

**PE/Cy7 anti-mouse CX3CR1**

**Catalog # / Size:** 1345080 / 100 µg  
1345075 / 25 µg

**Clone:** SA011F11

**Isotype:** Mouse IgG2a, κ

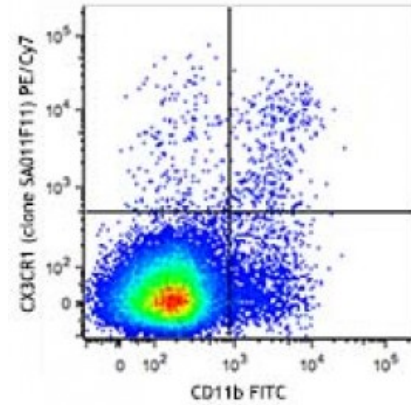
**Immunogen:** Mouse CX3CR1-transfected cells

**Reactivity:** Mouse

**Preparation:** The antibody was purified by affinity chromatography and conjugated with PE/Cy7 under optimal conditions. The solution is free of unconjugated PE/Cy7 and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

**Concentration:** 0.2



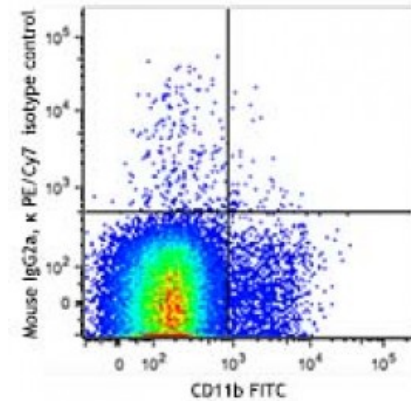
C57BL/6 mouse splenocytes were stained with CD11b FITC and CX3CR1 (clone SA011F11) PE/Cy7 (top) or mouse IgG2a, κ PE/Cy7 isotype control (bottom).

**Applications:**

**Applications:** Flow Cytometry

**Recommended Usage:** Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is ≤0.015 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

**Application Notes:** For *in vivo* studies or highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 149011) with a lower endotoxin limit than standard LEAF™ purified antibodies (Endotoxin <0.01 EU/microg).



**Description:** CX3CR1 is a 40 kD, G-protein coupled receptor, with seven transmembrane regions. CX3CR1 is expressed by resident and alternatively activated macrophages (M2), a subset of monocytes, dendritic cells (DCs), NK cells, a subset of memory T cells, and mast cells. CX3CR1 is involved in cell recruitment during inflammation and participates in cell adhesion and extravasation from blood vessels. Its ligand is CX3CL1, also known as fractalkine or neurotactin. CX3CR1 is also a coreceptor for HIV1 and variations in this gene leads to increased susceptibility to HIV. In the brain, it is expressed by glial cells, which interact with CX3CL1 expressed by neurons.

- Antigen References:**
1. Ponzetta A, *et al.* 2013. *J. Immunol.* 191:5684.
  2. Jacquelin S, *et al.* 2013. *Blood.* 122:674.
  3. Garcia JA, *et al.* 2013. *J. Immunol.* 191:1063.
  4. Lee YS, *et al.* 2013. *Ce*

