

RayBiotech, Inc.

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Certificate of Analysis and Data Sheet

Mouse Anti-BrdU

With HRP-conjugated Secondary Antibody

Catalog No.

DS-MB-00143

Target Species

Chemical

Isotype

IgG1

Synonyms

5- Bromodeoxyuridine

Preparation

Purification: Purified IgG prepared by ion exchange chromatography from tissue culture supernatant

Formulation

Product Type: Monoclonal Antibody

Product Form: Purified Purified IgG - liquid

Buffer: Phosphate buffered saline

Preservative Stabilizers: 0.09% Sodium Azide

Specificity

DS-MB-00143 recognizes the thymidine analogue bromodeoxyuridine (BrdU), which can be incorporated into DNA during S phase of the cell cycle. The antibody is suitable for detecting incorporated BrdU in a wide variety of cell types and is suitable for use on tissue sections in double-labeling techniques.

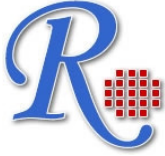
Storage

Store at +4°C or at -20°C if preferred. This product should be stored undiluted.

Storage in frost free freezers is not recommended. This product is photosensitive and should be protected from light. Avoid repeated freezing and thawing as this may denature the antibody. Should this product contain a precipitate we recommend microcentrifugation before use.

Shelf Life: 18 months from date of shipment

**The products are furnished for LABORATORY RESEARCH USE ONLY.
Not for diagnostic or therapeutic use.**



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Recommended Protocol

FLOW CYTOMETRY ANALYSIS

Prepare the following solutions before proceeding:

Phosphate buffered saline (PBS)

2N HCl containing 0.5% Triton X-100

PBS containing 0.05% Tween-20

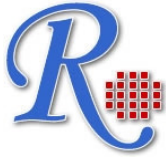
PBS containing 1% BSA (PBS/BSA)

10mg/ml Propidium iodide (PI)

0.1M Na₂B₄O₇, pH 8.5

1. Add BrdU to the cell suspension in culture medium to a final concentration of 10 µmol/L and incubate for 30 minutes in a CO₂ incubator at 37°C.
2. Wash cells twice with PBS/BSA by centrifuging at 500g for 10 minutes, decant supernatant and resuspend in a minimum volume of PBS.
3. Add cells slowly into 5ml of 70% ethanol at -20°C, mixing continuously (vortex preferred). Incubate on ice for 30 minutes.
4. Centrifuge at 500g for 10 minutes, decant supernatant, and resuspend cell pellet.
5. Add 2ml of 2N HCl containing 0.5% Triton X-100 and incubate the cells for 30 minutes at room temperature (preferably on a rocking platform).
6. Centrifuge at 500g for 10 minutes, decant supernatant and resuspend in 3 ml of 0.1M Na₂B₄O₇, pH 8.5.
7. Centrifuge at 500g for 10 minutes, decant supernatant and resuspend the cells in PBS/BSA + 0.05% Tween-20. Adjust cell concentration to 1 x 10⁷/ml.
8. Aliquot 100ul of cell suspension into required number of 12 x 75mm tubes.
9. Incubate the cells with the BrdU antibody at the recommended dilution for 45 minutes at room temperature or overnight at 4°C.
10. Add 2 ml of PBS/BSA and centrifuge the cells at 1000rpm for 5 minutes.
11. If a secondary antibody layer is required then decant the supernatant and incubate the cells with the secondary antibody for 30 minutes at room temperature. If no secondary antibody layer is required then proceed to step 13.
12. Wash the cells after the secondary antibody layer by repeating step 10.

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13. Decant the supernatant and add 1ml of PBS containing 10 μ g/ml PI (Dilute the 10mg/ml solution of PI 1/1000 in a suitable volume of PBS).

14. Analyze cells by flow cytometry following the manufacturer's instructions. The PI should be read on the appropriate channel set to the Peak/Area and not log scale.

Applications

Table Summary of antibody applications and working conditions

| Options / Functions | YES | NO | Not determined | Recommended Work dilution or concentration |
|---------------------|-----|----|----------------|--|
| ELISA | | | • | |
| Western Blotting | | | • | |
| IHC – Paraffin | • | | | |
| IHC - Frozen | • | | | |
| Flow Cytometry | • | | | 1/25-1/100 |

Note: Other applications are not tested yet. Optimal dilutions should be determined by each laboratory for each application.

Secondary Antibody Applications

| Options / Functions | YES | NO | Not determined | Recommended Work dilution or concentration |
|-----------------------------------|-----|----|----------------|--|
| Immunoassay (ELISA, Western blot) | • | | | 1:5000-1:10000 |

Reference

1. Magaud, J.P. et al. (1989) Double Immunocytochemical labelling of cell and tissue samples with monoclonal anti-bromodeoxyuridine. *J Histochem Cytochem.* 37:1517 – 1527.
2. Innis, S.M. et al. (2010) Perinatal Lipid Nutrition Alters Early Intestinal Development and Programs the Response to Experimental Colitis in Young Adult Rats. *Am J Physiol Gastrointest Liver Physiol.* 2010 Sep 23.
3. Caronia, G. et al. (2010) Bone morphogenetic protein signaling in the developing telencephalon controls formation of the hippocampal dentate gyrus and modifies fear-related behavior. *J Neurosci.* 30: 6291-301.

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