

Product datasheet for **TP323199L**

MBNL1 (NM_207297) Human Recombinant Protein

Product data:

Product Type:	Recombinant Proteins
Description:	Purified recombinant protein of Homo sapiens muscleblind-like (Drosophila) (MBNL1), transcript variant 7, 1 mg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC223199 representing NM_207297 Red =Cloning site Green =Tags(s) MAVSVTPIRDTKWLTLEVCREFQRGTC SRPDTECKFAHPSKSCQVENGRVIACFDSLKGRC SRENCKYLH PPPHLKTQLEINGRNLIQQKNMAMLAQQMQLANAMMPGAPLQPVP MFSVAPSLATNASAAAFNPYLG PV SPSLVPAEILPTAPMLVTGNPGVVPAAAAAAAAA QKLMRTDRLEVCREYQRGNCNRGENDCRFAHPADSTM IDTNDNTVTVCM DYIKGRCSREKCKYFHPPAHLQAKIKAAQYQVNQAAAAQAAATAAAMGIPQAVLPLPLP KRPAL EKTNGATAVFNTGIFQYQQALANMQLQHTAFLPPGSILCMTPATSV DTHNICRTSD TRTRPLEQKLISEEDLAANDILDYKDDDDKV
Tag:	C-Myc/DDK
Predicted MW:	36.9 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:	<u>NP_997180</u>
Locus ID:	4154



[View online »](#)

UniProt ID: [Q9NR56](#), [Q86VM6](#)

RefSeq Size: 5277

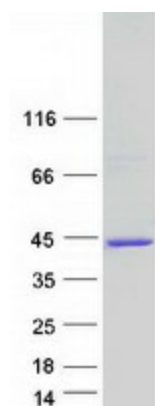
Cytogenetics: 3q25.1-q25.2

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 nature 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]).