

## Product datasheet for **TP304694M**

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

#### Product data:

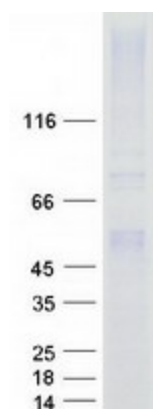
<b>Product Type:</b>	Recombinant Proteins
<b>Description:</b>	Recombinant protein of human phenylalanine hydroxylase (PAH), 100 µg
<b>Species:</b>	Human
<b>Expression Host:</b>	HEK293T
<b>Expression cDNA Clone or AA Sequence:</b>	>RC204694 protein sequence <b>Red</b> =Cloning site <b>Green</b> =Tags(s)
	MSTAVLENPGLGRKLSDFGQETSYIEDNCNQNGAISLIFSLKEEVGALAKVLRLEENDVNLTHIESRPS RLKKDEYEFFTHLDKRSLPALTNIILRHLDIGATVHELSDKKKDTVPWFPRTIQELDRFANQILSYGA ELDADHPGFKDPVYRARRKQFADIAYNYRHGQPIPRVEYMEEGKKTWGTVEFKTLKSLYKTHACYEYNHIF PLLEKYCGFHEDNIPQLEDVVSQFLQTCTGFRLRPVAGLLSSRDFLGGLAFRVFHCTQYIRHGSKPMYTP PDICHELLGHVPLFSDRSFAQFSQEIGLASLGAPDEYIEKLATIWFTVEFGLCKQGDSIKAYGAGLLSS FGELQYCLSEKPKLLPLELEKTAIQNYTVTEFQPLYVAESFNDAKEKVRNFAATIPRPFVRYDPYTOR IEVLDNTQQLKILADSINSEIGILCSALQKIK
	<b>TRTRPLEQKLISEEDLAANDILDYKDDDDKV</b>
<b>Tag:</b>	C-Myc/DDK
<b>Predicted MW:</b>	51.7 kDa
<b>Concentration:</b>	>0.1 µg/µL as determined by microplate BCA method
<b>Purity:</b>	> 80% as determined by SDS-PAGE and Coomassie blue staining
<b>Buffer:</b>	25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol
<b>Preparation:</b>	Recombinant protein was captured through anti-DDK affinity column followed by conventional chromatography steps.
<b>Note:</b>	For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process.
<b>Storage:</b>	Store at -80°C.
<b>Stability:</b>	Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles.
<b>RefSeq:</b>	<u><a href="#">NP_000268</a></u>



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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]).