#### **Anti-CXCL12 Rabbit Polyclonal Antibody**



## **Catalog #: 50631**

#### **Aliases**

SDF1; SDF1A; SDF1B; Stromal cell-derived factor 1; SDF-1; hSDF-1; C-X-C motif chemokine 12; Intercrine reduced in hepatomas; IRH; hIRH; Pre-B cell growth-stimulating factor; PBSF

# **Background**

Gene Name: CXCL12 NCBI Gene Entry: 6387 UniProt Entry: P48061

# **Application Information**

Molecular Weight: Predicted, 10 kDa; observed, 10 kDa

Clonality: Rabbit polyclonal antibody Species Reactivity: Human, dog

Applications Tested: Western blotting (WB), immunohistochemistry (IHC), immunocytochemistry

(IC)

## **Immunogen**

A synthesized peptide derived from human CXCL12

## **Isotype**

Rabbit IgG

# **Storage Buffer**

Supplied in PBS (pH 7.3) containing 30% glycerol, and 0.01% sodium azide.

## **Storage**

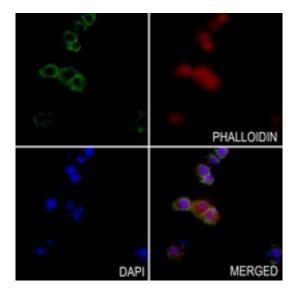
Store at -20 °C for one year.

#### **Recommended Dilutions**

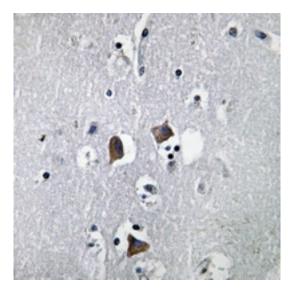
Western Blotting (WB): 1:500-1:1,000 Immunohistochemistry (IHC): 1:50-1:100 Immunocytochemistry (IC): 1:50-1:200

**Note:** This product is for research use only.

#### Validation Data

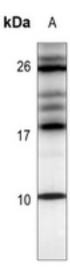


Immunocytochemical analysis of CXCL12 staining in Hela cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C 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).



Immunohistochemical analysis of CXCL12 staining in human brain 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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Western blotting analysis of CXCL12 expression in Hela (A) whole cell lysates.