

Product Data Sheet

Anti-MEF2D (Phospho-S444) Antibody

Catalog # Source Reactivity Applications

CPA4462 Rabbit H, M, R, B, P WB, IH

Description Rabbit polyclonal antibody to MEF2D (Phospho-S444)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S444 of human MEF2D protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of MEF2D protein only when phosphorylated at S444.

Clonality Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IH (1/100 - 1/200)

Gene Symbol MEF2D

Alternative Names Myocyte-specific enhancer factor 2D

Entrez Gene 4209 (Human); 17261 (Mouse); 81518 (Rat)

SwissProt Q14814 (Human); Q63943 (Mouse); O89038 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

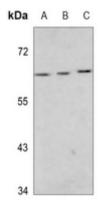
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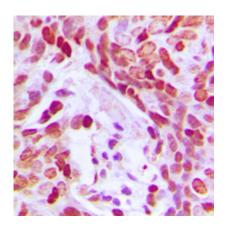




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Western blot analysis of MEF2D (Phospho-S444) expression in HEK293T (A), Jurkat (B), U2OS (C) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 70 kD)



Immunohistochemical analysis of MEF2D (Phospho-S444) staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody 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.

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