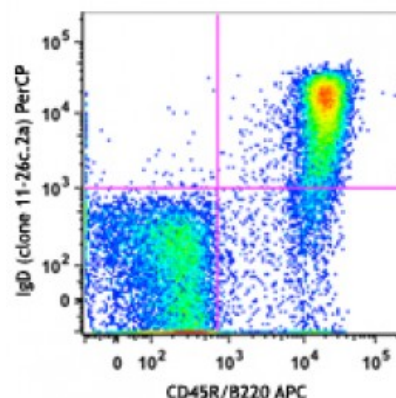


**PerCP anti-mouse IgD**

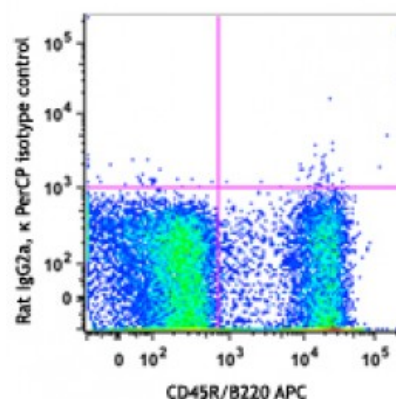
|                          |  |
|--------------------------|--|
| <b>Catalog # / Size:</b> | 2628675 / 25 µg<br>2628680 / 100 µg  |
| <b>Clone:</b>            | 11-26c.2a  |
| <b>Isotype:</b>          | Rat IgG2a, κ   |
| <b>Reactivity:</b>       | Mouse  |
| <b>Preparation:</b>      | The antibody was purified by affinity chromatography and conjugated with PerCP under optimal conditions. The solution is free of unconjugated PerCP and unconjugated antibody. |
| <b>Formulation:</b>      | Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.  |
| <b>Concentration:</b>    | 0.2  |



C57BL/6 mouse splenocytes were stained with CD45R/B220 APC and IgD (clone 11-26c.2a) PerCP (top) or rat IgG2a, κ PerCP isotype control (bottom).

**Applications:**

|                           |   |
|---------------------------|---|
| <b>Applications:</b>      | Flow Cytometry  |
| <b>Recommended Usage:</b> | 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. |



\* PerCP has a maximum absorption of 482 nm and a maximum emission of 675 nm.

|                           |   |
|---------------------------|---|
| <b>Application Notes:</b> | The 11-26c.2a antibody reacts with immunoglobulin D in all tested mouse haplotypes. The antibody binds membrane IgD expressed on most B cells. The 11-26c.2a antibody neither induces proliferation of splenic B cells nor induces B cell activation. Additional reported applications (for the relevant formats) include: immunohistochemical staining of acetone-fixed frozen sections <sup>2,3</sup> . |
|---------------------------|---|

|                                |  |
|--------------------------------|--|
| <b>Application References:</b> | <ol style="list-style-type: none"> <li>1. Nitschke L, <i>et al.</i> 1993. <i>P. Natl. Acad. Sci. USA</i> 90:1887. (FC)</li> <li>2. Weih D, <i>et al.</i> 2001. <i>J. Immunol.</i> 167:1909. (IHC)</li> <li>3. Koni PA, <i>et al.</i> 2001. <i>J. Exp. Med.</i> 193:741. (IHC)</li> <li>4. Ahuja A, <i>et al.</i> 2007. <i>J. Immunol.</i> 179:3351. (FC) <a href="#">PubMed</a></li> <li>5. Haynes NM, <i>et al.</i> 2007. <i>J. Immunol.</i> 179:5099. (FC)</li> <li>6. Good-Jacobson KL, <i>et al.</i> 2010. <i>Nat. Immunol.</i> 11:535. (FC) <a href="#">PubMed</a></li> </ol> |
|--------------------------------|--|

7. Tomayko MM, *et al.* 2010. *J. Immunol.* 185:7146. [PubMed](#)  
8. Park SY, *et al.* 2013. *J. Immunol.* 190:1094. [PubMed](#)
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**Description:** Surface IgD is an important B cell differentiation marker.