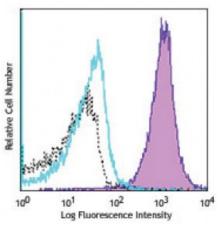
## **Product Data Sheet**

## PE anti-mouse H-2Kb bound to SIINFEKL

Catalog # / Size:	1308015 / 25 μg 1308020 / 100 μg
Clone:	25-D1.16
Isotype:	Mouse IgG1, к
Immunogen:	SIINFEKL pulsed RMA-S cells
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



C57BL/6 mouse splenocytes were pulsed with (filled histogram) or without (cyan open histogram) SIINFEKL for 2 hours and then stained with 25-D1.16 PE or mouse IgG1,  $\kappa$  PE isotype control (broken line 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 $\leq 0.125$ microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.
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 sections3, and inhibition of T cell response to H-2Kb-SIINFEKL <i>in vitro</i> .
Application References:	<ol> <li>Mareeva T, <i>et al.</i> 2010. <i>J. Immunol. Methods</i> 353:78. (WB)</li> <li>Dolan BP, <i>et al.</i> 2010. <i>J. Immunol.</i> 184:1419. (IF)</li> <li>Porgador A, <i>et al.</i> 1997. <i>Immunity</i> 6:715. (WB, IF, IHC)</li> <li>Herve J, <i>et al.</i> 2013. <i>J. Immunol.</i> 190:3163. <u>PubMed.</u></li> </ol>
Description:	This antibody has been proven to be very useful in tracking the quantity and localization of these specific antigen-presenting cells (APC) <i>in vivo</i> .
Antigen References:	<ol> <li>Mareeva T, <i>et al.</i> 2010. <i>J. Immunol. Methods</i> 353:78.</li> <li>Mareeva T, <i>et al.</i> 2008. <i>J. Biol. Chem.</i> 283:29053.</li> <li>Deng Y, <i>et al.</i> 1998. <i>J. Immunol.</i> 161:1677.</li> </ol>

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