

# Product datasheet for TP301788M

### PPOX (NM\_000309) Human Recombinant Protein

#### **Product data:**

#### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Froduct data.	
Product Type:	Recombinant Proteins
Description:	Recombinant protein of human protoporphyrinogen oxidase (PPOX), nuclear gene encoding mitochondrial protein, transcript variant 1, 100 μg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC201788 protein sequence <mark>Red</mark> =Cloning site Green=Tags(s)
	MGRTVVVLGGGISGLAASYHLSRAPCPPKVVLVESSERLGGWIRSVRGPNGAIFELGPRGIRPAGALGAR TLLLVSELGLDSEVLPVRGDHPAAQNRFLYVGGALHALPTGLRGLLRPSPPFSKPLFWAGLRELTKPRGK EPDETVHSFAQRRLGPEVASLAMDSLCRGVFAGNSRELSIRSCFPSLFQAEQTHRSILLGLLLGAGRTPQ PDSALIRQALAERWSQWSLRGGLEMLPQALETHLTSRGVSVLRGQPVCGLSLQAEGRWKVSLRDSSLEAD HVISAIPASVLSELLPAEAAPLARALSAITAVSVAVVNLQYQGAHLPVQGFGHLVPSSEDPGVLGIVYDS VAFPEQDGSPPGLRVTVMLGGSWLQTLEASGCVLSQELFQQRAQEAAATQLGLKEMPSHCLVHLHKNCIP QYTLGHWQKLESARQFLTAHRLPLTLAGASYEGVAVNDCIESGRQAAVSVLGTEPNS
	TRTRPLEQKLISEEDLAANDILDYKDDDDKV
Tag:	C-Myc/DDK
Predicted MW:	50.6 kDa
Concentration:	>0.05 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.



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	PPOX (NM_000309) Human Recombinant Protein – TP301788M
RefSeq:	<u>NP 000300</u>
Locus ID:	5498
UniProt ID:	<u>P50336</u>
RefSeq Size:	1716
Cytogenetics:	1q23.3
RefSeq ORF:	1431
Synonyms:	PPO; V290M; VP
Summary: This gene encodes the penultimate enzyme of heme biosynthesis, which catalyze electron oxidation of protoporphyrinogen IX to form protoporphyrin IX. Mutation cause variegate porphyria, an autosomal dominant disorder of heme metabolism from a deficiency in protoporphyrinogen oxidase, an enzyme located on the inne mitochondrial membrane. Alternatively spliced transcript variants encoding the s have been identified. [provided by RefSeq, Jul 2008]	
Protein Families:	Druggable Genome
Protein Pathways	s: Metabolic pathways, Porphyrin and chlorophyll metabolism

## **Product images:**

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Coomassie blue staining of purified PPOX protein (Cat# [TP301788]). The protein was produced from HEK293T cells transfected with PPOX cDNA clone (Cat# [RC201788]) using MegaTran 2.0 (Cat# [TT210002]).

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