## Purified anti-human CD58 (LFA-3)

Catalog # / Size: 2254510 / 100 μg

Clone: TS2/9

**Isotype:** Mouse IgG1, κ

Immunogen: Human cytolytic T cells

Reactivity: Human

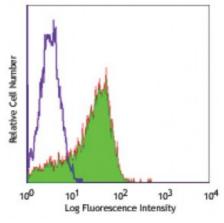
**Preparation:** The antibody was purified by affinity

chromatography.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

**Concentration:** 0.5



Human peripheral blood lymphocytes stained with purified TS2/9, followed by anti-mouse IgG FITC

## **Applications:**

**Applications:** Other

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  $\leq 1.0$  microg per  $10^6$  cells in 100 microL volume or 100 microL of

whole blood. It is recommended that the reagent be titrated for optimal

performance for each application.

Application Notes:

Additional reported applications include: immunoprecipitation1, inhibition of

cytolytic activity1, augment of IL-1 release by TE cells2

Application

1. Sanchez-Madrid F, et al. 1982. Proc. Nati Acad. Sci. USA. 79:7489

**References:** 2. Le PT, *et al.* 1990. *J. Immunol.* 144:4541

**Description:** CD58, also known as lymphocyte function-associated antigen 3 (LFA-3) is a 45-70

kD cell surface protein that is a member of the immunoglobulin superfamily.

Alternative splicing of CD58 gives rise to transmembrane and

glycosylphosphatidylinositol (GPI)-anchored forms on cell surface. CD58 is

expressed on both hematopoietic and non-hematopoietic cells including B cells, T cells, monocytes, erythrocytes, endothelial cells, epithelial cells, and fibroblasts. High levels are observed on memory T cells and dendritic cells. CD58 expressed on antigen presenting cells and target cells enhances T cell recognition via the binding of it's cognate ligand, CD2, on the T cell surface. The HCD58 antibody recognizes human CD58 and has been shown to be useful for flow cytometry.

Antigen References:

1. Springer TA, et al. 1987. Annu. Rev. Immunol. 5:223.

2. Dustin ML, et al. 1987. Nature 329:846.

3. Arulanandam AR, et al. 1994. J. Exp. Med. 180:1861.

4. Sanders ME, et al.