

Product Data Sheet

Anti-CXCL1 Antibody

Catalog # Source Reactivity Applications

CPA4952 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to CXCL1

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human CXCL1. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of CXCL1 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 CXCL1

Alternative Names GRO; GRO1; GROA; MGSA; SCYB1; Growth-regulated alpha protein; C-X-C motif

chemokine 1; GRO-alpha(1-73); Melanoma growth stimulatory activity; MGSA;

Neutrophil-activating protein 3; NAP-3

Entrez Gene 2919 (Human)

SwissProt P09341 (Human)

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

freeze/thaw cycles.

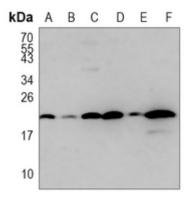
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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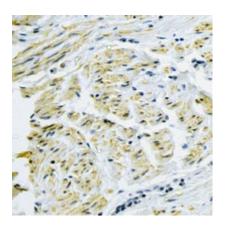
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Western blot analysis of CXCL1 expression in HEK293T (A), H1688 (B), HCT116 (C), mouse spleen (D), mouse lung (E), rat spleen (F) whole cell lysates. (Predicted band size: 11 kD; Observed band size: 20 kD)



Immunohistochemical analysis of CXCL1 staining in human lung 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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