PANK2 polyclonal antibody

Catalog # PAB2719 Size 400 uL

Applications



Western Blot (Tissue lysate)

Western blot analysis of PANK2 polyclonal antibody (Cat # PAB2719) in mouse liver tissue lysate (35 ug/lane). PANK2 (arrow) was detected using the purified polyclonal abtibody.

Specification	
Product Description	Rabbit polyclonal antibody raised against synthetic peptide of PANK2.
Immunogen	A synthetic peptide (conjugated with KLH) corresponding to amino acids 65-95 at N-terminus of hum an PANK2.
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Protein G purification
Recommend Usage	ELISA Western Blot (1:1000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS (0.09% sodium azide)
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.

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Product Information

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

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Gene Info — PANK2

Entrez GenelD	80025
Protein Accession#	AAN32907;Q9BZ23
Gene Name	PANK2
Gene Alias	C20orf48, FLJ17232, HARP, HSS, MGC15053, NBIA1, PKAN
Gene Description	pantothenate kinase 2
Omim ID	<u>234200 606157 607236</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a protein belonging to the pantothenate kinase family and is the only member of that family to be expressed in mitochondria. Pantothenate kinase is a key regulatory enzyme in the biosynthesis of coenzyme A (CoA) in bacteria and mammalian cells. It catalyzes the first com mitted step in the universal biosynthetic pathway leading to CoA and is itself subject to regulation t hrough feedback inhibition by acyl CoA species. Mutations in this gene are associated with HAR P syndrome and pantothenate kinase-associated neurodegeneration (PKAN), formerly Hallervord en-Spatz syndrome. Alternative splicing, involving the use of alternate first exons, results in multipl e transcripts encoding different isoforms. [provided by RefSeq
Other Designations	Hallervorden-Spatz syndrome OTTHUMP00000030143 OTTHUMP00000030148 pantothenic aci d kinase

Publication Reference

• Compound heterozygous PANK2 mutations confirm HARP and Hallervorden-Spatz syndromes are allelic.

Houlden H, Lincoln S, Farrer M, Cleland PG, Hardy J, Orrell RW. Neurology 2003 Nov; 61(10):1423.

<u>An isoform of hPANK2, deficient in pantothenate kinase-associated neurodegeneration, localizes to</u> <u>mitochondria.</u>

Hortnagel K, Prokisch H, Meitinger T.

Human Molecular Genetics 2003 Feb; 12(3):321.

Pathway

- <u>Metabolic pathways</u>
- Pantothenate and CoA biosynthesis

Disease

- <u>Neurodegenerative Diseases</u>
- Parkinson disease