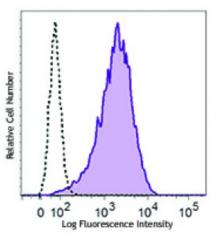
Product Data Sheet

Biotin anti-mouse IL-33Rα (IL1RL1, ST2)

Catalog # / Size:	1326535 / 25 μg 1326540 / 100 μg
Clone:	DIH9
Isotype:	Rat IgG2a, к
Immunogen:	IL-33R α -hFc γ 1 fusion protein.
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with biotin under optimal conditions. The solution is free of unconjugated biotin.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
Concentration:	0.2



Mouse Th2 clone D10.G4.1 was stained with biotinylated anti-mouse IL-33R α /ST2 (clone DIH9) (filled histogram) or biotinylated rat IgG2a, κ isotype control (open histogram), followed by SAV-PE.

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 ≤ 0.5 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.
Application References:	1. Hashiguchi M, <i>et al.</i> 2014. <i>Eur. J. Immunolog</i> y. (FC) <u>PubMed</u>
Description:	IL-33R α , also known as ST2 or IL-1RL1, is a member of the Toll/IL-1 receptor family. It binds IL-33 and is structurally similar to IL-1R1. Two forms of the protein exist - a soluble form known as ST2 and a membrane anchored form known as ST2L. The membrane form is expressed by Th2 cells and bone marrow derived mast cells, whereas the soluble form is expressed by serum-stimulated fibroblasts.
	Blocking with anti-ST2 antibodies has been shown to alleviate experimental arthritis and airway inflammation. The IL-33-ST2 axis has been reported to be important across a range of diseases including asthma, allergies, and cardiac disease.
Antigen References:	 Yanagisawa K, <i>et al.</i> 1993. <i>FEBS Lett.</i> 318:83. Schmitt E, <i>et al.</i> 1990. <i>Cytokine</i> 6:407. Yanagisawa K, <i>et al.</i> 1992. <i>FEBS Lett.</i> 302:51. Takagi T, <i>et al.</i> 1993.

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