## Anti-IL-8 Antibody [PSH10-96] - BSA and Azide free HA751386



Species reactivity: Human
Applications: WB, FC

Molecular Wt: Predicted band size: 11 kDa

Clone number: PSH10-96

**Description:** Interleukin 8 (IL-8 or chemokine (C-X-C motif) ligand 8, CXCL8) is a chemokine produced

by macrophages and other cell types such as epithelial cells, airway smooth muscle cells and endothelial cells. Endothelial cells store IL-8 in their storage vesicles, the Weibel–Palade bodies. In humans, the interleukin-8 protein is encoded by the CXCL8 gene. IL-8 is initially produced as a precursor peptide of 99 amino acids which then undergoes cleavage to create several active IL-8 isoforms. In culture, a 72 amino acid peptide is the major form secreted by macrophages. There are many receptors on the surface membrane capable of binding IL-8; the most frequently studied types are the G protein-coupled serpentine receptors CXCR1 and CXCR2. Expression and affinity for IL-8 differs between the two receptors (CXCR1 > CXCR2). Through a chain of biochemical reactions, IL-8 is secreted and is an important mediator of the immune reaction in the innate immune system response.

**Immunogen:** Recombinant protein within human IL-8 aa 1-99.

Positive control: U-937 treated with 100ng/mL TPA for 24 hours then add 5µg/mL LPS for 4 hours then add

300ng/mL BFA for 3 hours cell lysate, U-937 treated with 100ng/mL TPA for 24 hours then

add 5µg/mL LPS for 4 hours then add 300ng/mL BFA for 3 hours.

Subcellular location: Secreted.

Database links: SwissProt: P10145 Human

Recommended Dilutions:

**WB** 1:2,000 **FC** 1:1,000

**Storage Buffer:** PBS (pH7.4).

**Storage Instruction:** Store at  $+4^{\circ}$ C after thawing. Aliquot store at  $-20^{\circ}$ C. Avoid repeated freeze / thaw cycles.

**Purity:** Protein A affinity purified.

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## **Images**

 Fig1: Western blot analysis of IL-8 on different lysates with Rabbit anti-IL-8 antibody (HA751386) at 1/2,000 dilution.

Lane 1: U-937 cell lysate

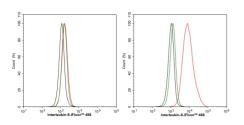
Lane 2: U-937 treated with 100ng/mL TPA for 24 hours then add 5 $\mu$ g/mL LPS for 4 hours then add 300ng/mL BFA for 3 hours cell lysate

Lysates/proteins at 10 µg/Lane.

Predicted band size: 11 kDa Observed band size: 11 kDa

Exposure time: 3 minutes; ECL: K1801;

4-20% SDS-PAGE gel.



**Fig2:** Flow cytometric analysis of U-937 cells untreated (left) / treated with 100ng/mL TPA for 24 hours then add  $5\mu$ g/mL LPS for 4 hours then add 300ng/mL BFA for 3 hours (right) labeling IL-8.

Cells were fixed and permeabilized. Then stained with the primary antibody (HA751386, 1/1,000) (red) compared with Rabbit IgG Isotype Control (green). After incubation of the primary antibody at +4  $^{\circ}$ C for an hour, the cells were stained with a iFluor <sup>TM</sup> 488 conjugate-Goat anti-Rabbit IgG Secondary antibody (HA1121) at 1/1,000 dilution for 30 minutes at +4  $^{\circ}$ C. Unlabelled sample was used as a control (cells without incubation with primary antibody; black).

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

## **Background References**

- 1. Ramachandra N et al. Role of IL8 in myeloid malignancies. Leuk Lymphoma. 2023 Nov-Dec
- 2. Jiang H et al. Targeting IL8 as a sequential therapy strategy to overcome chemotherapy resistance in advanced gastric cancer. Cell Death Discov. 2022 Apr

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