

Product Data Sheet

Anti-MST1/2 (Phospho-T183) Antibody

Catalog # Source Reactivity Applications

CPA4262 Rabbit H, M, R, B, C, P, Z WB, IH

Description Rabbit polyclonal antibody to MST1/2 (Phospho-T183)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

T183 of human MST1/2 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of MST1/2 protein only when phosphorylated at T183.

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 STK3; STK4

Alternative Names STK3; KRS1; MST2; Serine/threonine-protein kinase 3; Mammalian STE20-like

protein kinase 2; MST-2; STE20-like kinase MST2; Serine/threonine-protein kinase

Krs-1; STK4; KRS2; MST1; Serine/threonine-protein kinase 4; Mammalian STE20-like

protein kinase 1; M

Entrez Gene 6789 (Human); 56274, 58231 (Mouse); 65189 (Rat)

SwissProt Q13188, Q13043 (Human); Q9JI10, Q9JI11 (Mouse); O54748 (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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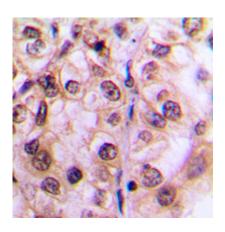
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Western blot analysis of MST1/2 (Phospho-T183) expression in HEK293T (A) whole cell lysates. (Predicted band size: 56; 55 kD; Observed band size: 65 kD)



Immunohistochemical analysis of MST1/2 (Phospho-T183) 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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