FXR1 (D10A2) XP® Rabbit mAb

Small 100 µl (10 western blots)

Petite 40 ul (4 western blots)

Cell Signaling

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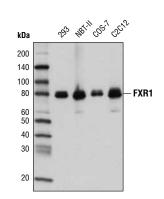
rev. 01/05/15

For Research Use Only. Not For Use In Diagnostic Procedures.

Applications Species Cross-Reactivity* Molecular Wt. Isotype W. IF-IC H. M. R. Mk 78-80. 82-84 kDa Rabbit IgG** Endogenous

Background: Fragile X syndrome is a genetic disorder characterized by a spectrum of physical and behavioral features and is a frequent form of inherited mental retardation (1). X-linked FMRP (FMR-1) and its two autosomal homologs, FXR1 and FXR2, are polyribosome-associated RNA-binding proteins that are involved in the pathogenesis of fragile X syndrome (1-3). Each of the fragile X proteins can self-associate, as well as form heteromers with the other two related proteins (3). FMRP can act as a translation regulator and is a component of RNAi effector complexes (RISC), suggesting a role in gene silencing (4). The Drosophila homolog of FMRP (dFMRP) associates with Argonaute 2 (Ago2) and Dicer and can coimmunoprecipitate with miRNA and siRNA (5). These results suggest that fragile X syndrome is related to abnormal translation caused by defects in RNAi-related pathways. In addition, FMRP. FXR1, and FXR2 are components of stress granules (SG) and have been implicated in the translational regulation of mRNAs (6)

Specificity/Sensitivity: FXR1 (D10A2) XP® Rabbit mAb recognizes endogenous levels of total FXR1 protein.



Western blot analysis of extracts from various cell lines using FXR1 (D10A2) XP® Rabbit mAb.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly574 of human FXR1 protein.

Entrez-Gene ID #8087 UniProt ID #P51114

Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

*Species cross-reactivity is determined by western blot.

**Anti-rabbit secondary antibodies must be used to detect this antibody.

Recommended Antibody Dilutions:

1:1000 Western blotting Immunofluorescence (IF-IC) 1:50

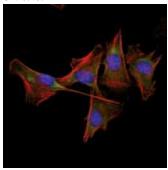
For product specific protocols please see the web page for this product at www.cellsignal.com.

Please visit www.cellsignal.com for a complete listing of recommended companion products.

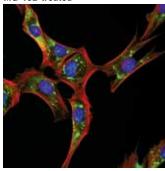
Background References:

- (1) Verkerk, A.J. et al. (1991) Cell 65, 905-14.
- (2) Siomi, M.C. et al. (1995) EMBO J 14, 2401-8.
- (3) Zhang, Y. et al. (1995) EMBO J 14, 5358-66.
- (4) Caudy, A.A. et al. (2002) Genes Dev 16, 2491-6.
- (5) Siomi, H. et al. (2004) Ment Retard Dev Disabil Res Rev 10, 68-74.
- (6) Linder, B. et al. (2008) Hum Mol Genet 17, 3236-46.

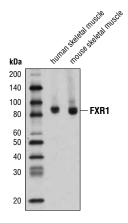
Untreated



MG-132-treated



Confocal immunofluorescent analysis of C2C12 cells, untreated (left) or MG-132 treated (10 µg/ mL, 3 hr; right), using FXR1 (D10A2) XP® Rabbit mAb (green). Actin filaments were labeled with DyLight™ 554 Phalloidin #13054 (red). Blue pseudocolor = DRAQ5® #4084 (fluorescent DNA dye).



Western blot analysis of extracts from human and mouse skeletal muscle tissues using FXR1 (D10A2) XP® Rabbit mAb.

IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v BSA, 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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IF-Immunofluorescence Applications Key: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation F—Flow cytometry E-P—ELISA-Peptide Species Cross-Reactivity Key: H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenogus Z—zebrafish

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