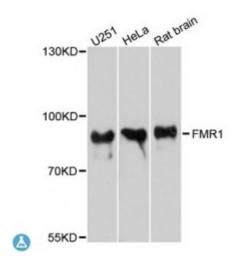


Anti-FMR1 Antibody



Description The protein encoded by this gene binds RNA and is associated with

polysomes. The encoded protein may be involved in mRNA trafficking from the nucleus to the cytoplasm. A trinucleotide repeat (CGG) in the 5' UTR 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 been

described for this gene.

Model STJ113567

Host Rabbit

Reactivity Human, Rat

Applications WB

Immunogen A synthetic peptide corresponding to a sequence within amino acids 500-600

of human FMR1 (NP_002015.1).

Gene ID 2332

Gene Symbol FMR1

Dilution range WB 1:500 - 1:1000

Tissue Specificity Expressed in the brain, cerebellum and testis

Purification Affinity purification

Note For Research Use Only (RUO).

Protein Name Synaptic functional regulator FMR1 Fragile X mental retardation protein 1

FMRP Protein FMR-1

Molecular Weight 71.174 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Storage Instruction Store at -20C. Avoid freeze / thaw cycles.

Database Links HGNC:37750MIM:300623

Alternative Names Synaptic functional regulator FMR1 Fragile X mental retardation protein 1

FMRP Protein FMR-1

Function Multifunctional polyribosome-associated 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

components,

Cellular Localization Nucleus, Isoform 11: Nucleus

Post-translational Phosphorylated,

Modifications

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