Anti-CXCR4 Antibody

ER1802-28



Product Type: Rabbit polyclonal IgG, primary antibodies

Species reactivity: Human, Rat

Applications: WB, IF-Cell, FC

Molecular Wt: 45 kDa

Description: The C-X-C or a chemokine family is characterized by a pair of cysteine residues separated

by a single amino acid and primarily functions as chemoattractants for neutrophils. The C-X-C family includes IL-8, NAP-2, MSGA and stromal cell derived factor-1 (SDF-1). SDF-1 was originally described as a pre-B cell stimulatory factor, but has now been shown to function as a potent chemo-attractant for T cells and monocytes but not neutrophils. Receptors for the C-X-C family are G protein-coupled, seven pass transmembrane domain proteins which include IL-8RA, IL-8RB and CXCR-4 (also known as LESTR or fusin). CXCR-4 is highly homologous to the IL-8 receptors, sharing 37% sequence identity at the amino acid level. The IL-8 receptors bind to IL-8, NAP-2 and MSGA, while fusin binds to its cognate ligand, SDF-1. CXCR-4 has been identified as the major co-receptor for T-tropic HIV-1 and SDF-1

has been shown to inhibit HIV-1 infection.

Immunogen: Synthetic peptide within N-terminal of human CXCR4.

Positive control: 293, K562, Hela, HUVEC, THP-1.

Subcellular location: Cell junction. Plasma membrane. Endosome. Lysosome.

Database links: SwissProt: P61073 Human | 008565 Rat

Recommended Dilutions:

WB 1:500-1:1,000
IF-Cell 1:500-1:2,000
FC 1:50-1:100

Storage Buffer: 1*PBS (pH7.4), 0.2% BSA, 50% Glycerol. Preservative: 0.05% Sodium Azide.

Storage Instruction: Shipped at 4° C. Store at $+4^{\circ}$ C short term (1-2 weeks). It is recommended to aliquot into

single-use upon delivery. Store at -20 ℃ long term.

Purity: Immunogen affinity purified.

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Images

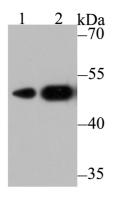


Fig1: Western blot analysis of CXCR4 on different cell lysates using anti-CXCR4 antibody at 1/500 dilution.

Positive control:

Lane 1: 293 Lane 2: K562

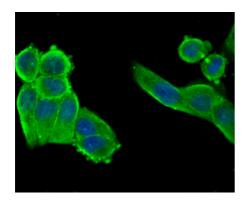


Fig2: ICC staining CXCR4 in Hela cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

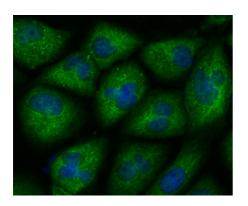


Fig3: ICC staining CXCR4 in HUVEC cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

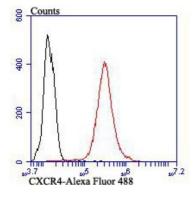


Fig4: Flow cytometric analysis of THP-1 cells with CXCR4 antibody at 1/100 dilution (red) compared with an unlabelled control (cells without incubation with primary antibody; black). Alexa Fluor 488-conjugated goat anti-rabbit IgG was used as the secondary antibody.

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