# abcam

## Product datasheet

# Anti-FMRP antibody [1D10] ab230915

# 1 References 2 Images

Overview

Product name Anti-FMRP antibody [1D10]

**Description** Mouse monoclonal [1D10] to FMRP

Host species Mouse

Tested applications
Suitable for: IHC-P, WB
Species reactivity
Reacts with: Human

Predicted to work with: Mouse, Rat, Orangutan

Immunogen Recombinant fragment corresponding to Human FMRP aa 1-300. Produced in E. coli.

(NP\_002015).

Database link: Q06787

Run BLAST with Run BLAST with

Positive control WB: Untransfected and pCMV6-ENTRY FMRP-transfected HEK-293T whole cell lysate. IHC-P:

Human lung tissue.

**General notes**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.

If 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

found 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 long

term. Avoid freeze / thaw cycle.

Storage buffer Preservative: 0.02% Sodium azide

Constituents: PBS, 1% BSA, 50% Glycerol (glycerin, glycerine)

Purity Protein A/G purified

**Purification notes** Purified from tissue culture supernatant.

**Clonality** Monoclonal

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Clone number 1D10 lsotype lgG2b

#### **Applications**

#### The Abpromise guarantee

Our Abpromise quarantee covers the use of ab230915 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 10 µg/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
WB		1/500 - 1/2000. Predicted molecular weight: 71 kDa.

#### **Target**

#### **Function**

Translation repressor. Component of the CYFIP1-EIF4E-FMR1 complex which binds to the mRNA cap and mediates translational repression. In the CYFIP1-EIF4E-FMR1 complex this subunit mediates translation repression (By similarity). RNA-binding protein that plays a role in intracellular RNA transport and in the regulation of translation of target mRNAs. Associated with polysomes. May play a role in the transport of mRNA from the nucleus to the cytoplasm. Binds strongly to poly(G), binds moderately to poly(U) but shows very little binding to poly(A) or poly(C).

#### **Tissue specificity**

Highest levels found in neurons, brain, testis, placenta and lymphocytes. Also expressed in

epithelial tissues and at very low levels in glial cells.

#### Involvement in disease

Defects in FMR1 are the cause of fragile X syndrome (FRAX) [MIM:300624]. Fragile X syndrome is a common genetic disease (has a prevalence of one in every 2000 children) which is characterized by moderate to severe mental retardation, macroorchidism (enlargement of the testicles), large ears, prominent jaw, and high-pitched, jocular speech. The defect in most fragile X syndrome patients results from an amplification of a CGG repeat region which is directly in front of the coding region.

Defects in FMR1 are the cause of fragile X tremor/ataxia syndrome (FXTAS) [MIM:300623]. In FXTAS, the expanded repeats range in size from 55 to 200 repeats and are referred to as 'premutations'. Full repeat expansions with greater than 200 repeats results in fragile X mental retardation syndrome [MIM:300624]. Carriers of the premutation typically do not show the full fragile X syndrome phenotype, but comprise a subgroup that may have some physical features of fragile X syndrome or mild cognitive and emotional problems.

Defects in FMR1 are the cause of premature ovarian failure syndrome type 1 (POF1)

[MIM:311360]. An ovarian disorder defined as the cessation of ovarian function under the age of 40 years. It is characterized by oligomenorrhea or amenorrhea, in the presence of elevated levels of serum gonadotropins and low estradiol.

#### Sequence similarities

Belongs to the FMR1 family.
Contains 2 KH domains.

# Post-translational modifications

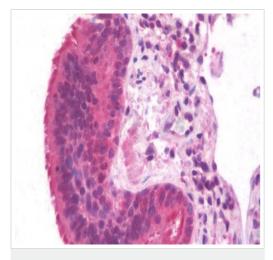
**Cellular localization** 

Phosphorylated on several serine residues.

modifications

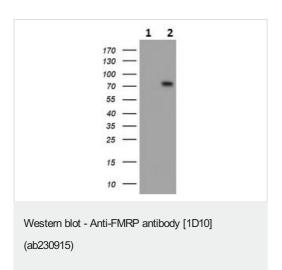
Cytoplasm. Nucleus > nucleolus.

### **Images**



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-FMRP antibody [1D10] (ab230915)

Formalin-fixed, paraffin-embedded human lung tissue stained for FMRP using ab230915 at 10  $\mu$ g/ml in immunohistochemical analysis.



**All lanes :** Anti-FMRP antibody [1D10] (ab230915) at 1/500 dilution

**Lane 1**: pCMV6-ENTRY control-transfected HEK-293T (human epithelial cell line from embryonic kidney transformed with large T antigen) whole cell lysate

**Lane 2**: pCMV6-ENTRY FMRP-transfected HEK-293T whole cell lysate

Lysates/proteins at 5 µg per lane.

Predicted band size: 71 kDa

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