

TPK1 Protein, Human (HEK293, His)

Cat. No.:	HY-P71376
Synonyms:	Thiamin pyrophosphokinase 1; hTPK1; Placental protein 20; PP20; Thiamine pyrophosphokinase 1; TPK1
Species:	Human
Source:	HEK293
Accession:	AAH68460.1 (M1-S243)
Gene ID:	27010
Molecular Weight:	Approximately 30.0 kDa

PROPERTIES

AA Sequence	<pre> MEHAFTPLEP LLSTGNLKYC LVILNQPLDN YFRHLWNKAL LRACADGGAN RLYDITEGER ESFLPEFING DFDSIRPEVR EYYATKGCEL ISTPDQDHTD FTKCLKMLQK KIEEKDLKVD VIVTLGGLAG RFDQIMASVN TLFQATHITP FPIIIQEE S LIYLLQPGKH RLHVDTGMEG DWCGLIPVGGQ PCSQVTTTGL KWNLTNDVLA FGTLVSTSN T YDGS GVV TVE TDHPLLWTMA IKS </pre>
Appearance	Solution.
Formulation	Supplied as a 0.2 µm filtered solution of 20 mM PB, 150 mM NaCl, pH 7.4.
Endotoxin Level	<1 EU/µg, determined by LAL method.
Reconstitution	N/A
Storage & Stability	Stored at -80°C for 1 year. It is stable at -20°C for 3 months after opening. It is recommended to freeze aliquots at -80°C for extended storage. Avoid repeated freeze-thaw cycles.
Shipping	Shipping with dry ice.

DESCRIPTION

Background	<p>Thiamin pyrophosphokinase 1 (TPK1) functions as a homodimer and catalyzes the conversion of thiamine (Vitamin B1) to thiamine pyrophosphate (TDP), a cofactor for some enzymes of the glycolytic and energy production pathways. TDP serves as a transient intermediate carrier of the aldehyde group in the reactions of TDP-dependent enzymes such as pyruvate dehydrogenase, 2-oxoglutarate dehydrogenase, transketolase, and others. TPK1 can also catalyze the phosphorylation of pyrithiamine to pyrithiamine pyrophosphate. Defects in TPK1 gene are a cause of thiamine metabolism dysfunction syndrome-5^[1].</p>
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Caution: Product has not been fully validated for medical applications. For research use only.

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