTP  
PDICHELLGHVPLFSDRSFAQFSQEIGLASLGAPDEYIEKLATYWFVTEFGLCKQGDSIKAYGAGLLSS  
FGELQYCLSEKPKLLPLELEKTAIQNYTVTEFQPLYVAESFNDAKEKVRNFAATIPRPFVRYDPYTQR  
IEVLDNTQQLKILADSINSEIGILCSALQKIK

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV**

**Tag:** C-Myc/DDK

**Predicted MW:** 51.7 kDa

**Concentration:** >0.1 µg/µL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process.

**Storage:** Store at -80°C.

**Stability:** Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles.

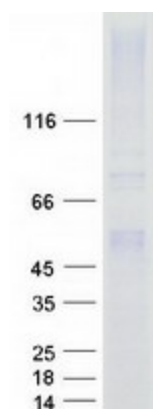
**RefSeq:** [NP\\_000268](#)



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Locus ID:	5053
UniProt ID:	<a href="#">P00439</a> , <a href="#">A0A024RBG4</a>
RefSeq Size:	2680
Cytogenetics:	12q23.2
RefSeq ORF:	1356
Synonyms:	PH; PKU; PKU1
Summary:	This gene encodes a member of the bipterin-dependent aromatic amino acid hydroxylase protein family. The encoded phenylalanine hydroxylase enzyme hydroxylates phenylalanine to tyrosine and is the rate-limiting step in phenylalanine catabolism. Deficiency of this enzyme activity results in the autosomal recessive disorder phenylketonuria. [provided by RefSeq, Aug 2017]
Protein Families:	Druggable Genome
Protein Pathways:	Metabolic pathways, Phenylalanine, tyrosine and tryptophan biosynthesis, Phenylalanine metabolism

### Product images:



Coomassie blue staining of purified PAH protein (Cat# [TP304694]). The protein was produced from HEK293T cells transfected with PAH cDNA clone (Cat# [RC204694]) using MegaTran 2.0 (Cat# [TT210002]).