KIRAVIA Blue 520™ anti-mouse CD163

Catalog # / $1376590 / 100 \mu g$

Size: 1376585 / 25 μg

Clone: S15049I

Isotype: Rat IgG2a, κ

Immunogen: Recombinant mouse CD163

extracellular domain

Reactivity: Mouse

Preparation: The antibody was purified by affinity

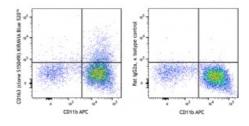
chromatography and conjugated with KIRAVIA Blue 520™ under optimal

conditions.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide

Concentration: 0.2 mg/mL



C57BL/6 mouse bone marrow cells were stained with CD11b APC and anti-mouse CD163 (clone S15049I) KIRAVIA Blue 520™ (left) or rat IgG2a, κ KIRAVIA Blue 520™ isotype control (right). Data shown were from myeloid population

population.

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 $\leq 0.5~\mu g$ per million cells in $100~\mu L$ volume. It is recommended that the reagent be titrated for optimal performance for each application.

* KIRAVIA Blue 520™ has an excitation maximum of 495 nm, and a maximum emission of 520 nm.

Application Notes:

Additional reported applications (for the relevant formats) include: immunoprecipitation 1 , in vitro costimulation of T and NK cells 1 , in vitro blocking of allogeneic mixed leukocyte response and inhibition of MHC-

unrestricted CTL cytotoxicity^{3,4}, in vitro induction of thymocyte

 $\ differentiation^{2,5-9,11},\ and\ immunohistochemical\ staining\ of\ acetone-fixed$

frozen sections. For *in vivo* studies or highly sensitive assays, we

recommend Ultra-LEAF $^{\text{\tiny TM}}$ purified antibody (Endotoxin < 0.01 EU/ μ g, Azide-

Free, 0.2 µm filtered) (Cat. No. 102116).

Application References:

- 1. Gross JA, et al. 1992. J. Immunol. 149:380. (IP, Costim)
 - 2. Cibotti R, et al. 1997. Immunity 6:245. (Costim)
 - 3. Masten BJ, et al. 1997. Am. J. Respir. Cell Mol. Biol. 16:335. (Block)
 - 4. Nishio M, et al. 1996. J. Immunol. 157:4347. (Block)
 - 5. Zhang N and He Y-W, 2005. J. Exp. Med. 202:395. (Costim)
 - 6. Terrazas LI, et al. 2005. Intl. J. Parasitology. 35:1349. (Costim)
 - 7. Perchonock CE, et al. 2006. Mol Cell Biol. 26(16):6005. (Costim)
 - 8. Wang W, et al. 2007. J. Immunol. 178:4885. (Costim)
 - 9. Pua HH, et al. 2007. J. Exp. Med. 204:25. (Costim)
- 10. Perchonock CE, et al. 2007. J. Immunol. 179:1768.
- 11. Barbi J, et al. 2007. Blood 110:2215.
- 12. Milpied P, et al. 2011. Blood 118:2993. PubMed
- 13. Cunningham NR, et al. 2011. Int Immunol. 23:693. PubMed
- 14. Crispin JC, et al. 2012. J. Immunol. 188:3567. PubMed
- 15. Li CR, et al. 2014. J Immunol. 192:1425. PubMed
- 16. Blankenhaus B, et al. 2014. PLoS Pathog. 10:1003913. PubMed

Description:

CD163 is a member of the group B scavenger receptor cysteine-rich superfamily, also known as GHI/61, M130, RM3/1, p155, hemoglobin-haptoglobin complex receptor, or macrophage-associated antigen. It is a 134 kD (non-reduced)/155 kD (reduced) glycoprotein primarily expressed on macrophages, Kupffer cells, monocytes, a subset of dendritic cells, and a subset of hematopoietic stem/progenitor cells. CD163 binds to haptoglobin-hemoglobin complex and TWEAK, and plays a role in clearing hemoglobin and regulating cytokine production by macrophages. Membrane CD163 can be cleaved by metalloproteinases (MMP), resulting in a soluble form. Elevated serum level of sCD163 has been implicated in many kinds of inflammatory diseases.

Antigen References:

- 1. Kim CH, et al. 2001. J. Clin. Invest. 107:595.
- 2. Heesch K, et al. 2014. PLoS One. 9:5.
- 3. Wehr A, et al. 2013. J. Immunol. 190:5226.