

Product information

ACP2, 31-380aa

Human, His-tagged, Recombinant, *E.coli*

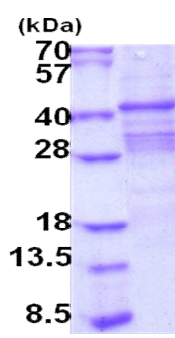
Cat. No. IBATGP2959

Full name: Lysosomal acid phosphatase isoform 1

NCBI Accession No.: NP_001601

Synonyms: LAP

Description: ACP2 also known as Lysosomal acid phosphatase isoform 1. ACP2 is composed of two subunits, alpha and beta, and is chemically and genetically distinct from red cell acid phosphatase. It is a member of a family of distinct isoenzymes which hydrolyze orthophosphoric monoesters to alcohol and phosphate. Acid phosphatase deficiency is caused by mutations in the ACP2 (beta subunit) and ACP3 (alpha subunit) genes. Recombinant human ACP2, fused to His-tag at N-terminus, was expressed in *E.coli*.

<p>Form: Liquid, In 20mM Tris-HCl (pH8.0) containing 10% glycerol</p> <p>Molecular Weight: 42.9kDa (373aa)</p> <p>Purity: > 85% by SDS - PAGE</p> <p>Concentration: 1mg/ml (determined by Bradford assay)</p>	 <p>15% SDS-PAGE (3ug)</p>
<p>Sequences of amino acids:</p> <p>MGSSHHHHH SSGLVPRGSH MGSRLRFVT LLYRHGDRSP VKTYPKDPYQ EEEWPQGFQQ LTKEGMLQHW ELGQALRQRY HGFLNTSYHR QEVYVRSTDF DRTLMSAEAN LAGLFPPNGM QRFNPNISWQ PIPVHTVPIT EDRLKFLPLG PCPRYEQLQN ETRQTPEYQN ESSRNAQFLD MVANETGLTD LTLETVWVNY DTLFCEQTHG LRLPPWASPQ TMQRLSRLKD FSRFLFGIY QQA EKARLQG GVLLAQIRKN LTLMATTSQL PKLLVYSAHD TTLVALQMAL DVYNGEQAPY ASCHIFELYQ EDSGNFSVEM YFRNESDKAP WPLSLPGCPH RCPLQDFLRL TEPVVPKDWQ QEQQLASGPA DTE</p>	

General references:

Moss DW., et al. (1995) *Critical reviews in clinical laboratory sciences*, 32 (4): 431–67

Storage: Can be stored at +4°C short term (1-2 weeks). For long term storage, aliquot and store at -20°C or -70°C.

Avoid repeated freezing and thawing cycles

For research use only. This product is not intended or approved for human, diagnostics or veterinary use.



Manufactured for:
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