

Product Data Sheet

Anti-SMAD3 Antibody

Catalog # Source Reactivity Applications

CPA4246 Rabbit H, M, R, D, Mk, P, S WB, IH

Description Rabbit polyclonal antibody to SMAD3

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human SMAD3. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of SMAD3 protein.

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 SMAD3

Alternative Names MADH3; Mothers against decapentaplegic homolog 3; MAD homolog 3; Mad3;

Mothers against DPP homolog 3; hMAD-3; JV15-2; SMAD family member 3; SMAD 3;

Smad3: hSMAD3

Entrez Gene 4088 (Human); 17127 (Mouse); 25631 (Rat)

SwissProt P84022 (Human); Q8BUN5 (Mouse); P84025 (Rat)

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

freeze/thaw cycles.

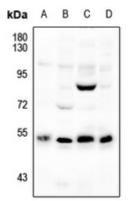
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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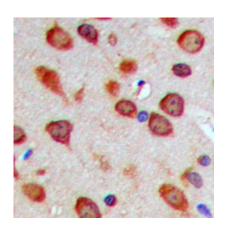
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Western blot analysis of SMAD3 expression in MG63 (A), Myla2059 (B), HCT116 (C), A549 (D) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 52 kD)



Immunohistochemical analysis of SMAD3 staining in human brain 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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