

### **Product Data Sheet**

# Anti-c-Met (Phospho-Y1349) Antibody

Catalog # Source Reactivity Applications

CPA4463 Rabbit H, M, R, B, P WB, IH, IF/IC

**Description** Rabbit polyclonal antibody to c-Met (Phospho-Y1349)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

Y1349 of human c-Met protein. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of c-Met protein only when phosphorylated at Y1349.

**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 MET

Alternative Names Hepatocyte growth factor receptor; HGF receptor; HGF/SF receptor; Proto-oncogene

c-Met; Scatter factor receptor; SF receptor; Tyrosine-protein kinase Met

**Entrez Gene** 4233 (Human); 24553 (Rat)

SwissProt P08581 (Human); P16056 (Mouse); P97523 (Rat)

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

freeze/thaw cycles.

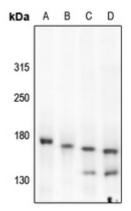
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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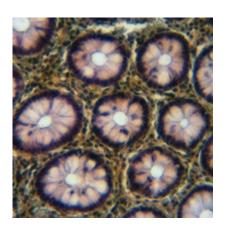
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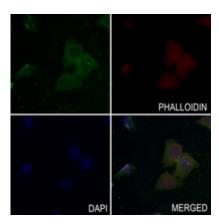
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Western blot analysis of c-Met (Phospho-Y1349) expression in CT26 (A), rat liver (B), HEK293T (C), LO2 (D) whole cell lysates. (Predicted band size: 155 kD; Observed band size: 145; 170 kD)



Immunohistochemical analysis of c-Met (Phospho-Y1349) staining in human colorectal 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.



Immunofluorescent analysis of c-Met (Phospho-Y1349) staining in LO2 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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