## Brilliant Violet 421™ Armenian Hamster IgG Isotype Ctrl

**Catalog # / Size:** 2604680 / 500 μl

2604675 / 125 µl

2604745 / 50 µg

Clone: HTK888

**Isotype:** Hamster IgG

Immunogen: Trinitrophenol + KLH

**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: 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.

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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Application Notes:

The HTK888 immunoglobulin is useful as an isotype-matched control (for the relevant formats) for Western blotting, immunoprecipitation, functional assay, immunofluorescence microscopy, immunocytochemistry and immunofluorescent staining (surface or intracellular) for flow cytometric analysis. The LEAF  $^{\text{TM}}$  purified antibody (Endotoxin <0.1 EU/ $\mu$ g, Azide-Free, 0.2  $\mu$ m filtered) is recommended for functional assays (Cat. No. 400916) as negative control. For *in vivo* studies or highly sensitive assays, we recommend Ultra-LEAF  $^{\text{TM}}$  purified antibody (Cat. No. 400940) with a lower endotoxin limit than standard LEAF  $^{\text{TM}}$  purified antibodies (Endotoxin <0.01 EU/microg).

Application References:

1. Lesley R, et al. 2006. P. Natl. Acad. Sci. USA 103:10717.

2. Yu R, et al. 2006. Obesity 14:1353.

3. Yang JH, et al. 2005. Rheumatology (Oxford). 44:1245. PubMed

3. Mina-Osorio P, et al. 2008. J. Leukocyte Biol. 84:448. PubMed

4. Shen H, et al. 2009. J. Am Soc Nephrol. 20:1032. PubMed

Description:	This antibody was chosen as an isotype control after screening on a variety of resting, activated, live, and fixed mouse, rat and human tissues.