

### Anti-CD11b + CD11c antibody [OX42] ab1211

★★★★★ [6 Abreviews](#) [143 References](#) [4 Images](#)

#### Overview

<b>Product name</b>	Anti-CD11b + CD11c antibody [OX42]
<b>Description</b>	Mouse monoclonal [OX42] to CD11b + CD11c
<b>Host species</b>	Mouse
<b>Tested applications</b>	<b>Suitable for:</b> IHC-Fr, ICC/IF <b>Unsuitable for:</b> IHC-P
<b>Species reactivity</b>	<b>Reacts with:</b> Rat <b>Does not react with:</b> Mouse
<b>Immunogen</b>	Tissue, cells or virus corresponding to CD11b. Resident peritoneal macrophages from (PVG.RT1[c] x PVG.RT1[u]) and (PVG.RT1[c] x PVG.RT1[a]) F1-hybrid rat.
<b>Positive control</b>	ICC/IF: NR8383 cells. IHC-Fr: Rat spleen and lung tissue sections (10µm).
<b>General notes</b>	<p><b><u>IHC protocol advice:</u></b></p> <p>This antibody is not suitable for IHC on paraffin-embedded samples.</p> <p>For best results in IHC on frozen tissue, the following may help detection:</p> <ol style="list-style-type: none"> <li>1) Paraformaldehyde perfusion fixed samples have worked well for many customers.</li> <li>2) For non-perfused tissue, either snap freeze or immerse in periodate-lysine-paraformaldehyde (PLP) fixative for 24 hours at 4°C. Reduce the concentration of paraformaldehyde to 0.25-0.5% since this increases the staining intensity for immune cell surface markers (PMID: 7868861).</li> <li>3) PFA-fixed samples will require cryoprotection by sucrose infiltration.</li> <li>4) For snap frozen tissue, fix sections in cold acetone for 10 min. Allow to dry for 10 min at room temperature. Wash with water for 10 min.</li> <li>5) Do not heat the samples or sections.</li> <li>6) During the staining procedure, do not allow the sections to dry out.</li> </ol> <p>Our Technical team (<a href="mailto:technical@abcam.com">technical@abcam.com</a>) will be happy to provide further information and advice.</p> <p>This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or conjugation for your experiments, please contact <a href="mailto:orders@abcam.com">orders@abcam.com</a>.</p> <p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be</p>

found below, along with publications, customer reviews and Q&As

## Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.
<b>Storage buffer</b>	pH: 7.40 Preservative: 0.02% Sodium azide Constituent: PBS  Some batches contain 6.97% L-Arginine as a stabilizing agent. For lot-specific buffer information, please contact our Scientific Support team.
<b>Purity</b>	Protein G purified
<b>Primary antibody notes</b>	The principal use of this antibody is in the study of functional heterogeneity of macrophages; it may be used to follow macrophage differentiation and to investigate the function of the CD11b + CD11c equivalent antigen.
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	OX42
<b>Isotype</b>	IgG2a
<b>Light chain type</b>	kappa

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab1211 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

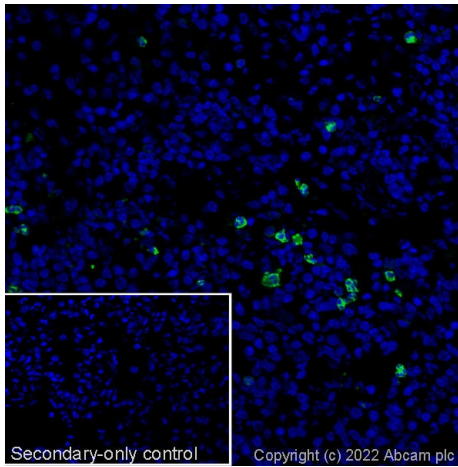
Application	Abreviews	Notes
IHC-Fr	★★★★★ (2)	Use at an assay dependent concentration.
ICC/IF		Use a concentration of 1 - 5 µg/ml.

