

PRODUCT INSERT

MONOCLONAL ANTIBODY TO THE MOUSE H-2K<sup>k</sup> ANTIGEN

Product	Form	Volume	Antibody*	Excitation	Peak Emission	Matching Isotype Controls	
				(nm)	(nm)		
MM4201	FITC	1.0 ml	500 µg	488	525	Mouse IgG2a FITC	MG2a01
MM4204	R-PE	1.0 ml	100 µg	488	575	Mouse IgG2a R-PE	MG2a04

**PRODUCT DESCRIPTION**

Mouse monoclonal antibody to the mouse H-2K<sup>k</sup> antigen

**Clone:** 36-7-5

**Isotype:** Mouse IgG<sub>2a</sub>K

**Lot No.:** See label      **Expiration:** See label

**Buffer:** Phosphate buffered saline (PBS)

**Preservatives:** 0.1% *sodium azide*. Sodium azide is an extremely toxic and dangerous compound particularly when combined with acids or metals. Solutions containing sodium azide should be disposed of properly.

**Stabilizer:** Sucrose.

**PRODUCT CHARACTERIZATION**

**Antigen Specificity:** MAb 36-7-5 reacts with the H-2K<sup>k</sup> MHC class I alloantigen. Cross-reactivity with splenocytes of SJL/Hsd mice has been observed by flow cytometric analysis. The antibody does not react with other (e.g., *b, d, q*) haplotypes.

**Research Applications:**

- Flow cytometry
- Complement-dependent cytotoxicity
- Immunohistochemistry (acetone-fixed, frozen sections only)

**Note:** Flow cytometric data shown may not necessarily have been generated using the enclosed lot of reagent. For this reason, and due to differences in flow cytometers and cytometer settings, results may vary from those illustrated above. It is suggested that investigators titrate reagents to determine optimal conditions for use in their systems.

**STORAGE & HANDLING**

Store reagents at 2-8°C. Light exposure should be avoided for fluorochrome-conjugated reagents. Use dim light during handling, incubation with cells and prior to analysis. It is recommended that cells be analyzed within 18 hours of staining. If the reagent is being diluted, it is recommended that only the quantity to be used within one week be diluted.

**PRODUCT QUALITY CONTROL**

To ensure lot-to-lot consistency, each batch of monoclonal antibody is tested by flow cytometry to conform to characteristics of a standard reference reagent. From this testing it is recommended that between 0.1 and 0.2 µg of antibody be used per 1 x 10<sup>6</sup> cells in a 100 µl staining volume. Because conditions may vary, it is recommended that each investigator determine the optimal amount of antibody to be used for each application.

\* The amount of antibody is determined by measuring the optical density using a spectrophotometer. The antibody titer is verified by immunofluorescent staining and flow cytometric analysis.

**REFERENCES:**

1. Sachs, D.H. Personal communication

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