

## **Product datasheet for TP319704**

## OriGene Technologies, Inc.

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## MBNL1 (NM\_207295) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human muscleblind-like (Drosophila) (MBNL1), transcript variant 5,

20 µg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone >RC219704 representing NM\_207295

or AA Sequence: Red=Cloning site Green=Tags(s)

MAVSVTPIRDTKWLTLEVCREFQRGTCSRPDTECKFAHPSKSCQVENGRVIACFDSLKGRCSRENCKYLH PPPHLKTQLEINGRNNLIQQKNMAMLAQQMQLANAMMPGAPLQPVVCREYQRGNCNRGENDCRFA

**HPADS** 

TMIDTNDNTVTVCMDYIKGRCSREKCKYFHPPAHLQAKIKAAQYQVNQAAAAQAAATAAAMGIPQAVLP

Ρ

LPKRPALEKTNGATAVFNTGIFQYQQALANMQLQQHTAFLPPGSILCMTPATSVVPMVHGATPATVSAAT

TSATSVPFAATATANQIPIISAEHLTSHKYVTQM

**TRTRPL**EQKLISEEDLAANDILDYKDDDDK**V** 

Tag: C-Myc/DDK

**Predicted MW:** 34 kDa

Concentration: >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.

Storage: Store at -80°C.

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 997178

**Locus ID:** 4154

 UniProt ID:
 Q9NR56

 RefSeq Size:
 5168

**Cytogenetics:** 3q25.1-q25.2

RefSeg ORF: 942

Synonyms: EXP; MBNL

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

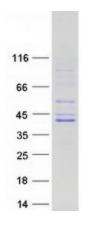
in Drosophila melanogaster. The encoded protein is a C3H-type zinc finger protein that modulates alternative splicing of pre-mRNAs. Muscleblind proteins bind specifically to

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# TP319704). The protein was produced from HEK293T cells transfected with MBNL1 cDNA clone (Cat# [RC219704]) using MegaTran 2.0 (Cat# [TT210002]).