

DESCRIPTION

Species Reactivity	Mouse
Specificity	Detects mouse EphA1 in direct ELISAs. In direct ELISAs, no cross-reactivity with recombinant mouse EphA2, recombinant human (rh) EphA3, or rhEphA5 is observed.
Source	Monoclonal Rat IgG _{2A} Clone # 404517
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Mouse myeloma cell line NS0-derived recombinant mouse EphA1 Glu27-Glu548 Accession # AAH71215
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

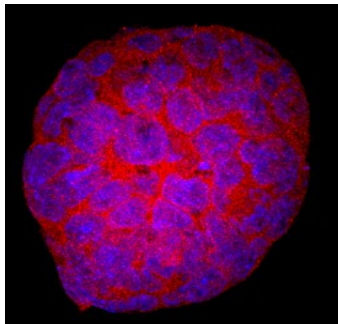
APPLICATIONS

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
Immunocytochemistry	8-25 µg/mL	See Below

DATA

Immunocytochemistry



EphA1 in D3 Mouse Cell Line. EphA1 was detected in immersion fixed D3 mouse embryonic stem cell line using Rat Anti-Mouse EphA1 Monoclonal Antibody (Catalog # MAB30341) at 10 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Rat IgG Secondary Antibody (red; Catalog # NL013) and counterstained with DAPI (blue). Specific staining was localized to cytoplasm and cell secretion. View our protocol for [Fluorescent ICC Staining of Stem Cells on Coverslips](#).

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> ● 12 months from date of receipt, -20 to -70 °C as supplied. ● 1 month, 2 to 8 °C under sterile conditions after reconstitution. ● 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

EphA1, also known as Eph and Esk, is a member of the Eph receptor tyrosine kinase family and binds several Ephrin-A ligands. The A and B class Eph proteins share a common structural organization (1-4). The mouse EphA1 cDNA encodes a 977 amino acid (aa) precursor that includes a 24 aa signal sequence, a 524 aa extracellular domain (ECD), a 21 aa transmembrane segment, and a 408 aa cytoplasmic domain. The ECD contains an N-terminal globular domain, a cysteine-rich domain, and two fibronectin type III domains. The cytoplasmic domain contains a juxtamembrane motif with two tyrosine residues which are the major autophosphorylation sites, a kinase domain, and a conserved sterile alpha motif (SAM) (5, 6). Within the ECD, mouse EphA1 shares 84% aa sequence identity with human EphA1, approximately 40% aa sequence identity with mouse EphA2, 3, 4, 6, 7, and 8, and 27% aa sequence identity with mouse EphA5. A splice variant of mouse EphA1 lacks the transmembrane segment and is predicted to exist as a soluble molecule (7). Membrane bound or clustered Ephrin ligands interact with EphA1 and activate its kinase domain which is capable of Ser, Thr, and Tyr phosphorylation (7). Reverse signaling is propagated through the Ephrin ligand. EphA1 is widely expressed in differentiated epithelial cells, particularly in bone marrow, spleen, thymus, and testes (5, 7). It is expressed on CD4⁺ T cells but not on CD19⁺ B cells (8). On T cells, EphA1 induces Tyr phosphorylation of PYK2 and promotes chemokine-induced chemotaxis (8). EphA1 is upregulated or downregulated in a variety of human carcinomas and is implicated in tumor invasiveness (2, 9, 10).

References:

1. Poliakov, A. *et al.* (2004) *Dev. Cell* **7**:465.
2. Surawska, H. *et al.* (2004) *Cytokine Growth Factor Rev.* **15**:419.
3. Pasquale, E.B. (2005) *Nat. Rev. Mol. Cell Biol.* **6**:462.
4. Davy, A. and P. Soriano (2005) *Dev. Dyn.* **232**:1.
5. Coulthard, M.G. *et al.* (2001) *Growth Factors* **18**:303.
6. Lickliter, J.D. *et al.* (1996) *Proc. Natl. Acad. Sci. USA* **93**:145.
7. Douville, E.M.J. *et al.* (1992) *Mol. Cell. Biol.* **12**:2681.
8. Aasheim, H-C. *et al.* (2005) *Blood* **105**:2869.
9. Iwase, T. *et al.* (1993) *Biochim. Biophys. Res. Commun.* **194**:698.
10. Hirai, H. *et al.* (1987) *Science* **238**:1717.