

Product datasheet for PH303122

OriGene Technologies, Inc.

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Myosin light chain 3 (MYL3) (NM 000258) Human Mass Spec Standard

Product data:

Product Type: Mass Spec Standards

Description: MYL3 MS Standard C13 and N15-labeled recombinant protein (NP_000249)

Species: Human **HEK293 Expression Host:**

Expression cDNA Clone

or AA Sequence:

RC203122

Predicted MW: 21.9 kDa

>RC203122 protein sequence **Protein Sequence:**

Red=Cloning site Green=Tags(s)

MAPKKPEPKKDDAKAAPKAAPAPAPPPEPERPKEVEFDASKIKIEFTPEQIEEFKEAFMLFDRTPKCEMK ITYGQCGDVLRALGQNPTQAEVLRVLGKPRQEELNTKMMDFETFLPMLQHISKNKDTGTYEDFVEGLRVF

DKEGNGTVMGAELRHVLATLGERLTEDEVEKLMAGQEDSNGCINYEAFVKHIMSS

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

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

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

RefSeq: NP 000249

RefSeg Size: 942 RefSeq ORF: 585

Synonyms: CMH8; MLC-IV/sb; MLC1SB; MLC1V; VLC1; VLCI

Locus ID: 4634

UniProt ID: P08590, A0A024R2Q5





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Cytogenetics: 3p21.31

Summary: MYL3 encodes myosin light chain 3, an alkali light chain also referred to in the literature as

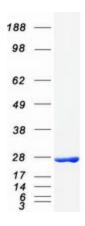
both the ventricular isoform and the slow skeletal muscle isoform. Mutations in MYL3 have been identified as a cause of mid-left ventricular chamber type hypertrophic cardiomyopathy.

[provided by RefSeq, Jul 2008]

Protein Families: Druggable Genome

Protein Pathways: Cardiac muscle contraction, Dilated cardiomyopathy, Hypertrophic cardiomyopathy (HCM)

Product images:



Coomassie blue staining of purified MYL3 protein (Cat# [TP303122]). The protein was produced from HEK293T cells transfected with MYL3 cDNA clone (Cat# [RC203122]) using MegaTran 2.0 (Cat# [TT210002]).