

32-4218: Recombinant Human Methylmalonic Aciduria cbID type, with Homocystinuria

Alternative Name : Chromosome 2 Open Reading Frame 25,Methylmalonic Aciduria (Cobalamin Deficiency) CbID Type With Homocystinuria,Methylmalonic Aciduria And Homocystinuria Type D Protein Mitochondrial,Protein C2orf25 Mitochondrial,CL25022,C2orf25,cbID.

Description

Source : Escherichia Coli. MMADHC Human Recombinant produced in E.coli is a single, non-glycosylated polypeptide chain containing 281 amino acids (39-296) and having a molecular mass of 31.0 kDa. MMADHC is fused to a 23 amino acid His-tag at N-terminus. MMADHC is a mitochondrial protein which takes part in an early step of vitamin B12 metabolism. Vitamin B12 (cobalamin) is vital for regular development and existence in humans. Mutations in MMADHC can result in methylmalonic aciduria and homocystinuria type cbID, a cobalamin metabolism syndrome which is characterized by decreased levels of the coenzymes methylcobalamin and adenosylcobalamin.

Product Info

Amount : 20 µg
Purification : Greater than 90% as determined by SDS-PAGE.
Content : The MMADHC solution (1mg/ml) contains 20mM Tris-HCl buffer (pH 8.0), 0.15M NaCl, 1mM DTT and 10% glycerol.
Storage condition : Store at 4°C if entire vial will be used within 2-4 weeks. Store, frozen at -20°C for longer periods of time. For long term storage it is recommended to add a carrier protein (0.1% HSA or BSA). Avoid multiple freeze-thaw cycles.
Amino Acid : MGSSHHHHHH SSGLVPRGSH MGSSDESHVA AAPDICSRT VWPDETMGPF GPQDQRFQLP
GNIGFDCHLN GTASQKKS LV HKTLPDVLA E PLSSERHEFV MAQYVNEFQG NDAPVEQEIN SAETYFESAR
VECAIQTCPE LLRKDFESLF PEVANGK LMI LTVTQTKND MTWVSEEVEI EREVLLEKFI NGAKEICYAL
RAEGYWADFI DPSSGLAFFG PYTNNTLFET DERYRHLGFS VDDLGCCKVI RHSLWGTHVV VGSIFTNATP
DSHIMKKLSG N