

## DESCRIPTION

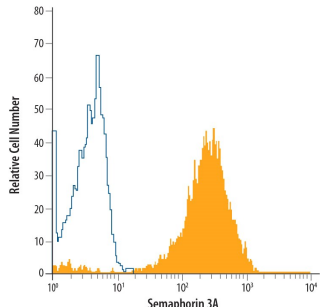
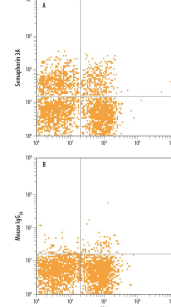
<b>Species Reactivity</b>	Human/Mouse
<b>Specificity</b>	Detects human Semaphorin 3A in direct ELISAs and Western blots. In direct ELISAs and Western blots, no cross-reactivity with recombinant human Semaphorin 3B, 3E, 6A, 6B, 6C, 6D, 7A, recombinant mouse Semaphorin 3C, or 3F is observed.
<b>Source</b>	Monoclonal Mouse IgG <sub>2B</sub> Clone # 215803
<b>Purification</b>	Protein A or G purified from hybridoma culture supernatant
<b>Immunogen</b>	Mouse myeloma cell line NS0-derived recombinant human Semaphorin 3A Lys26-Val771 (Arg555Ala, Arg552Ala) Accession # Q14563
<b>Conjugate</b>	Phycoerythrin Excitation Wavelength: 488 nm Emission Wavelength: 565-605 nm
<b>Formulation</b>	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details.  *Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.

## APPLICATIONS

**Please Note:** Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
<b>Intracellular Staining by Flow Cytometry</b>	10 µL/10 <sup>6</sup> cells	See Below

## DATA

<p><b>Intracellular Staining by Flow Cytometry</b></p>  <p><b>Detection of Semaphorin 3A in CD3<sup>+</sup>-gated Human PBMCs by Flow Cytometry.</b> CD3<sup>+</sup>-gated human peripheral blood mononuclear cells (PBMCs) treated with PHA were stained with Mouse Anti-Human/Mouse Semaphorin 3A PE-conjugated Monoclonal Antibody (Catalog # IC1250P, filled histogram) or isotype control antibody (Catalog # IC0041P, open histogram). To facilitate intracellular staining, cells were fixed with Flow Cytometry Fixation Buffer (Catalog # FC004) and permeabilized with Flow Cytometry Permeabilization/Wash Buffer I (Catalog # FC005). View our protocol for <a href="#">Staining Intracellular Molecules</a>.</p>	<p><b>Intracellular Staining by Flow Cytometry</b></p>  <p><b>Detection of Semaphorin 3A in Mouse Splenocytes by Flow Cytometry.</b> Mouse splenocytes treated with PHA were stained with Rat Anti-Mouse CD3 APC-conjugated Monoclonal Antibody (Catalog # FAB4841A) and either (A) Mouse Anti-Human/Mouse Semaphorin 3A PE-conjugated Monoclonal Antibody (Catalog # IC1250P) or (B) Mouse IgG<sub>2B</sub> Phycoerythrin Isotype Control (Catalog # IC0041P). To facilitate intracellular staining, cells were fixed with Flow Cytometry Fixation Buffer (Catalog # FC004) and permeabilized with Flow Cytometry Permeabilization/Wash Buffer I (Catalog # FC005). View our protocol for <a href="#">Staining Intracellular Molecules</a>.</p>
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## PREPARATION AND STORAGE

<b>Shipping</b>	The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.
<b>Stability &amp; Storage</b>	<b>Protect from light. Do not freeze.</b> <ul style="list-style-type: none"> <li>12 months from date of receipt, 2 to 8 °C as supplied.</li> </ul>

## BACKGROUND

The Semaphorins constitute a large family of secreted, GPI-anchored and transmembrane cell signaling molecules. Depending on their domain organization and species origin, these proteins can be classified into eight groups. To date, at least 19 vertebrate Semaphorins belonging to five groups (class 3 through 7) have been identified. All Semaphorins contain a conserved, 500 amino acid (aa) Sema domain at the amino terminus. Semaphorins are best known for their roles in axon guidance during neuronal development. Semaphorins are also expressed in non-neuronal tissues and are involved in angiogenesis, hematopoiesis, organogenesis, and the regulation of immune functions (1, 2).

Class 3 Semaphorins (Sema3) are secreted proteins containing a Sema domain, an immunoglobulin c2-like domain and a basic domain near the carboxyl tail. Sema3A (also referred to as SemIII, SemD and Collapsin) cDNA predicts a 771 aa precursor protein with a putative 25 aa signal peptide (1-3). Bioactive Sema3A is a disulfide-linked dimer (4). The bioactivity is increased after proteolytic processing by a furin-like endoprotease near the carboxy-terminus (1). The functional receptor complex for Sema3 is composed of two distinct transmembrane proteins: Neuropilin-1 (Npn-1) and Plexin-A. Npn-1 binds directly to Sema3A with high-affinity and confers specificity. Plexin-A interacts with Npn-1 to increase the affinity of the complex for Sema3A and serves as the signaling subunit in the receptor complex (1, 2, 5).

## References:

1. Nakamura, F. *et al.* (2000) *J Neurobiol.* **44**:219.
2. Goshima, Y. *et al.* (2002) *J. Clin. Invest.* **109**:993.
3. Kolodkin, A.L. *et al.* (1993) *Cell* **75**:1389.
4. Koppel, A.M. *et al.* (1998) *J. Biol. Chem.* **273**:15708.
5. Yu, T.W. *et al.* (2001) *Nature Neurosci. Supplement* **4**:1169.
6. Luo Y. *et al.* (1993) *Cell* **75**:217.