Product Data Sheet

Brilliant Violet 421™ Rat IgG1, κ Isotype Ctrl

Catalog # / Size: 2602145 / 125 μl

2602150 / 500 µl

2602195 / 50 μg

Clone: RTK2071 Isotype: Rat IgG1, κ

Immunogen: Trinitrophenol + KLH

Preparation: The immunoglobulin was purified by

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

unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and BSA

(origin USA).

Concentration: microg sizes: 0.2 mg/ml

microL sizes: lot-specific

Applications:

Applications: Flow Cytometry

Recommended

Usage:

Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis as negative control. Use at concentrations comparable to those of the specific antibody of interest. Use our Concentration Lookup tool to find the exact concentrations of your lots of product.

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

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Application Notes:

The RTK2071 immunoglobulin is useful as an isotype-matched control (for the relevant formats) for Western blotting, immunoprecipitation,

immunohistochemistry, functional assay, and immunofluorescence microscopy. The LEAF $^{\text{\tiny IM}}$ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 400414) as negative control1. For in vivo studies or highly sensitive assays, we recommend Ultra-LEAF $^{\text{\tiny IM}}$ purified antibody (Cat. No. 400432) with a lower endotoxin limit than standard LEAF $^{\text{\tiny IM}}$

purified antibodies (Endotoxin < 0.01 EU/microg).

Application 1. Riemann M, et al. 2005. J. Immunol. 175:3560. PubMed 2. Wondimu Z, et al. 2010. Am. J. Pathol. 177:2334. PubMed

Description: The isotype of RTK2071 immunoglobulin is rat IgG1, κ. This antibody was chosen

as an isotype control after screening on a variety of resting, activated, live, and fixed mouse, rat and human tissues.