## **Product Data Sheet**

## Brilliant Violet 421™ anti-mouse CD192 (CCR2)

Catalog # / Size: 1353025 / 50 μg

Clone: SA203G11
Isotype: Rat IgG2b, κ

Immunogen: Mouse CCR2 transfectants.

Reactivity: Mouse

**Preparation:** The antibody was purified by affinity

chromatography and conjugated with Brilliant Violet  $421^{™}$  under optimal conditions. The solution is free of unconjugated Brilliant Violet  $421^{™}$  and

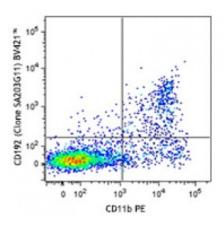
unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and BSA

(origin USA).

Concentration: Lot-specific



C57BL/6 bone marrow was stained with Ly-6G APC/Cy7, CD11b PE, and CD192 (clone SA203G11) Brilliant Violet  $421^{\text{TM}}$  (top) or rat IgG2b,  $\kappa$  Brilliant Violet  $421^{\text{TM}}$  isotype control (bottom). Dot plots are gated on Ly- $6G^{-}$  cells.

## **Applications:**

**Applications:** Flow Cytometry

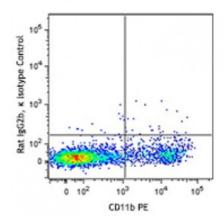
Recommended

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.

Brilliant Violet 421™ excites at 405 nm and emits at 421 nm. The standard bandpass filter 450/50 nm is recommended for detection. Brilliant Violet 421™ is a trademark of Sirigen Group Ltd.

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**Description:** CD192, also known as CCR2, is a 42 kD G-protein coupled receptor that is

associated with bone marrow homeostasis. Specifically, CD192 mediates monocyte chemotaxis and acts as a receptor for monocyte chemoattractant protein 1 (MCP-1). CD192 is primarily expressed on monocytes and macrophages, with some expression on basophils. It is involved in monocyte infiltration in

inflammatory diseases such as rheumatoid arthritis and cancer.

Antigen References:

1. Mack M, et al. 2001. J. Immunol. 166:4697.

2. Dutta P, et al. 2015. Cell Stem Cell. 16:477.

3. Li L, et al. 2008. Kidney Int. 74:1526.