

Product Data Sheet

Anti-Midkine Antibody

Catalog # Source Reactivity Applications

CPA1732 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to Midkine

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

region of human Midkine. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Midkine 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 (1/50 - 1/100)

Gene Symbol MDK

Alternative Names MK1; NEGF2; Midkine; MK; Amphiregulin-associated protein; ARAP; Midgestation

and kidney protein; Neurite outgrowth-promoting factor 2; Neurite

outgrowth-promoting protein

Entrez Gene 4192 (Human); 17242 (Mouse); 81517 (Rat)

SwissProt P21741 (Human); P12025 (Mouse); Q9R1S9 (Rat)

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

freeze/thaw cycles.

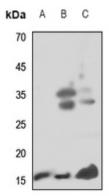
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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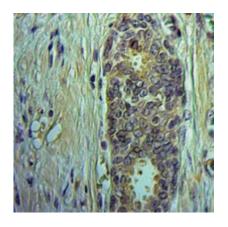
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Western blot analysis of Midkine expression in mouse heart (A), rat liver (B), rat heart (C) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 15 kD)



Immunohistochemical analysis of Midkine staining in human prostate 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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