



Anti-FMR1 Antibody

Alternative Names: FMRP, FRAXA, POF, POF1, Synaptic functional regulator FMR1, Fragile X Mental Retardation 1

Catalogue Number: AB18-10051

Size: 100ug

Background Information

Fragile X Mental Retardation 1 (FMR1) is an RNA binding protein that plays a central role in neuronal development and synaptic plasticity through the regulation of alternative mRNA splicing, mRNA stability, mRNA dendritic transport and postsynaptic local protein synthesis of a subset of mRNAs. A trinucleotide repeat (CGG) in the gene encoding this protein is normally found at 6-53 copies, but an expansion to 55-230 repeats is the cause of fragile X syndrome. Expansion of the trinucleotide repeat may also cause one form of premature ovarian failure (POF1). Multiple alternatively spliced transcript variants that encode different protein isoforms and which are located in different cellular locations have also been described for this gene.

FMR1 has been shown to play a translation-independent role in the modulation of presynaptic action potential (AP) duration and neurotransmitter release via large-conductance calcium-activated potassium (BK) channels in hippocampal and cortical excitatory neurons [1]. FMR1 may be involved in the control of DNA damage response (DDR) mechanisms through the regulation of ATR-dependent signaling pathways such as histone H2AFX/H2A.x and BRCA1 phosphorylations [2].

Product Information

Antibody Type:	Polyclonal	Host:	Rabbit
Isotype:	IgG	Species Reactivity:	Human Mouse Rat
Immunogen:	Recombinant human FMR1		
Format:	100 µg in 100 µl PBS containing 0.02% sodium azide.		
Storage Conditions:	6 months: 4°C. Long-term storage: -20°C. Avoid multiple freeze and thaw cycles.		
Applications:	WB IHC IF WB 1:500-2000. IHC 1:50-200. IF 1:50-200.		

Additional Information

Subcellular location:	Cytoplasm, Nucleus, nucleolus	MW:	71kDa
Gene ID	2332	Uniprot ID:	Q06787



References

1. Myrick L.K. et al. Proc. Natl. Acad. Sci. U.S.A. 112:949-956(2015),
 2. Alpatov R. et al Cell 157:869-881(2014)
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