

Instructions For Use

RA0375-C.5-IFU-RUO

Rev. Date: Jan. 2, 2015

Revision: 1

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P.O. Box 3286 - Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Tel. (435) 755-9848 - Fax (435) 755-0015 - www.scytek.com

Bromodeoxyuridine (BrdU) (Proliferation Marker); Clone BU20a (Concentrate)

Availability/Contents: Item #_ RA0375-C.5 Volume 0.5 ml

Description:

Species: Mouse

Immunogen: Bromodeoxyuridine (BrdU) conjugated to KLH

Clone: BU20a Isotype: IgG1

Entrez Gene ID: Not Applicable
Hu Chromosome Loc.: Not Applicable

Synonyms: Bromodeoxyuridine, BUdr Mol. Weight of Antigen: Depends on the target

Format: 200µg/ml of Ab purified from Bioreactor Concentrate by Protein A/G. Prepared in 10mM PBS

with 0.05% BSA & 0.05% azide.

Specificity: This antibody reacts with Bromodeoxyuridine (BrdU) in single stranded DNA (produced by

partial denaturation of double stranded DNA), BrdU coupled to a protein carrier, as well as free

BrdU.

Background: BrdU is a thymidine analog, incorporated into cell nuclei during DNA synthesis prior to mitosis.

An antibody to BrdU is helpful in detecting S-phase cells, providing useful information on the

aggressiveness of tumors.

Species Reactivity: All species

Positive Control: Cells grown in presence of BrdU or tissues from experimental animals injected with BrdU

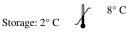
Cellular Localization: Nuclear

Titer/ Working Dilution: Immunohistochemistry (Frozen and Formalin-fixed): 0.5-1 μg/ml

Flow Cytometry: 0.5-1 µg/million cells

Immunofluorescence: $0.5-1 \mu g/ml$

Microbiological State: This product is not sterile.







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Uses/Limitations: Not to be taken internally.

For Research Use Only.

This product is intended for qualitative immunohistochemistry with normal and neoplastic formalin-fixed, paraffin-embedded

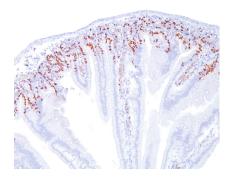
tissue sections, to be viewed by light

microscopy.

Do not use if reagent becomes cloudy. Do not use past expiration date.

Non-Sterile.

Ordering Information and Current Pricing at www.scytek.com



Formalin-fixed, paraffin-embedded mouse intestine tissue (10X) stained with BrdU; Clone BU20a.

Procedure:

- 1. **Tissue Section Pretreatment (Required):** Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by incubating sections in 4N HCl for 30 minutes at room temperature followed by digestion with Trypsin (Two Component Solution) (ScyTek catalog# TSS).
- Primary Antibody Incubation Time: We suggest an incubation period of 30 minutes at room temperature.
 However, depending upon the fixation conditions and the staining system employed, optimal incubation should be determined by the user.
- 3. **Visualization:** For maximum staining intensity we recommend the "UltraTek HRP Anti-Polyvalent Lab Pack" (ScyTek catalog# UHP125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).

Precautions:

Contains Sodium Azide as a preservative (0.09% w/v).

Do not pipette by mouth.

Avoid contact of reagents and specimens with skin and mucous membranes.

Avoid microbial contamination of reagents or increased nonspecific staining may occur.

This product contains no hazardous material at a reportable concentration according to U.S. 29 CFR 1910.1200,

OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

References:

- 1. Welberg JW et. al. Journal of Clinical Pathology, 1990, 43(6):453-6.
- 2. Williams LS et. al. Cytometry, 1990, 11(4):490-7.

Warranty:

No products or "Instructions For Use (IFU)" are to be construed as a recommendation for use in violation of any patents. We make no representations, warranties or assurances as to the accuracy or completeness of information provided on our IFU or website. Our warranty is limited to the actual price paid for the product. ScyTek Laboratories, Inc. is not liable for any property damage, personal injury, time or effort or economic loss caused by our products. Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining can cause inconsistent results. Variations in fixation and embedding or the inherent nature of the tissue specimen may cause variations in results. Endogenous peroxidase activity or pseudoperoxidase activity in erythrocytes and endogenous biotin may cause non-specific staining depending on detection system used.

Storage: 2° C

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