# Biotin Anti-Mouse NK1.1 (CD161) Monoclonal Antibody

Catalog Number	Vial Size
M100N2-08B	50 µg
M100N2-08E	500 µg



Web | www.sungenebiotech.com

**Important Note:** Centrifuge before opening to ensure complete recovery of vial contents. This product is guaranteed up to one year from purchase.

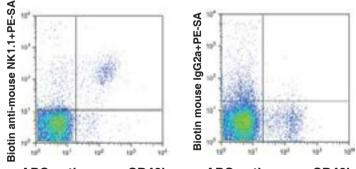
#### **Purified Antibody Characterization**

Clone	Isotype	Reactivity
PK136	Mouse IgG2a	Mouse

#### Description

NK-1.1 surface antigen is encoded by the NKR-P1B/NKR-P1C gene, also known as CD161b/CD161c and Ly-55. It is expressed on NK cells and NK-T cells in some mouse strains, including C57BL/6, FVB/N, and NZB, but not AKR, BALB/c, CBA/J, C3H, DBA/1, DBA/2, NOD, SJL, and 129. Expression of NKR-P1C antigen has been correlated with lysis of tumor cells in vitro and rejection of bone marrow allografts in vivo. NK-1.1 has also been shown to play a role in NK cell activation, IFN- $\gamma$  production, and cytotoxic granule release. NK-1.1 and DX5 are commonly used as mouse NK cell markers.

## Illustration of Immunofluorescent Staining



APC anti-mouse CD49b

APC anti-mouse CD49b

C57BL/6 mouse splenocytes stained with APC anti-mouse CD49b and Biotin anti-mouse NK1.1(left) or mouse IgG2a isotype control(right), followed by PE-SA

## **Product Information**

Conjugation: Biotin

**Formulation:** PBS pH 7.2, 0.09% NaN<sub>3</sub>, 0.2% BSA

Concentration: 0.5 mg/ml

**Storage:** Keep as concentrated solution. Store at 4°C and protected from prolonged exposure to light. **Do not freeze.** 

Application: Recommended Application: FC

**Usage:** Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis (The amount of the reagent is suggested to be used  $\leq 0.25$  µg /10<sup>6</sup> cells in 100 µl). Since applications vary, the appropriate dilutions must be determined for individual use.

## References

- [1] Carlyle, J.R., et al. 1999. J. Immunol. 162:5917.
- [2] Sentman, C.L., etal. 1989. Hybridoma 8:605.
- [3] Koo, G.C., et al. 1984. Hybridoma 3:301.
- [4] Sentman, C.L., et al. 1989. J. Immunol. 142:1847.
- [5] Koo, G.C., et al. 1986. J. Immunol. 137:3742.
- [6] Karlhofer, F.M., et al. 1991. J. Immunol. 146:3662.

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