Product Data Sheet

APC anti-mouse CD253 (TRAIL)

Catalog # / Size: 1146545 / 25 μg

1146550 / 100 µg

Clone: N2B2

Isotype: Rat IgG2a, κ

Immunogen: Mouse TRAIL-transfected 2PK-3 cells

Reactivity: Mouse

Preparation: The antibody was purified by affinity

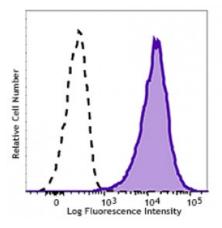
chromatography and conjugated with APC under optimal conditions. The solution is free of unconjugated APC and

unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.2 mg/ml



Mouse TRAIL transfected L5718Y cells were stained with CD253 (clone N2B2) APC (filled histogram) or Rat IgG2a

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 $\leq 1.0 \,\mu g$ per million cells in 100 $\,\mu$ l volume. It is recommended that

the reagent be titrated for optimal performance for each application.

Application Notes:

Additional reported applications (for the relevant formats) include: in vitro

blocking of NK cell cytotoxicity^{1,2}.

Application

1. Kayagaki N, et al. 1999. J. Immunol. 163:1906.

References: 2. Wiley SR, et al. 1995. Immunity 3:673.

3. Wu GS, et al. 1999. Cancer Res. 59:2770.

4. Mariani SM, et al. 1998. Eur. J. Immunol.

Description: CD253 is a 40 kD TNF superfamily member known as TRAIL, Apo-2 ligand, and

Apo-2L. TRAIL is expressed on a variety of cells, including IL-2 and IL-15 activated NK cells and activated T cells. However, it is undetectable on resting T and B cells. TRAIL has been reported to induce apoptosis in tumor and transformed cell lines by a caspase-dependent process. The N2B2 antibody has been reported to be useful for flow cytometric staining and blocking NK cell cytotoxicity *in vitro*.

Antigen References:

1. Kayagaki N, et al. 1999. J. Immunol. 163:1906.

2. Wiley SR, et al. 1995. Immunity 3:673.

3. Wu GS, et al. 1999. Cancer Res. 59:2770.

4. Mariani SM, et al. 1998. Eur. J. Immunol.