**Application notes** Is unsuitable for IHC-P.

## Target

**Cellular localization** CD11b: Membrane. CD11c: Membrane.

## Images



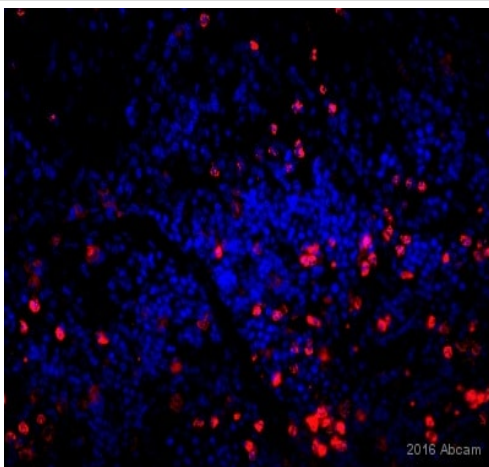
Immunohistochemistry (Frozen sections) - Anti-  
CD11b + CD11c antibody [OX42] (ab1211)

Immunofluorescence staining of CD11b + CD11c staining in a section of 10% formalin fixed (10 mins, RT) frozen rat spleen.

Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab1211 at 1µg/ml. The section was then incubated with **ab150117** (Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) preabsorbed, (Shown in green) 1.5µg/ml) for 1 hour at room temperature. Nuclear DNA was labelled with DAPI (shown in blue). The section was then mounted using Fluoromount®.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

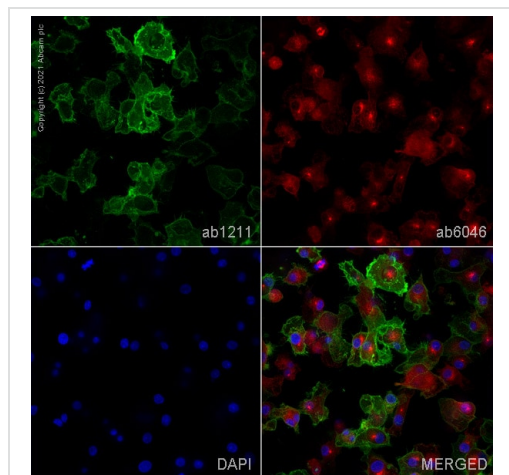
For other IHC staining systems (automated and non-automated), customers should optimize variable parameters such as antigen retrieval conditions, antibody concentrations and incubation times.



Immunohistochemistry (Frozen sections) - Anti-  
CD11b + CD11c antibody [OX42] (ab1211)

Image courtesy of Carl Hobbs.

ab1211 staining CD11b + CD11c (shown in red) in rat spleen tissue sections by IHC (PFA perfusion fixed frozen sections). Tissue samples were fixed with formaldehyde and blocked with BSA for 10 minutes at 21°C. The sample was incubated with primary antibody (1/2000 in TBS/BSA/azide) at 21°C for 16 hours. A Biotin-conjugated Goat anti mouse polyclonal (1/300) was used as the secondary antibody.

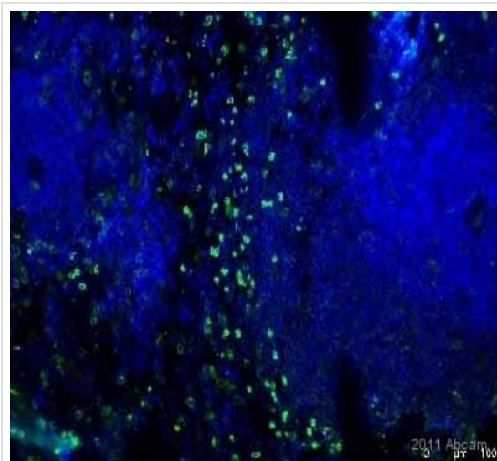


Immunocytochemistry/ Immunofluorescence - Anti-CD11b + CD11c antibody [OX42] (ab1211)

ab1211 staining CD11b + CD11c in NR8383 cells. The cells were fixed with 4% paraformaldehyde (10 min), permeabilized with 0.1% PBS-Tween for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab1211 at 1 µg/ml and **ab6046**, Rabbit polyclonal to beta Tubulin - Loading Control. Cells were then incubated with **ab150117**, Goat polyclonal Secondary Antibody to Mouse IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (shown in green) and **ab150080**, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 594) at 1/1000 dilution (shown in pseudocolour red). Nuclear DNA was labelled with DAPI (shown in blue).

Also suitable in cells fixed with 100% methanol (5 min).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal sections is shown.



Immunohistochemistry (Frozen sections) - Anti-CD11b + CD11c antibody [OX42] (ab1211)  
Image courtesy of an anonymous Abreview.

ab1211 staining CD11b + CD11c (shown in green) in rat spleen tissue by IHC (Frozen sections). Tissue was fixed in formaldehyde, blocked with 10% serum for 30 minutes at 24°C, then incubated with ab1211 at a 1/100 dilution. The secondary used was an Alexa-Fluor 488 conjugated goat anti-mouse polyclonal, used at a 1/1000 dilution.

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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