

## **Product datasheet for PH310562**

## OriGene Technologies, Inc.

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## Acid Phosphatase 2 (ACP2) (NM\_001610) Human Mass Spec Standard

**Product data:** 

**Product Type:** Mass Spec Standards

**Description:** ACP2 MS Standard C13 and N15-labeled recombinant protein (NP\_001601)

Species:HumanExpression Host:HEK293

**Expression cDNA Clone** 

e RC210562

or AA Sequence:

Predicted MW:

44.45 kDa

Protein Sequence: >RC210562 representing NM\_001610

Red=Cloning site Green=Tags(s)

MAGKRXXGWSRAALLQLLLGVNLVVMPPTRARSLRFVTLLYRHGDRSPVKTYPKDPYQEEEWPQGFGQLT KEGMLQHWELGQALRQRYHGFLNTSYHRQEVYVRSTDFDRTLMSAEANLAGLFPPNGMQRFNPNISWQPI PVHTVPITEDRLLKFPLGPCPRYEQLQNETRQTPEYQNESSRNAQFLDMVANETGLTDLTLETVWNVYDT LFCEQTHGLRLPPWASPQTMQRLSRLKDFSFRFLFGIYQQAEKARLQGGVLLAQIRKNLTLMATTSQLPK LLVYSAHDTTLVALQMALDVYNGEQAPYASCHIFELYQEDSGNFSVEMYFRNESDKAPWPLSLPGCPHRC PLQDFLRLTEPVVPKDWQQECQLASGPADTEVIVALAVCGSILFLLIVLLLTVLFRMQAQPPGYRHVADG

EDHA

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

Concentration: >0.05 µg/µL as determined by microplate BCA method

Labeling Method: Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3

**Storage:** Store at -80°C. Avoid repeated freeze-thaw cycles.

**Stability:** Stable for 3 months from receipt of products under proper storage and handling conditions.

**RefSeq:** <u>NP 001601</u>

RefSeq Size: 2218
RefSeq ORF: 1270
Synonyms: LAP



Locus ID: 53

UniProt ID: P11117

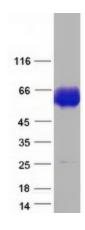
**Cytogenetics:** 11p11.2 | 11p12-p11

**Summary:** The protein encoded by this gene belongs to the histidine acid phosphatase family, which

hydrolyze orthophosphoric monoesters to alcohol and phosphate. This protein is localized to the lysosomal membrane, and is chemically and genetically distinct from the red cell acid phosphatase. Mice lacking this gene showed multiple defects, including bone structure alterations, lysosomal storage defects, and an increased tendency towards seizures. An enzymatically-inactive allele of this gene in mice showed severe growth retardation, hair-follicle abnormalities, and an ataxia-like phenotype. Alternatively spliced transcript variants have been found for this gene. A C-terminally extended isoform is also predicted to be produced by the use of an alternative in-frame translation termination codon via a stop codon readthrough mechanism. [provided by RefSeq, Oct 2017]

Protein Families: Druggable Genome, Transmembrane
Protein Pathways: Lysosome, Riboflavin metabolism

## **Product images:**



Coomassie blue staining of purified ACP2 protein (Cat# [TP310562]). The protein was produced from HEK293T cells transfected with ACP2 cDNA clone (Cat# [RC210562]) using MegaTran 2.0 (Cat# [TT210002]).