PE anti-human MERTK

Catalog # / Size: 2438040 / 100 tests

2438035 / 25 tests

Clone: 590H11G1E3

Isotype: Mouse IgG1, κ

Immunogen: MERTK extracellular domain/Fc fusion.

Reactivity: Human

Preparation: The antibody was purified by affinity

chromatography and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and

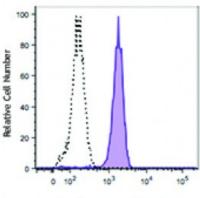
unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

0.2% (w/v) BSA (origin USA).

Concentration: Lot-specific



Log Fluorescence Intensity

Human peripheral blood monocytes were stimulated and cultured with M-CSF for seven days and stained with human MERTK (clone

590H11G1E3) PE (filled histogram) or mouse IgG1, κ isotype control

(open histogram).

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.

Application References:

1. Rogers AE, et al. 2012. Oncogene 31:4171.

Description: MERTK plays a role in the retinal pigment epithelium as a regulator of rod outer

segments fragments phagocytosis. MERTK also plays a role in the inhibition of Toll-like receptor-mediated innate immune responses through the activation of STAT1. Upregulation of MERTK seems to also promote the survival of certain cancer cells, such as t(1;19)-positive acute lymphoblastic leukemias (ALL). MERTK also has a role in cellular migration, as MERTK KO macrophages demonstrate cytoskeletal disruptions that impacts its shape and directional migration. Melanoma cells express high levels of MERTK, which makes this molecule an

attractive therapeutic target.

Antigen

1. Schlegel J, et al. 2013. J. Clin. Invest. 123:2257.

References: 2. Chen J, et al. 1997. Oncogene 14:2033.

3. Yefimova MG, et al. 2013. Autophagy 9:653.

4. Zhang W, et al. 2013. J.