

**PE/Dazzle™ 594 anti-mouse H-2Kb bound to SIINFEKL**

**Catalog # / Size:** 1308055 / 25 µg  
1308060 / 100 µg

**Clone:** 25-D1.16

**Isotype:** Mouse IgG1, κ

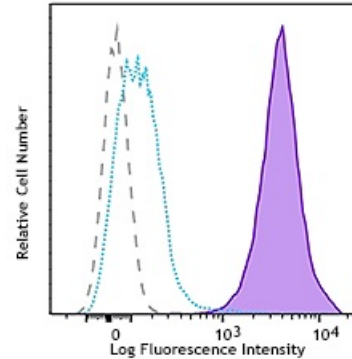
**Immunogen:** SIINFEKL pulsed RMA-S cells

**Reactivity:** Mouse

**Preparation:** The antibody was purified by affinity chromatography and conjugated with PE/Dazzle™ 594 under optimal conditions. The solution is free of unconjugated PE/Dazzle™ 594 and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

**Concentration:** 0.2 mg/ml



C57BL/6 mouse splenocytes were pulsed with or without SIINFEKL for 2 hours, and then stained with anti-mouse SIINFEKL bound H-2K<sup>b</sup> (clone 25-D1.16) PE/Dazzle™ 594 (purple filled histogram indicates the pulsed cells and cyan open histogram

**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 ≤ 0.5 µg per million cells in 100 µl volume. It is recommended that the reagent be titrated for optimal performance for each application.

\* PE/Dazzle™ 594 has a maximum excitation of 566 nm and a maximum emission of 610 nm.

**Application Notes:** The 25-D1.16 monoclonal antibody specifically reacts with ovalbumin-derived peptide SIINFEKL bound to H-2Kb of MHC class I, but not with unbound H-2Kb or H-2Kb bound with an irrelevant peptide. Additional reported applications (for relevant formats) include: Western Blotting<sup>1,3</sup>, immunofluorescence microscopy<sup>2,3</sup>, immunohistochemical staining of frozen tissue sections<sup>3</sup>, and inhibition of T cell response to H-2K<sup>b</sup>-SIINFEKL *in vitro*.

**Application References:**

1. Mareeva T, et al. 2010. *J. Immunol. Methods* 353:78.
2. Mareeva T, et al. 2008. *J. Biol. Chem.* 283:29053.
3. Deng Y, et al. 1998. *J. Immunol.* 161:1677.

**Description:** This antibody has been proven to be very useful in tracking the quantity and localization of these specific antigen-presenting cells (APC) *in vivo*.

**Antigen References:**

1. Mareeva T, et al. 2010. *J. Immunol. Methods* 353:78.
2. Mareeva T, et al. 2008. *J. Biol. Chem.* 283:29053.
3. Deng Y, et al. 1998. *J. Immunol.* 161:1677.