# abcam

## Product datasheet

## Anti-CD4 antibody [SP35] - BSA and Azide free ab238798



## 9 Images

#### Overview

**Product name** Anti-CD4 antibody [SP35] - BSA and Azide free

**Description** Rabbit monoclonal [SP35] to CD4 - BSA and Azide free

**Host species** Rabbit

**Tested applications** Suitable for: IHC-P, Flow Cyt, mIHC

Reacts with: Human **Species reactivity** 

Predicted to work with: Pig ...

**Immunogen** Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control IHC-P: Human Hodgkin's lymphoma and tonsil tissue. Flow Cyt: THP-1 cells

General notes ab238798 is the carrier-free version of ab213215.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our **conjugation kits** for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

This product is FOR RESEARCH USE ONLY. For commercial use, please contact partnerships@abcam.com.

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#### **Properties**

Form Liquid

**Storage instructions** Shipped at 4°C. Store at +4°C. Do Not Freeze.

Storage buffer pH: 7.20

Constituent: PBS

Carrier free Yes

Purity Protein A/G purified

**Purification notes** Purified from TCS by protein A/G.

**Clonality** Monoclonal

Clone number SP35

**Isotype** IgG

## **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab238798 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 concentration. (For 30 minutes at room temperature. Antigen Retrieval: Boil tissue section in EDTA buffer for 10 min followed by cooling at room temperature for 20 min).
Flow Cyt		Use at an assay dependent concentration. (For 30 minutes at 4°C).
mIHC		1/500. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2)

## **Target**

Function Accessory protein for MHC class-II antigen/T-cell receptor interaction. May regulate T-cell

activation. Induces the aggregation of lipid rafts.

Sequence similarities Contains 3 lg-like C2-type (immunoglobulin-like) domains.

Contains 1 lg-like V-type (immunoglobulin-like) domain.

Post-translational modifications

Palmitoylation and association with LCK contribute to the enrichment of CD4 in lipid rafts.

Cellular localization Cell membrane. Localizes to lipid rafts. Removed from plasma membrane by HIV-1 Nef protein

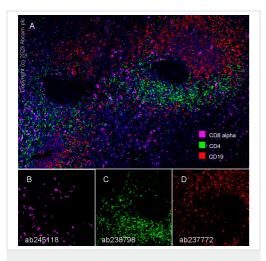
that increases clathrin-dependent endocytosis of this antigen to target it to lysosomal degradation.

Cell surface expression is also down-modulated by HIV-1 Envelope polyprotein gp160 that

interacts with, and sequesters CD4 in the endoplasmic reticulum.

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#### **Images**



Multiplex immunohistochemistry - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)

Multiplex immunohistochemistry analysis of formalin/PFA-fixed paraffin-embedded Human spleen tissue labeling CD8 alpha with **ab245118** at 1/500 dilution, CD4 with ab238798 at 1/500, and CD19 with **ab237772** at 1/5000 dilution.

Panel A: merged staining of anti-CD8 alpha (magenta; Opal™690), anti-CD4 (green; Opal™520) and anti-CD19 (red; Opal™570) on human spleen.

Panel B: anti-CD8 alpha stained on cytotoxic T cells.

Panel C: anti-CD4 stained on T helper cells.

Panel D: anti-CD19 stained on B cells.

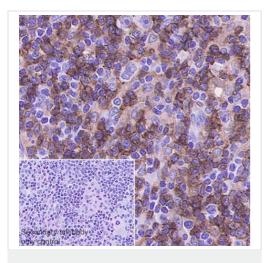
Sections were treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins before antibody incubation. The section was incubated in three rounds of staining: in the order of <a href="mailto:ab245118">ab245118</a> for 30 mins, then ab238798 and <a href="mailto:ab237772">ab237772</a> for 10 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

DAPI was used as a nuclear counterstain.

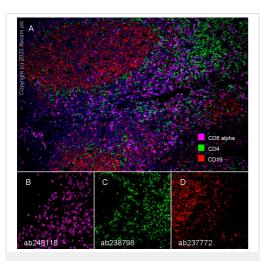
The immunostaining was performed on a Leica Biosystems
BOND® RX instrument with an Opal™ 4-color kit. Image acquisition
was performed with Leica SP8 confocal microscope.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human Hodgkin lymphoma tissue sections labeling CD4 with <u>ab213215</u> at 1/50 dilution (3.1 μg/ml). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 10mins. Rabbit specific IHC polymer detection kit HRP/DAB (<u>ab209101</u>) was used as the secondary antibody. Hematoxylin was used as a counterstain. Positive staining on the human Hodgkin's lymphoma, performed on a Leica Biosystems BOND<sup>TM</sup> RX instrument.

