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MCEE, 37-176aa, Human, His tag, E.coli

产品货号: TP02927

第三版

别名: Methylmalonyl CoA epimerase, GLOD2.

描述: MCEE catalyzes the interconversion of D- and L-methylmalonyl-CoA during the degradation of branched chain amino acids, odd chain-length fatty acids, and other metabolites. This protein deficiency is an autosomal recessive inborn error of amino acid metabolism, involving valine, threonine, isoleucine and methionine. This organic aciduria may present in the neonatal period with life-threatening metabolic acidosis, hyperammonemia, feeding difficulties, pancytopenia and coma.

配方: Liquid. In 20 mM Tris-HCl buffer (pH8.0) containing 0.2M NaCl, 1mM DTT, 0.1mM PMSF, 10% glycerol

分子量: 17.3 kDa (161aa) confirmed by MALDI-TOF

序列:

MGSSHHHHHHSSGLVPRGSHMQVTGSVWNLGRLNHVAIAVPDLEKAAAFYKNILGAQVSEAVPLPEHGVSVVFNLG  
NTKMELLHPLGRDPIAGFLQKNKAGGMHHICIEVDNINAAMDLKKKIRSLSEEVKIGAHGKPVIFLHPKDCGGVLVE  
LEQA

纯度: > 95% by HPLC

浓度: 1 mg/ml (determined by Bradford assay)

内毒素: <1.0 EU per 1 ug of protein (determined by LAL method)

存储: +4 ° C 保存 (1-2 周). 长期保存在 -20 ° C 或者 -70 ° C. 避免反复冻融.