

## **TECHNICAL DATA SHEET**

# PE Anti-Mouse CD178 (FasL) (MFL3)

Catalog Number: 50-5911

### PRODUCT INFORMATION

Contents: PE Anti-Mouse CD178 (FasL) (MFL3)

Isotype: Armenian Hamster IgG

Concentration: 0.2 mg/mL

Clone: MFL3

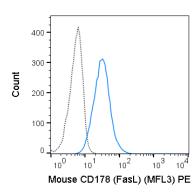
Reactivity: Mouse

Use By: 12 months from date of receipt

Storage Conditions: 2-8°C protected from light

Formulation: 10 mM NaH<sub>2</sub>PO<sub>4</sub>, 150 mM NaCl, 0.09% NaN<sub>3</sub>,

0.1% gelatin, pH7.2



Mouse Fas Ligand transected cells were stained with 0.25 ug PE Anti-Mouse CD178 (50-5911) (solid line) or 0.25 ug PE Armenian Hamster IgG isotype control (dashed line).

Rev. 20190712

#### **DESCRIPTION**

The MFL3 monoclonal antibody reacts with mouse CD178, a 40 kD member type-II transmembrane protein and member of the TNF family of proteins. CD178 is also known as Fas ligand, FasL, Apo-1 ligand, and CD95 ligand. CD178 is expressed on activated T cells and and natural killer (NK) cells and also tissue at immune privileged sites such as the eye and testis. FasL interacts with its receptor CD95 (Fas) to initiate apoptotic cell death and is thought to play a role in T cell development, immune response regulation, and cell-mediated cytotoxic responses. This MFL3 clone has been reported to block CD178/CD96 induced apoptosis.

## **PREPARATION & STORAGE**

This monoclonal antibody was purified from tissue culture supernatant via affinity chromatography. The purified antibody was conjugated under optimal conditions, with unreacted dye removed from the preparation. It is recommended to store the product undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

## **APPLICATION NOTES**

This antibody preparation has been quality-tested for flow cytometry using mouse spleen cells, or an appropriate cell type (where indicated). Please refer to the figure legend for the optimal concentration used to stain the tissue shown. We recommend titrating the antibody under your specific conditions to determine the optimal concentration of antibody needed in your experimental system.

#### **REFERENCES**

Takahashi T, Tanaka M, Brannan CI, et al. 1994. Cell. 76(6):969-976. Suda T, Okazaki T, Naito Y, et al. 1995. J Immunol. 154(8):3806-3813. Vignaux F, Vivier E, Malissen B, Depraetere V, Nagata S, Golstein P. 1995. J Exp Med. 181(2):781-786. Griffith TS, Ferguson TA. 1997. Immunol Today. 18(5):240-244.

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