# abcam

# Product datasheet

# Anti-CDA antibody [EPR20525] - BSA and Azide free ab227815



#### 5 Images

#### Overview

Product name Anti-CDA antibody [EPR20525] - BSA and Azide free

**Description** Rabbit monoclonal [EPR20525] to CDA - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: Flow Cyt (Intra), WB, IHC-P

Species reactivity Reacts with: Human

**Immunogen** Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

**Positive control** IHC-P: Human spleen tissue.

**General notes** ab227815 is the carrier-free version of <u>ab222515</u>.

Our <u>carrier-free</u> 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 cell-based 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.

Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **patents**.

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#### **Properties**

Form Liquid

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

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

**Purity** Protein A purified

Clonality Monoclonal
Clone number EPR20525

**Isotype** IgG

#### **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab227815 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
Flow Cyt (Intra)		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 16 kDa.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

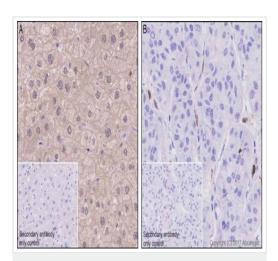
### **Target**

Relevance CDA (Cytidine deaminase) scavengers exogenous and endogenous cytidine and 2'-

deoxycytidine for UMP synthesis. Growth inhibition of granulocyte-macrophage colony foring cells by human cytidine deaminase requires the catalytic function of the protein. It is highly expressed in

granulocytes.

# **Images**



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CDA antibody

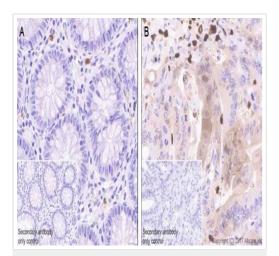
[EPR20525] - BSA and Azide free (ab227815)

Immunohistochemical analysis of paraffin-embedded human normal liver (panel A) and liver cancer (panel B) tissues labeling CDA with ab222515 at 1/2000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic and nuclear staining on normal human liver tissues, with only sporadic stromal cells showing positive staining in human liver cancer. The IHC signal on human liver cancer tissue was much lower than its corresponding normal tissue (PMID: 9849491). Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab222515).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CDA antibody

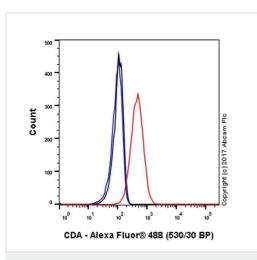
[EPR20525] - BSA and Azide free (ab227815)

Immunohistochemical analysis of paraffin-embedded human normal colon (panel A) and colon cancer (panel B) tissues labeling CDA with <a href="mailto:ab222515">ab222515</a> at 1/2000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Cytoplasmic and nuclear staining on sporadic stromal cells of human normal colon tissue, while adjacent cancer cells and some stromal cells show moderate positive staining. The IHC signal on human colon cancer tissue was higher than its corresponding normal tissue (PMID: 9849491). Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab222515).

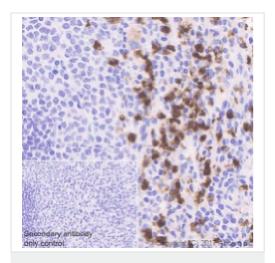
Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-CDA antibody [EPR20525] - BSA and Azide free (ab227815)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol permeabilized HeLa (human epithelial cell line from cervix adenocarcinoma) cell line labeling CDA with <a href="mailto:ab222515">ab222515</a> at 1/70 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730) (black) and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150077) at 1/2000 dilution was used as the secondary antibody.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab222515).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CDA antibody

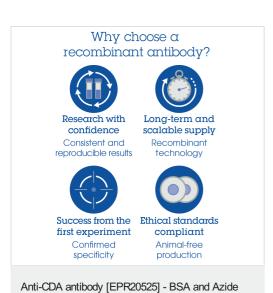
[EPR20525] - BSA and Azide free (ab227815)

Immunohistochemical analysis of paraffin-embedded human spleen tissue labeling CDA with <u>ab222515</u> at 1/2000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Nuclear and cytoplasmic staining on neutrophils of human spleen (PMID: 11069255). Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) Ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab222515).

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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#### Terms and conditions

free (ab227815)

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