

# **Product Data Sheet**

# **Anti-EMP Antibody**

Catalog # Source Reactivity Applications

CPA3379 Rabbit H, M, R, B, C, Mk WB, IH

**Description** Rabbit polyclonal antibody to EMP

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

region of human EMP. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of EMP 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 MAEA

Alternative Names EMP; Macrophage erythroblast attacher; Cell proliferation-inducing gene 5 protein;

Erythroblast macrophage protein; Human lung cancer oncogene 10 protein; HLC-10

**Entrez Gene** 10296 (Human); 59003 (Mouse); 298982 (Rat)

SwissProt Q7L5Y9 (Human); Q4VC33 (Mouse); Q5RKJ1 (Rat)

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

freeze/thaw cycles.

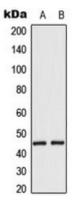
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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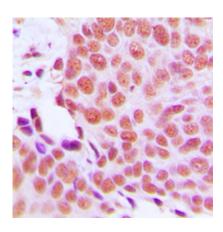
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Western blot analysis of EMP expression in HepG2 (A), HeLa (B) whole cell lysates. (Predicted band size: 45 kD; Observed band size: 45 kD)



Immunohistochemical analysis of EMP 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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