

Product datasheet for TP323199

OriGene Technologies, Inc.

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MBNL1 (NM 207297) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Purified recombinant protein of Homo sapiens muscleblind-like (Drosophila) (MBNL1), transcript Description:

variant 7, 20 µg

Species: Human **Expression Host:** HEK293T

Expression cDNA

Clone or AA Sequence:

Concentration:

>RC223199 representing NM 207297 Red=Cloning site Green=Tags(s)

MAVSVTPIRDTKWLTLEVCREFQRGTCSRPDTECKFAHPSKSCQVENGRVIACFDSLKGRCSRENCKYLH PPPHLKTQLEINGRNNLIQQKNMAMLAQQMQLANAMMPGAPLQPVPMFSVAPSLATNASAAAFNPYLGPV SPSLVPAEILPTAPMLVTGNPGVPVPAAAAAAAQKLMRTDRLEVCREYQRGNCNRGENDCRFAHPADSTM IDTNDNTVTVCMDYIKGRCSREKCKYFHPPAHLQAKIKAAQYQVNQAAAAQAAATAAAMGIPQAVLPPLP

KRPALEKTNGATAVFNTGIFQYQQALANMQLQQHTAFLPPGSILCMTPATSVDTHNICRTSD

TRTRPLEOKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK Predicted MW: 36.9 kDa

>0.05 µg/µL as determined by microplate BCA method

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

Buffer: 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

Preparation: Recombinant protein was captured through anti-DDK affinity column followed by conventional

chromatography steps.

Note: For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Store at -80°C. Storage:

Stability: Stable for 12 months from the date of receipt of the product under proper storage and handling

conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 997180

Locus ID: 4154



MBNL1 (NM_207297) Human Recombinant Protein - TP323199

UniProt ID: Q9NR56, Q86VM6

RefSeq Size: 5277

Cytogenetics: 3q25.1-q25.2

RefSeq ORF: 1026

Synonyms: EXP; MBNL

Summary: This gene encodes a member of the muscleblind protein family which was initially described in

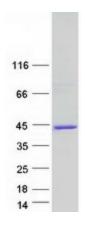
Drosophila melanogaster. The encoded protein is a C3H-type zinc finger protein that modulates alternative splicing of pre-mRNAs. Muscleblind proteins bind specifically to expanded dsCUG RNA

but not to normal size CUG repeats and may thereby play a role in the pathophysiology of

myotonic dystrophy. Mice lacking this gene exhibited muscle abnormalities and cataracts. Several alternatively spliced transcript variants have been described but the full-length natures of only some have been determined. The different isoforms are thought to have different binding

specificities and/or splicing activities. [provided by RefSeq, Sep 2015]

Product images:



Coomassie blue staining of purified MBNL1 protein (Cat# TP323199). The protein was produced from HEK293T cells transfected with MBNL1 cDNA clone (Cat# [RC223199]) using MegaTran 2.0 (Cat# [TT210002]).