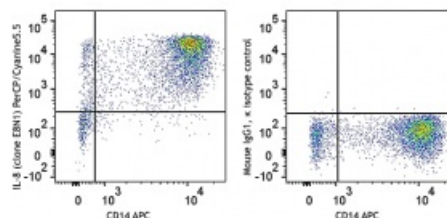


# PerCP/Cyanine5.5 anti-human IL-8

<b>Catalog # /</b>	3157110 / 100 tests
<b>Size:</b>	3157105 / 25 tests
<b>Clone:</b>	E8N1
<b>Isotype:</b>	Mouse IgG1, $\kappa$
<b>Immunogen:</b>	Human TRAIL-transfected mouse cell line
<b>Reactivity:</b>	Human
<b>Preparation:</b>	The antibody was purified by affinity chromatography and conjugated with PerCP/Cyanine5.5 under optimal conditions.
<b>Formulation:</b>	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA)
<b>Workshop Number:</b>	HCDM listed
<b>Concentration:</b>	lot-specific



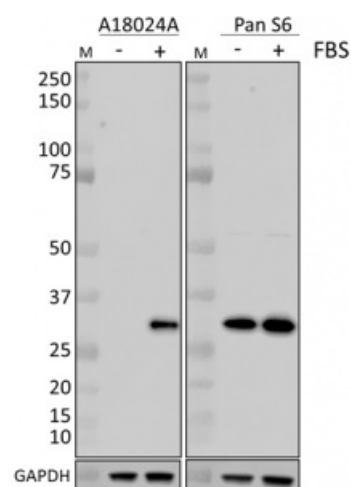
LPS-stimulated (6 hours, in the presence of monensin) human peripheral blood monocytes were surface stained with CD14 APC and then intracellularly stained with IL-8 (clone E8N1) PerCP/Cyanine5.5 (left) or mouse IgG1,  $\kappa$  PerCP/Cyanine5.5 isotype control (right).

## Applications:

<b>Applications:</b>	Intracellular Flow Cytometry
<b>Recommended Usage:</b>	Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is 5 $\mu$ L per million cells in 100 $\mu$ L staining volume or 5 $\mu$ L per 100 $\mu$ L of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

<b>Application Notes:</b>	<b>ELISA Detection:</b> The biotinylated E8N1 antibody is useful as the detection antibody in a sandwich ELISA assay, when used in conjunction with the purified H8A5 antibody as the capture antibody.
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\* PerCP/Cyanine5.5 has a maximum absorption of 482 nm and a maximum emission of 690 nm.



Total cell lysates (15  $\mu$ g protein) from serum-starved NIH/3T3 cells treated without (-) or with (+) 20% FBS for 30 minutes were resolved by 4-12% Bis-Tris gel electrophoresis, transferred to a PVDF membrane, and probed with 0.25  $\mu$ g/mL (1:2000 dilution) of purified anti-RPS6 Phospho (Ser244) antibody (clone A18024A). Proteins were visualized by chemiluminescence detection using HRP goat anti-mouse IgG antibody at a 1:3000 dilution. Equal protein loading was confirmed using a purified anti-RPS6 antibody and Direct-Blot™ HRP anti-GAPDH antibody used at a 1:25000 dilution.

(lower). Lane M: molecular weight ladder.

- Application References:**
1. Kayagaki N, *et al.* 1999. *J. Immunol.* 162:2639. (Block)
  2. Uno K, *et al.* 2003. *Blood* 101:3658. (Block)
  3. Sato K, *et al.* 2005. *J. Immunol.* 174:4025. (Block)
  4. Denny MF, *et al.* 2007. *Blood* 110:2907. (Block)
  5. Kemter E, *et al.* 2011. *Xenotransplantation.* 19:40. [PubMed](#)
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**Description:** IL-8, also known as neutrophil chemotactic factor, neutrophil activating protein, and monocyte-derived neutrophil chemotactic factor, is a member of the alpha (C-X-C) subfamily of chemokines called CXCL8. In response to proinflammatory stimuli, IL-8 is produced by monocytes, macrophages, T cells, neutrophils, and fibroblasts. IL-8 promotes neutrophil chemotaxis and degranulation. The 72 amino acid IL-8 is the predominant form secreted by monocytes and lymphocytes. The E8N1 antibody recognizes the human IL-8 protein and has been shown to be useful for intracellular immunofluorescence flow cytometric analysis and as ELISA detection antibody.

- Antigen References:**
1. Fitzgerald K, *et al.* Eds. 2001. *The Cytokine FactsBook.* Academic Press San Diego.
  2. Baggiolini M, *et al.* 1994. *Adv. Immunol.* 55:97.
  3. Schröder J, *et al.* 1992. *Immunology Ser.* 57:387.
  4. Zwahlen R, *et al.* 1993. *Intl. Rev. Expt. Pathol.* 34:27.