# abcam

### Product datasheet

## Anti-MBNL1 antibody ab45899

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Overview	

Anti-MBNL1 antibody
Rabbit polyclonal to MBNL1
Rabbit
Suitable for: IHC-P, ICC/IF, WB
Reacts with: Human
Predicted to work with: Mouse, Rat 🛛 📤
Synthetic peptide corresponding to Human MBNL1 aa 250-350 conjugated to keyhole limpet naemocyanin. Peptide available as <u>ab48114, ab48115</u> )
Recombinant Human MBNL1 protein ( <b>ab114825</b> ) can be used as a positive control in WB. This antibody gave a positive signal in the following Human Lysates: Jurkat Whole Cell Jurkat Nuclear Lysate Human sketelal muscle FFPE tissue sections
The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.
f you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be ound below, along with publications, customer reviews and Q&As

Properties	
Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or - 80°C. Avoid freeze / thaw cycle.
Storage buffer	pH: 7.40 Preservative: 0.02% Sodium azide Constituent: PBS
	Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising agent. If you would like information about the formulation of a specific lot, please contact our scientific support team who will be happy to help.

Purity	Immunogen affinity purified
Clonality	Polyclonal
lsotype	lgG

#### Applications

The Abpromise guarantee

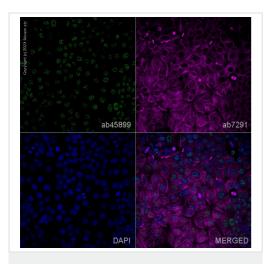
Our **Abpromise guarantee** covers the use of ab45899 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

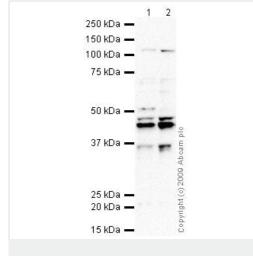
Application	Abreviews	Notes
IHC-P	★★★★	Use a concentration of 5 $\mu\text{g/ml}.$ Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
ICC/IF	★ ★ ★ ★ ★ <u>(2)</u>	Use a concentration of 1 µg/ml.
WB		Use a concentration of 1 $\mu$ g/ml. Detects a band of approximately 35,40,42 kDa (predicted molecular weight: 35,40,42 kDa).

#### Target

Function	Mediates pre-mRNA alternative splicing regulation. Acts either as activator or repressor of splicing on specific pre-mRNA targets. Inhibits cardiac troponin-T (TNNT2) pre-mRNA exon inclusion but induces insulin receptor (IR) pre-mRNA exon inclusion in muscle. Antagonizes the alternative splicing activity pattern of CELF proteins. Regulates the TNNT2 exon 5 skipping through competition with U2AF2. Inhibits the formation of the spliceosome A complex on intron 4 of TNNT2 pre-mRNA. Binds to the stem-loop structure within the polypyrimidine tract of TNNT2 intron 4 during spliceosome assembly. Binds to the 5'-YGCU(U/G)Y-3'consensus sequence. Binds to the IR RNA. Binds to expanded CUG repeat RNA, which folds into a hairpin structure containing GC base pairs and bulged, unpaired U residues.
Tissue specificity	Highly expressed in cardiac, skeletal muscle and during myoblast differentiation. Weakly expressed in other tissues (at protein level). Expressed in heart, brain, placenta, lung, liver, skeletal muscle, kidney and pancreas.
Involvement in disease	Plays a role in the pathogenesis of dystrophia myotonica type 1 (DM1) [MIM:160900]. A muscular disorder characterized by myotonia, muscle wasting in the distal extremities, cataract, hypogonadism, defective endocrine functions, male baldness and cardiac arrhythmias. Note=In muscle cells from DM1 patients, MBNL1 is sequestered by DMPK RNAs containing CUG triplet repeat expansions. MBNL1 binding is proportional to repeat length consistent with the direct correlation between the length of repeat expansion and disease severity.
Sequence similarities	Belongs to the muscleblind family. Contains 4 C3H1-type zinc fingers.
Cellular localization	Nucleus. Cytoplasm. Cytoplasmic granule. Localized with DDX1, TIAL1 and YBX1 in stress granules upon stress. Localized in the cytoplasm of multinucleated myotubes. Colocalizes with nuclear foci of retained expanded-repeat transcripts in myotubes from patients affected by myotonic dystrophy.



Immunocytochemistry/ Immunofluorescence - Anti-MBNL1 antibody (ab45899)



Western blot - Anti-MBNL1 antibody (ab45899)

ab45899 staining MBNL1 in MCF7 cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab45899 at 1µg/ml and <u>ab7291</u>, Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control. Cells were then incubated with <u>ab150081</u>, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor<sup>®</sup> 488), pre-adsorbed at 1/1000 dilution (shown in green) and <u>ab150120</u>, Goat polyclonal Secondary Antibody to Mouse IgG - H&L (Alexa Fluor<sup>®</sup> 594), preadsorbed at 1/1000 dilution (shown in pseudocolour magenta). Nuclear DNA was labelled with DAPI (shown in blue).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal sections is shown.

All lanes : Anti-MBNL1 antibody (ab45899) at 1 µg/ml

Lane 1 : Jurkat (Human T cell lymphoblast-like cell line) Whole Cell Lysate

Lane 2 : Jurkat nuclear extract lysate (ab14844)

Lysates/proteins at 10 µg per lane.

#### Secondary

All lanes : Goat polyclonal to Rabbit lgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

Performed under reducing conditions.

Predicted band size: 35,40,42 kDa Observed band size: 35,40,42 kDa Additional bands at: 110 kDa, 52 kDa. We are unsure as to the identity of these extra bands.

ab45899 recognises all three known isoforms to MBNL1



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-MBNL1 antibody (ab45899)

IHC image of ab45899 staining in human skeletal muscle formalin fixed paraffin embedded tissue section, performed on a Leica Bond<sup>TM</sup> system using the standard protocol F. The section was pretreated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab45899, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

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