

Product datasheet for TP319389M

OriGene Technologies, Inc.

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ME2 (NM 002396) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human malic enzyme 2, NAD(+)-dependent, mitochondrial (ME2),

nuclear gene encoding mitochondrial protein, 100 µg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone >RC219389 representing NM_002396 or AA Sequence: Red=Cloning site Green=Tags(s)

MLSRLRVVSTTCTLACRHLHIKEKGKPLMLNPRTNKGMAFTLQERQMLGLQGLLPPKIETQDIQALRFHR NLKKMTSPLEKYIYIMGIQERNEKLFYRILQDDIESLMPIVYTPTVGLACSQYGHIFRRPKGLFISISDR GHVRSIVDNWPENHVKAVVVTDGERILGLGDLGVYGMGIPVGKLCLYTACAGIRPDRCLPVCIDVGTDNI ALLKDPFYMGLYQKRDRTQQYDDLIDEFMKAITDRYGRNTLIQFEDFGNHNAFRFLRKYREKYCTFNDDI QGTAAVALAGLLAAQKVISKPISEHKILFLGAGEAALGIANLIVMSMVENGLSEQEAQKKIWMFDKYGLL VKGRKAKIDSYQEPFTHSAPESIPDTFEDAVNILKPSTIIGVAGAGRLFTPDVIRAMASINERPVIFALS

NPTAQAECTAEEAYTLTEGRCLFASGSPFGPVKLTDGRVFTPGQGNNVYIFPGVALAVILCNTRHISDSV FLEAAKALTSQLTDEELAQGRLYPPLANIQEVSINIAIKVTEYLYANKMAFRYPEPEDKAKYVKERTWRS

EYDSLLPDVYEWPESASSPPVITE

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK

Predicted MW: 63.4 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.





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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 002387

Locus ID: 4200

 UniProt ID:
 P23368

 RefSeq Size:
 2730

Cytogenetics: 18q21.2

RefSeq ORF: 1752 Synonyms: ODS1

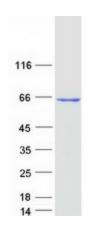
Summary: This gene encodes a mitochondrial NAD-dependent malic enzyme, a homotetrameric protein,

that catalyzes the oxidative decarboxylation of malate to pyruvate. It had previously been weakly linked to a syndrome known as Friedreich ataxia that has since been shown to be the result of mutation in a completely different gene. Certain single-nucleotide polymorphism haplotypes of this gene have been shown to increase the risk for idiopathic generalized epilepsy. Alternatively spliced transcript variants encoding different isoforms found for this

gene. [provided by RefSeq, Dec 2009]

Protein Pathways: Pyruvate metabolism

Product images:



Coomassie blue staining of purified ME2 protein (Cat# [TP319389]). The protein was produced from HEK293T cells transfected with ME2 cDNA clone (Cat# [RC219389]) using MegaTran 2.0 (Cat# [TT210002]).