

Product datasheet for TP308059M

OriGene Technologies, Inc.

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BCKDHB (NM_183050) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human branched chain keto acid dehydrogenase E1, beta polypeptide

(BCKDHB), nuclear gene encoding mitochondrial protein, transcript variant 1, 100 µg

Species: Human
Expression Host: HEK293T

Expression cDNA >RC208059 protein sequence Clone or AA Sequence: Red=Cloning site Green=Tags(s)

MAVVAAAAGWLLRLRAAGAEGHWRRLPGAGLARGFLHPAATVEDAAQRRQVAHFTFQPDPEPREYGQTQK MNLFQSVTSALDNSLAKDPTAVIFGEDVAFGGVFRCTVGLRDKYGKDRVFNTPLCEQGIVGFGIGIAVTG ATAIAEIQFADYIFPAFDQIVNEAAKYRYRSGDLFNCGSLTIRSPWGCVGHGALYHSQSPEAFFAHCPGI KVVIPRSPFQAKGLLLSCIEDKNPCIFFEPKILYRAAAEEVPIEPYNIPLSQAEVIQEGSDVTLVAWGTQ VHVIREVASMAKEKLGVSCEVIDLRTIIPWDVDTICKSVIKTGRLLISHEAPLTGGFASEISSTVQEECF

LNLEAPISRVCGYDTPFPHIFEPFYIPDKWKCYDALRKMINY

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK
Predicted MW: 37.8 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 898871



BCKDHB (NM_183050) Human Recombinant Protein - TP308059M

Locus ID: 594

UniProt ID: <u>P21953</u>, <u>A0A140VKB3</u>

RefSeq Size: 3712 Cytogenetics: 6q14.1 RefSeq ORF: 1176

Synonyms: BCKDE1B; BCKDH E1-beta; E1B

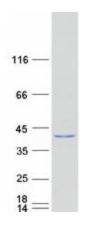
Summary: This gene encodes the E1 beta subunit of branched-chain keto acid dehydrogenase, which is a

multienzyme complex associated with the inner membrane of mitochondria. This enzyme complex functions in the catabolism of branched-chain amino acids. Mutations in this gene have been associated with maple syrup urine disease (MSUD), type 1B, a disease characterized by a maple syrup odor to the urine in addition to mental and physical retardation and feeding problems. Alternative splicing at this locus results in multiple transcript variants. [provided by

RefSeq, Jan 2016]

Protein Pathways: Metabolic pathways, Valine, leucine and isoleucine degradation

Product images:



Coomassie blue staining of purified BCKDHB protein (Cat# [TP308059]). The protein was produced from HEK293T cells transfected with BCKDHB cDNA clone (Cat# [RC208059]) using MegaTran 2.0 (Cat# [TT210002]).