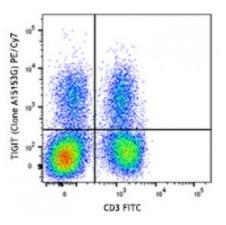
## **Product Data Sheet**

## PE/Cy7 anti-human TIGIT (VSTM3)

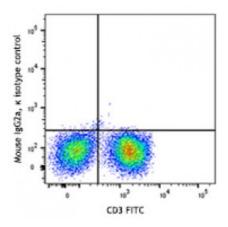
Catalog # / Size:	2463565 / 25 tests 2463570 / 100 tests
Clone:	A15153G
Isotype:	Mouse IgG2a, к
Immunogen:	Recombinant Human TIGIT.
<b>Reactivity:</b>	Human
Preparation:	The antibody was purified by affinity chromatography and conjugated with PE/Cy7 under optimal conditions. The solution is free of unconjugated PE/Cy7 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



Human peripheral blood leukocytes were stained with FITC CD3 and TIGIT (clone A15153G) PE/Cy7 (top) or PE/Cy7 mouse IgG2a, κ PE/Cy7 isotype control (bottom). Data shown was gated on the lymphocytes 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.
Application Notes:	This clone can suppress anti-CD3 induced T cell proliferation <i>in vitro</i> .



**Description:** T cell immunoreceptor with Ig and ITIM domains (TIGIT), also known as VSTM3 or WUCAM, is a 26 kD, type I transmembrane protein and is a member of the PVR (poliovirus receptor) family of immunoglobulin-like domain containing proteins. TIGIT is expressed on activated T cells, follicular T helper, memory, and regulatory T cells as well as on NK cells. TIGIT is a negative regulator of NK and T cell activation. Expression of TIGIT is associated with decreased functionality of CD8 T cells in chronic viral infection and tumors. TIGIT also promotes the differentiation of tolerogenic phenotype in dendritic cells with an increased secretion of IL-10 and a diminished production of IL-12.

 Antigen
 1. Stanietsky N, et al. 2009. Proc. Natl. Acad. Sci. 106:17858.

 References:
 2. Yu X, et al. 2009. Nat. Immunol. 10:48.

 3. Johnston R, et al. 2014. Cancer Cell. 26:923.

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