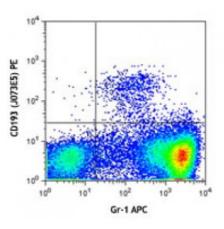
## **Product Data Sheet**

## PE anti-mouse CD193 (CCR3)

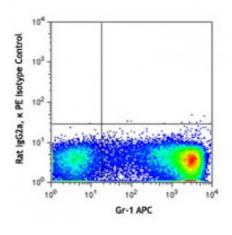
Catalog # / Size:	1322530 / 100 µg 1322525 / 25 µg
Clone:	J073E5
Isotype:	Rat IgG2a, к
Immunogen:	Mouse CCR3-transfectants
<b>Reactivity:</b>	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Concentration:</b>	0.2



C57BL/6 mouse peripheral blood myeloid cells were stained with Gr-1 APC and CD193 (clone J073E5) PE (top) or rat IgG2a, κ PE 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 $\leq 0.25$ microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.



Description:	CD193, also known as CC-chemokine receptor 3 (CCR3), CC CKR3, MIP1- $\alpha$ receptor like-2, and eotaxin receptor, is a member of the G protein-coupled, seven transmembrane receptor family. It binds to the CC chemokines eotaxin, eotaxin-2, and eotaxin-3 with high affinity. CD193 has also been reported to bind RANTES, MCP-3, and MCP-4 with low affinity. CD193 is expressed on mouse eosinophils, basophils, mast cells, mononuclear phagocytes, platelets, hematopoietic progenitor cells, and keratinocytes. It is thought to play a role in allergic diseases such as bronchial asthma and allergic rhinitis. CD193 also function as a co-receptor for HIV-1 and HIV-2, and the binding of eotaxin with CD193 has been shown to inhibit HIV infection in some cell types.
Antigen References:	1. Zlotnik A, <i>et al.</i> 2006. <i>Genome Biol.</i> 7:243. 2. Kodali RB, <i>et al.</i> 2004. <i>Arterioscler. Thromb. Vasc. Biol.</i> 24:1211. 3. Das AM, <i>et al.</i> 2006. <i>J. Pharmacol. Exp. Ther.</i> 318:411.

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