

# Rat IgG2a kappa Isotype Control (eBR2a), APC, eBioscience<sup>TM</sup>

## Product Details

|                         |                                    |
|-------------------------|------------------------------------|
| Size                    | 50 µg                              |
| Host/Isotype            | Rat / IgG2a, kappa                 |
| Class                   | Monoclonal                         |
| Type                    | Isotype Control                    |
| Clone                   | eBR2a                              |
| Conjugate               | APC                                |
| Excitation/Emission Max | 651/660 nm                         |
| Form                    | Liquid                             |
| Concentration           | 0.2 mg/mL                          |
| Purification            | Affinity chromatography            |
| Storage buffer          | PBS, pH 7.2                        |
| Contains                | 0.09% sodium azide                 |
| Storage conditions      | 4°C, store in dark, DO NOT FREEZE! |
| RRID                    | AB_470181                          |

## Applications

## Tested Dilution

## Publications

|   |                 |               |
|---|-----------------|---------------|
| Immunohistochemistry (Paraffin) (IHC (P)) | -               | 0 Publication |
| Flow Cytometry (Flow)                     | Assay-Dependent | 0 Publication |
| Control (Ctrl)                            | Assay-Dependent | 0 Publication |

## Product Specific Information

Description: The monoclonal rat IgG2a kappa is useful as an isotype control immunoglobulin.

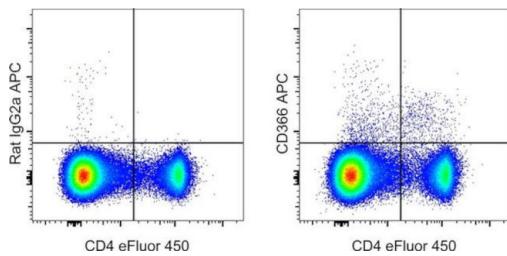
Applications Reported: This eBR2a antibody has been reported for use in flow cytometric analysis.

Applications Tested: This eBR2a antibody has been tested by flow cytometric analysis of mouse splenocytes and normal human peripheral blood cells and should be used at the same concentration as the experimental antibody.

Excitation: 633-647 nm; Emission: 660 nm; Laser: Red Laser.

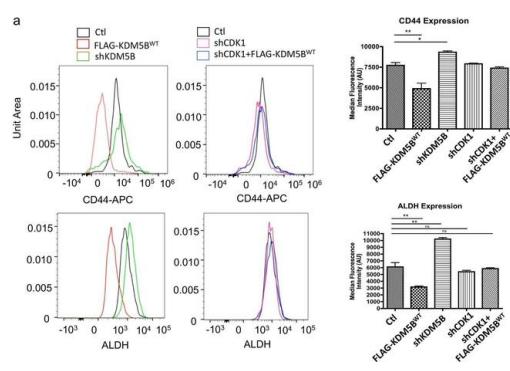
Filtration: 0.2 µm post-manufacturing filtered.

# Product Images For Rat IgG2a kappa Isotype Control (eBR2a), APC, eBioscience™



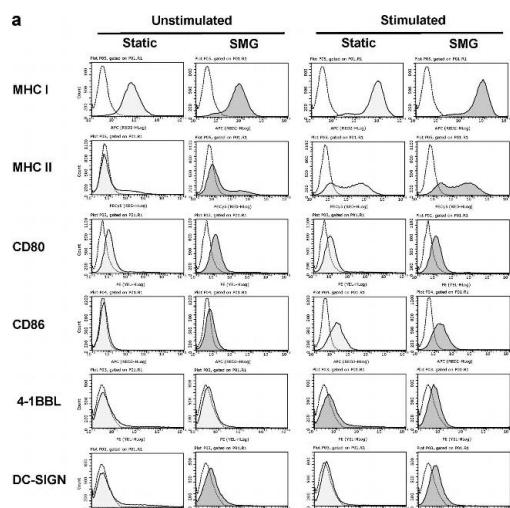
## Rat IgG2a kappa Isotype Control (17-4321-81) in Flow

Staining of C57BL/6 splenocytes with Anti-Mouse CD4 eFluor® 450 (Product # 48-5870) and 0.125 µg of Rat IgG2a K Isotype Control APC (Product # 17-4321-81) (left) or 0.125 µg of Anti-Mouse CD366 APC (right). Cells in the lymphocyte gate were used for analysis.



## Rat IgG2a kappa Isotype Control (17-4321-81) in Flow

KDM5B induces a reduction in the CSC population in TNBC and role of CDK1 in KDM5B action. MDA-MB-231 cells were transfected as indicated with expression vectors for FLAG-KDM5BWT, shKDM5B, and/or shCDK1. Shown are (a) flow cytometry analyses of ALDH+ (aldehyde dehydrogenase) and CD44-APC (allophycocyanin) positive populations including mean fluorescence values (MFI) and (b) mammosphere formation assay. (c) MDA-MB-231 cells were transfected with expression vectors for FLAG-KDM5BWT, -KDM5BS1456A or -KDM5BS1456D. Transfected cells were cultured in mammosphere media and colonies were quantified. Figures are representative of at least 3 independent experiments. Image collected and cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/31776402/>), licensed under a CC BY license.



## Rat IgG2a kappa Isotype Control (17-4321-81) in Flow

JAWS II DC undergo maturation when cultured in SMG. (a) Unstimulated and stimulated JAWS II DC ( $2 \times 10^5$ /ml) were cultured in Static (light gray) or SMG (dark gray) conditions for 72 h. JAWS II DC activated during Static or SMG culture were either incubated with (stimulated) or without (unstimulated) a cocktail of cytokines (IFN-, IL-4 and TNF). Afterwards, the cells were collected and stained with antibodies for MHC class I and II, CD80, CD86, 4-1BBL and DC-SIGN and their corresponding isotype controls. Open histograms with broken lines represent isotype controls. (b) Graph represents the MFI of each of the surface molecules examined in (a) for unstimulated Static (light gray bars) and SMG JAWS II DC (black bars). (c) Culture supernatants, collected from unstimulated and stimulated JAWS II DC ( $2 \times 10^5$ /ml) cultured as in (a), were assessed for IL-6 production by ELISA. One representation of two independent experiments with similar results is shown for (a,b). For (c), the data represents the mean + SD of quadruplicates of two independent experiments. In (b), \*p-value 0.05 comparing the expression of surface molecules by SMG and Static JAWS II DC. In (c), \*p-value 0.05 comparing the production of IL-6 by unstimulated and stimulated SMG to Static JAWS II DC. Image collected and cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/31554863/>), licensed under a CC BY license.

View more figures on [thermofisher.cn](http://thermofisher.cn)

## 93 References

Embryo-restricted responses to maternal IL-17A promote neurodevelopmental disorders in mouse offspring. *Mol Psychiatry* (2025)

A Novel Approach to Peripheral Nerve Regeneration: Local FK-506 Delivery Using a Reservoir Flap Model. *Yonsei Med J* (2024)

A synbiotic mixture of *Bifidobacterium breve* M16-V, oligosaccharides and pectin, enhances Short Chain Fatty Acid production and improves lung health in a preclinical model for pulmonary neutrophilia. *Front Nutr* (2024)

Endothelial cell signature in muscle stem cells validated by VEGFA-FLT1-AKT1 axis promoting survival of muscle stem cell. *Elife* (2024)

GM-CSF-dependent CD301b+ lung dendritic cells confer tolerance to inhaled allergens *Research Square* (2024)

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