Brilliant Violet 421™ anti-human CD303 (BDCA-2)

Catalog # / Size: 2371060 / 100 tests

2371055 / 25 tests

Clone: 201A

Isotype: Mouse IgG2a, κ

Reactivity: Human

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

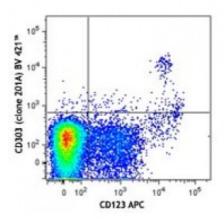
unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and BSA

(origin USA).

Concentration: Lot-specific



Human peripheral mononuclear cells were stained with CD123 APC and CD303 (clone 201A) Brilliant Violet 421™ (top) or mouse IgG2a, κ Brilliant Violet 421™ isotype control (bottom). Data shown was gated on the lymphocyte population.

Applications:

Applications: Flow Cytometry

Recommended

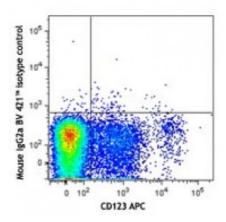
Usage:

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 ≤5 microL per million cells or 5 microL per 100 microL of whole blood. 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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U.S. Patent(s), pending patent applications and foreign equivalents.

Description: CD303, also known as BDCA-2 and CLEC4C, is a 38 kD type II transmembrane

glycoprotein. It is a member of the C-type lectin superfamily. CD303 is expressed by plasmacytoid dendritic cells (pDCs) and is involved in cell adhesion, signaling, and antigen capture and processing. Crosslinking of CD303 inhibits the production

of IFN- α/β and TLR-9 induced pDCs maturation.

Antigen References:

1. Jähn PS, et al. 2010. Cell Immunol. 265:15.

2. Graham LM and Brown GD. 2009. Cytokine 48:148.

3. Röck J, et al. 2007. Eur. J. Immunol. 37:3564.

4. Dzionek A, et al. 20