

Product Data Sheet

Anti-SMAD3 (Phospho-S204) Antibody

Catalog # Source Reactivity Applications

CPA1708 Rabbit H, M, R, C, Mk, P, S WB, IH, IF/IC

Description Rabbit polyclonal antibody to SMAD3 (Phospho-S204)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S204 of human SMAD3 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of SMAD3 protein only when phosphorylated at S204.

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 (1/50 - 1/100), IF/IC (1/50 - 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.

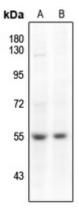
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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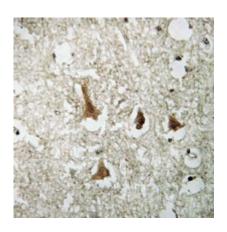
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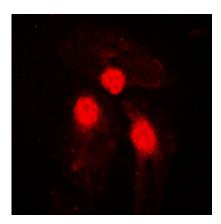
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Western blot analysis of SMAD3 (Phospho-S204) expression in HCT116 (A), A549 (B) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 52 kD)



Immunohistochemical analysis of SMAD3 (Phospho-S204) 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.



Immunofluorescent analysis of SMAD3 (Phospho-S204) staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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