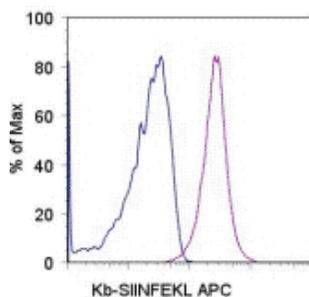


Anti-Mouse OVA257-264 (SIINFEKL) peptide bound to H-2Kb APC

Catalog Number: 17-5743

Also Known As: H-2Kb-SIINFEKL, OVA-Kb

RUO: For Research Use Only



Staining of unpulsed (blue histogram) or SIINFEKL-peptide-pulsed (purple histogram) C57BL/6 splenocytes with 0.06 µg of Anti-Mouse OVA₂₅₇₋₂₆₄ (SIINFEKL) peptide bound to H-2Kb APC. Total viable cells were used for analysis.

Product Information

Contents: Anti-Mouse OVA257-264 (SIINFEKL) peptide bound to H-2Kb APC

REF **Catalog Number:** 17-5743

Clone: eBio25-D1.16 (25-D1.16)

Concentration: 0.2 mg/ml

Host/Isotype: Mouse IgG1, κ

Formulation: aqueous buffer, 0.09% sodium azide, may contain carrier protein/stabilizer

Temperature Limitation: Store at 2-8°C. Do not freeze. Light sensitive material.

LOT **Batch Code:** Refer to Vial

Use By: Refer to Vial

Caution, contains Azide

Description

The 25-D1.16 monoclonal antibody reacts with the 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. This antibody has proven to be very useful tracking the quantity and localization of these specific antigen-presenting cells (APC) in vivo.

Applications Reported

This eBio25-D1.16 (25-D1.16) antibody has been reported for use in flow cytometric analysis.

Applications Tested

This eBio25-D1.16 (25-D1.16) antibody has been tested by flow cytometric analysis of SIINFEKL peptide pulsed mouse splenocytes. This can be used at less than or equal to 0.125 µg per test. A test is defined as the amount (µg) of antibody that will stain a cell sample in a final volume of 100 µL. Cell number should be determined empirically but can range from 10⁵ to 10⁸ cells/test. It is recommended that the antibody be carefully titrated for optimal performance in the assay of interest.

Cells can be pulsed with the SIINFEKL peptide according to the following protocol:

1. With cells in flow staining buffer, add SIINFEKL peptide to a final concentration of 30 µM.
2. Incubate cells at 37°C for 2 hours.
3. Wash cells with flow staining buffer.
4. Proceed with cell surface staining as normal.

For additional information see the references listed below.

References

Porgador A, Yewdell JW, Deng Y, Bennink JR, Germain RN. Localization, quantitation, and in situ detection of specific peptide-MHC class I complexes using a monoclonal antibody. *Immunity*. 1997 Jun;6(6):715-26. (25-D1.16, mAb development, PubMed)

Messaoudi I, LeMaout J, Nikolic-Zugic J. The mode of ligand recognition by two peptide:MHC class I-specific monoclonal antibodies. *J*

Immunol. 1999 Sep 15;163(6):3286-94.

Ackerman AL, Kyritsis C, Tampé R, Cresswell P. Access of soluble antigens to the endoplasmic reticulum can explain cross-presentation by dendritic cells. *Nat Immunol.* 2005 Jan;6(1):107-13.

Berwin B, Hart JP, Rice S, Gass C, Pizzo SV, Post SR, Nicchitta CV. Scavenger receptor-A mediates gp96/GRP94 and calreticulin internalization by antigen-presenting cells. *EMBO J.* 2003 Nov 17;22(22):6127-36.

Related Products

17-4714 Mouse IgG1 K Isotype Control APC

Not for further distribution without written consent.

Copyright © 2000-2010 eBioscience, Inc.

Tel: 888.999.1371 or 858.642.2058 • Fax: 858.642.2046 • www.eBioscience.com • info@eBioscience.com