The section was incubated with <u>ab213215</u> for 30 mins at room temperature. This image was generated using <u>ab213215</u>, the same clone, but with a different buffer formulation.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)



Multiplex immunohistochemistry - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)

Multiplex immunohistochemistry analysis of formalin/PFA-fixed paraffin-embedded Human tonsil tissue labeling CD8 alpha with **ab245118** at 1/500 dilution, CD4 with ab238798 at 1/500, and CD19 with **ab237772** at 1/5000 dilution.

Panel A: merged staining of anti-CD8 alpha (magenta; Opal™690), anti-CD4 (green; Opal™520) and anti-CD19 (red; Opal™570) on human tonsil.

Panel B: anti-CD8 alpha stained on cytotoxic T cells.

Panel C: anti-CD4 stained on T helper cells.

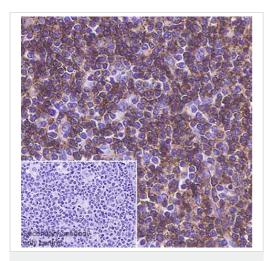
Panel D: anti-CD19 stained on B cells.

Sections were treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins before antibody incubation. The section was incubated in three rounds of staining: in the order of <a href="mailto:ab238798">ab238798</a> and <a href="mailto:ab2387772">ab237772</a> for 10 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

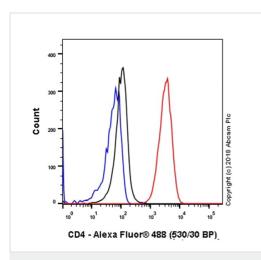
DAPI was used as a nuclear counterstain.

The immunostaining was performed on a Leica Biosystems
BOND® RX instrument with an Opal™ 4-color kit. Image acquisition
was performed with Leica SP8 confocal microscope.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human tonsil tissue sections labeling CD4 with ab213215 at 1/50 dilution (3.1 μg/ml). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 10mins. Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) was used as the secondary antibody. Hematoxylin was used as a counterstain. Positive staining on the human tonsil, performed on a Leica Biosystems BOND<sup>TM</sup> RX instrument. The section was incubated with ab213215 for 30 mins at room temperature. This image was generated using ab213215, the same clone, but with a different buffer formulation.



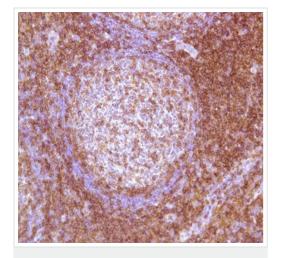
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)



Flow Cytometry - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)

Flow cytometry analysis of THP-1 (human acute monocytic leukemia) labeling CD4 with purified <u>ab213215</u> at 1/20 dilution (7.75  $\mu$ g/ml) (red). Goat anti rabbit lgG (Alexa Fluor<sup>®</sup> 488, <u>ab150081</u>) at 1/2000 dilution was used as a secondary antibody. lsotypecontrol - Rabbit monoclonal lgG (<u>ab172730</u>) (black). Unlableled control - Unlabelled cells (blue).

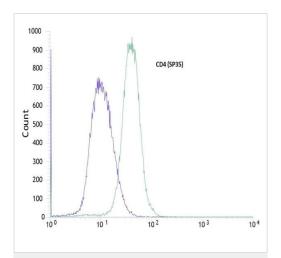
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab213215).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)

Immunohistochemical analysis of formalin-fixed, paraffin-embedded Human tonsil tissue labeling CD4 with <u>ab213215</u> at 1/50 dilution.

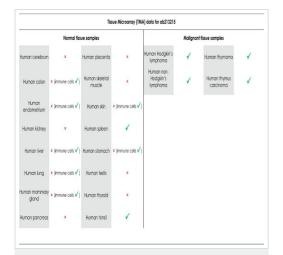
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, and sodium azide (ab213215).



Flow Cytometry - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)

Flow cytometric analysis of rabbit anti-CD4 (SP35) antibody, prediluted, <u>ab101530</u> in Jurkats cells (green) compared to negative control of rabbit lgG (blue)

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab101530).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD4 antibody [SP35] - BSA and Azide free (ab238798)

Tissue Microarrays stained for "Anti-CD4 antibody [SP35]" using "

ab213215" in immunohistochemical analysis. This table provides a

detailed overview of positive (tick mark) and negative (cross mark)

staining per sample type tested. The sections were pre-treated

using Heat mediated antigen retrieval using Bond™ Epitope

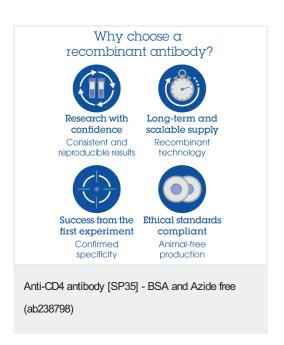
Retrieval Solution 2 (pH 9.0) for 10 minutes. The sections were

incubated with ab213215 for 30 mins at room temperature followed

by a ready to use Rabbit specific IHC polymer detection kit

HRP/DAB (ab209101). The immunostaining was performed on a

Leica Biosystems BOND® RX instrument.



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

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