

Product datasheet for TP316782L

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

TAT (NM_000353) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Homo sapiens tyrosine aminotransferase (TAT), nuclear gene

encoding mitochondrial protein, 1 mg

Species: Human Expression Host: HEK293T

Expression cDNA >RC216782 protein sequence Clone or AA Sequence: Red=Cloning site Green=Tags(s)

MDPYMIQMSSKGNLSSILDVHVNVGGRSSVPGKMKGRKARWSVRPSDMAKKTFNPIRAIVDNMKVKPNPN

KTMISLSIGDPTVFGNLPTDPEVTQAMKDALDSGKYNGYAPSIGFLSSREEIASYYHCPEAPLEAKDVIL TSGCSQAIDLCLAVLANPGQNILVPRPGFSLYKTLAESMGIEVKLYNLLPEKSWEIDLKQLEYLIDEKTA CLIVNNPSNPCGSVFSKRHLQKILAVAARQCVPILADEIYGDMVFSDCKYEPLATLSTDVPILSCGGLAK RWLVPGWRLGWILIHDRRDIFGNEIRDGLVKLSQRILGPCTIVQGALKSILCRTPGEFYHNTLSFLKSNA DLCYGALAAIPGLRPVRPSGAMYLMVGIEMEHFPEFENDVEFTERLVAEQSVHCLPATCFEYPNFIRVVI

TVPEVMMLEACSRIQEFCEQHYHCAEGSQEECDK

TRTRPLEQKLISEEDLAANDILDYKDDDDK**V**

Tag: C-Myc/DDK
Predicted MW: 50.2 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.





RefSeq: NP 000344

Locus ID: 6898

UniProt ID: <u>P17735</u>, <u>A0A140VKB7</u>

RefSeq Size: 2757
Cytogenetics: 16q22.2
RefSeq ORF: 1362

Summary: This nuclear gene encodes a mitochondrial protein tyrosine aminotransferase which is present

in the liver and catalyzes the conversion of L-tyrosine into p-hydroxyphenylpyruvate. Mutations in this gene cause tyrosinemia (type II, Richner-Hanhart syndrome), a disorder accompanied by major skin and corneal lesions, with possible cognitive disability. A regulator gene for tyrosine

aminotransferase is X-linked. [provided by RefSeq, Jul 2008]

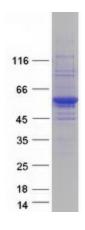
Protein Families: Druggable Genome, Embryonic stem cells, ES Cell Differentiation/IPS

Protein Pathways: Cysteine and methionine metabolism, Metabolic pathways, Phenylalanine, tyrosine and

tryptophan biosynthesis, Phenylalanine metabolism, Tyrosine metabolism, Ubiquinone and

other terpenoid-quinone biosynthesis

Product images:



Coomassie blue staining of purified TAT protein (Cat# [TP316782]). The protein was produced from HEK293T cells transfected with TAT cDNA clone (Cat# [RC216782]) using MegaTran 2.0 (Cat# [TT210002]).