

# Hydrogen Peroxide Blocking Reagent ab64218

★★★★★ [3 Abreviews](#) [23 References](#) [1 Image](#)

## Overview

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<b>Product name</b>	Hydrogen Peroxide Blocking Reagent
<b>Tested applications</b>	<b>Suitable for:</b> IHC-P, IHC-Fr
<b>General notes</b>	<p>Hydrogen Peroxide Blocking Reagent ab64218 for use in IHC with detection based on HRP / peroxidase.</p> <p>Note: The concentration of hydrogen peroxide in this product is 0.3%.</p> <p><b>IHC protocol suitable for use with Hydrogen Peroxide Blocking Reagent:</b> For frozen sections, skip steps 1 and 2.</p> <ol style="list-style-type: none"><li>1. Deparaffinize and rehydrate formalin-fixed paraffin-embedded tissue section.</li><li>2. Use appropriate <b>antigen retrieval buffer or enzyme</b> (primary antibody dependent) to treat sections. Wash 3 times in buffer.</li><li>3. <b>Add enough hydrogen peroxide blocking solution ab64218 to cover the sections. Incubate for 10 minutes.</b> Wash 2 times in buffer. If necessary, use <b>avidin biotin blocking</b>.</li><li>4. Apply <b>protein block</b> (or <b>normal serum</b> from same species as secondary antibody) and incubate for 5 minutes at room temperature to block nonspecific background staining. Wash once in buffer.</li><li>5. Apply primary antibody in <b>antibody diluent</b> and incubate.</li><li>6. Wash 4 times in buffer. Incubate slide with <b>biotinylated secondary antibody</b> (or <b>HRP polymer secondary antibody</b> and skip step 7). Wash 4 times in buffer.</li><li>7. Apply <b>streptavidin-HRP</b> and incubate for 10 minutes at room temperature.</li><li>8. Rinse 4 times in buffer. Place slide in <b>DAB substrate</b> or <b>AEC substrate</b> and incubate until desired color is achieved (1-10 mins). Rinse 4 times in buffer.</li><li>9. Add enough drops of <b>hematoxylin</b> to cover the section. Incubate for 1 minute.</li><li>10. Rinse 7-8 times in tap water. Add <b>mounting medium</b> to cover the section.</li></ol> <p>Find complete IHC kits, and reagents for antigen retrieval, blocking, signal amplification, visualization, counterstaining, and mounting in the <b><a href="#">IHC kits and reagents guide</a></b>.</p>

## Properties

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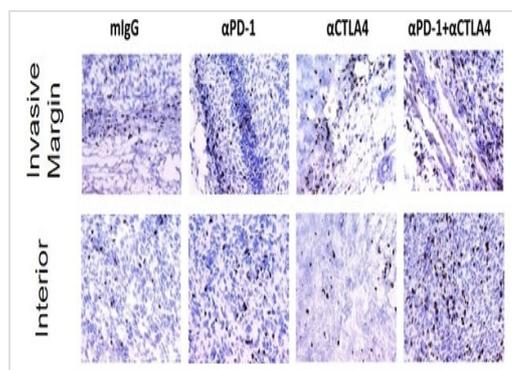
<b>Form</b>	Liquid
<b>Storage instructions</b>	Store at +4°C.
<b>Storage buffer</b>	Preservative: 0.1% Sodium azide Constituent: 0.3% Hydrogen peroxide
<b>Relevance</b>	Hydrogen Peroxide Blocking Reagent is intended for use in peroxidase-based immunohistochemical staining procedures on cell preparations, frozen tissue sections, and paraffin-embedded tissue sections.

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab64218 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		Use at an assay dependent dilution.
IHC-Fr		Use at an assay dependent dilution.

## Images



Immunohistochemical (frozen) analysis of MC38 tumor sections labeling CD3 with **ab16669** at 1/400 dilution. Sections were fixed with acetone, treated with peroxidase block (ab64218) to quench endogenous peroxidase, and then further blocked with a 10% goat serum and 5% BSA solution. CD3 positive T cells were detected using **ab80437**. Invasive margin (top) shows sections derived from the periphery of the tumor and interior (bottom) shows sections from within the tumor.

Immunohistochemistry (Frozen sections) - Hydrogen Peroxide Blocking Reagent (ab64218)

Selby MJ et al., PLoS One 11(9), Fig 2. doi: 10.1371/journal.pone.0161779. Reproduced under the Creative Commons license <http://creativecommons.org/licenses/by/4.0/>

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