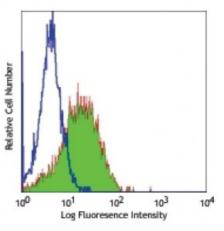
## **Product Data Sheet**

## Alexa Fluor<sup>®</sup> 488 anti-mouse CD309 (VEGFR2, Flk-1)

Catalog # / Size:	1209540 / 100 μg 1209535 / 25 μg
Clone:	89B3A5
Isotype:	Rat IgG2a, к
Immunogen:	Rat-1 cells transfected with full-length mouse Flk
<b>Reactivity:</b>	Mouse
Preparation:	The antibody was purified by affinity chromatography, and conjugated with Alexa Fluor® 488 under optimal conditions.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Concentration:</b>	0.5



Mouse FLK-1 transfected cells stained with 89B3A5 Alexa Fluor® 488

## **Applications:**

Applications:Flow CytometryRecommended<br/>Usage:Each lot of this antibody is quality control tested by immunofluorescent staining<br/>with flow cytometric analysis. For flow cytometric staining, the suggested use of<br/>this reagent is ≤2.0 microg per million cells in 100 microL volume. It is<br/>recommended that the reagent be titrated for optimal performance for each<br/>application.

 $\ast$  Alexa Fluor  $\circledast$  488 has a maximum emission of 519 nm when it is excited at 488 nm.

Application	1. Kaburn N, <i>et al.</i> 1997. <i>Development</i> 124:2039.
<b>References:</b>	

**Description:** The 89B3A5 antibody recognizes mouse CD309 also known as vascular endothelial growth factor receptor 2, VEGFR2, KDR, protein tyrosine kinase receptor flk-1, and fetal liver kinase-1. Flk-1 is a member of the tyrosine protein kinase family, sub-family CSF-1/PDGF, that contains a single pass transmembrane receptor with a protein kinase domain and seven immunoglobulin-like domains in the extracellular region. Flk-1 is expressed at high levels in adult heart, lung, kidney, brain, and skeletal muscle; other tissues express at lower levels. Flk-1 is a receptor for VEGF or VEGFC; ligand binding plays a key role in vascular development and vascular permeability. The 89B3A5 antibody has been shown to be useful for flow cytometry.

Antigen	1. Patterson C, <i>et al.</i> 1995. <i>J. Biol. Chem.</i> 270:23111.
<b>References:</b>	2. Quinn TP, et al. 1993. Proc. Natl. Acad. Sci. USA 90:7533.

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