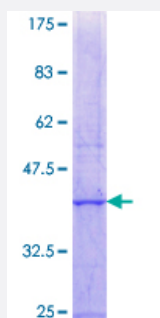


MMAA (Human) Recombinant Protein (Q01)

Catalog # H00166785-Q01

Size 25 ug, 10 ug

Applications



Specification

Product Description	Human MMAA partial ORF (NP_758454.1, 1 a.a. - 110 a.a.) recombinant protein with GST-tag at N-terminal.
Sequence	MPMLLPHPHQHFLKGLLRAPFRCYHFIFHSSTHLGSGIPCAQPFNSLGLHCTKWMLLSDGLKRKL CVQTTLKDHTTEGLSDKEQRFVDKLYTGLIQGQRACLAEAITLVES
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	37.84
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue.
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — MMAA

Entrez GeneID [166785](#)

GeneBank Accession# [NM_172250](#)

Protein Accession# [NP_758454.1](#)

Gene Name MMAA

Gene Alias MGC120010, MGC120011, MGC120012, MGC120013

Gene Description methylmalonic aciduria (cobalamin deficiency) cblA type

Omim ID [251100 607481](#)

Gene Ontology [Hyperlink](#)

Gene Summary The protein encoded by this gene is involved in the translocation of cobalamin into the mitochondrion, where it is used in the final steps of adenosylcobalamin synthesis. Adenosylcobalamin is a coenzyme required for the activity of methylmalonyl-CoA mutase. Defects in this gene are a cause of methylmalonic aciduria. [provided by RefSeq]

Other Designations methylmalonic aciduria type A