

Product Data Sheet

Anti-SMAD2 (Phospho-S465) Antibody

Catalog # Source Reactivity Applications

CPA4244 Rabbit H, M, R, B, C, Mk, P, Rb, S, Z WB, IH

Description Rabbit polyclonal antibody to SMAD2 (Phospho-S465)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S465 of human SMAD2 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of SMAD2 protein only when phosphorylated at S465.

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 SMAD2

Alternative Names MADH2; MADR2; Mothers against decapentaplegic homolog 2; MAD homolog 2;

Mothers against DPP homolog 2; JV18-1; Mad-related protein 2; hMAD-2; SMAD

family member 2; SMAD 2; Smad2; hSMAD2

Entrez Gene 4087 (Human); 17126 (Mouse); 29357 (Rat)

SwissProt Q15796 (Human); Q62432 (Mouse); O70436 (Rat)

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

freeze/thaw cycles.

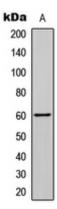
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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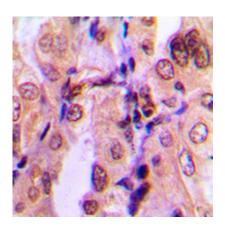
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Western blot analysis of SMAD2 (Phospho-S465) expression in HepG2 (A) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 60 kD)



Immunohistochemical analysis of SMAD2 (Phospho-S465) 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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