## PE anti-mouse CD66a (CEACAM1a)

**Catalog # / Size:**  $1272525 / 25 \mu g$ 

1272530 / 100 μg

Clone: MAb-CC1

**Isotype:** Mouse IgG1, κ

Immunogen: Balb/c mouse purified intestinal brush

border membrane

Reactivity: Mouse

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

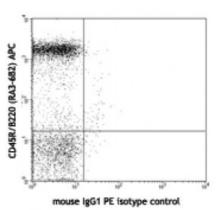
chromatography, and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and

unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.2



Balb/c mouse splenocytes stained with MAb-CC1 PE (lower panel) or mouse IgG1 PE isotype control (upper panel) and CD45R/B220 (RA3-6B2) APC

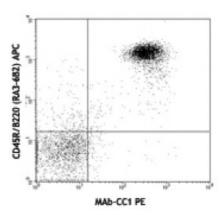
## **Applications:**

**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 ≤0.25 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.



Application References:

- 1. Turner BC, et al. 2004. J. Virol. 78 (10):5486
- 2. Williams RK, *et al.* 1990. *J. Virol.* 64:3817

3. Dveksler GS, et al. 1993. Proc. Natl. Acad. Sci. USA. 90:1716

**Description:** 

CD66a, known as CEACAM1a, carcinoembryonic antigen-related cell adhesion molecule 1a, is a glycoprotein of the immunoglobulin superfamily and the carcinoembryonic antigen family. Isoforms expressing either two or four alternatively spliced Ig-like domains in mice have been found in a number of epithelial, endothelial, or hematopoietic tissues. CEACAM1a functions as an intercellular adhesion molecule, an angiogenic factor, and a tumor cell growth inhibitor. It also serves as a signal regulatory protein influencing B cell receptor complex-mediated activation. The mouse and human CEACAM1a proteins are targets of viral or bacterial pathogens, respectively. It was reported that targeted disruption of the CEACAM1a gene resulting in a partial ablation of the protein in mice led to reduced susceptibility to virus infection. The antibody recognizes the N-terminal domain of murine CEACAM1a, it does not recognize murine CEACAM1b, an allele in SJL mice.

Antigen 1.

1. Nakagaki K, et al. 2005. J. Virol. 79(10):6102

References:	2. Greicius G <i>et al.</i> 2003. <i>J. Leukoc. Biol.</i> 74(1):126 3. Hemmila E <i>et al.</i> 2004. <i>J. Virol.</i> 78(18):10156
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