

# **Product datasheet**

# Anti-CD11b antibody [EPR19387] - BSA and Azide free ab232427

Recombinant RabMAb

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Overview		
Product name	Anti-CD11b antibody [EPR19387] - BSA and Azide free	
Description	Rabbit monoclonal [EPR19387] to CD11b - BSA and Azide free	
Host species	Rabbit	
Tested applications	Suitable for: ICC/IF, IP, Flow Cyt, WB	
Species reactivity	Reacts with: Mouse, Human	
Immunogen	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.	
Positive control	ICC/IF: RAW 264.7 cells.	
General notes	ab232427 is the carrier-free version of <u>ab184308</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 <b>conjugation kits</b> 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.	
	<ul> <li>This product is a recombinant monoclonal antibody, which offers several advantages including:</li> <li>High batch-to-batch consistency and reproducibility</li> <li>Improved sensitivity and specificity</li> <li>Long-term security of supply</li> <li>Animal-free production</li> <li>For more information see here.</li> <li>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.</li> </ul>	

# 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	EPR19387		
lsotype	lgG		

# Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab232427 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
ICC/IF		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.
Flow Cyt		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 170 kDa (predicted molecular weight: 127 kDa).

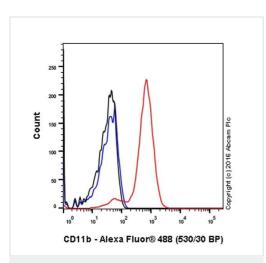
Target	
Function	Integrin alpha-M/beta-2 is implicated in various adhesive interactions of monocytes, macrophages and granulocytes as well as in mediating the uptake of complement-coated particles. It is identical with CR-3, the receptor for the iC3b fragment of the third complement component. It probably recognizes the R-G-D peptide in C3b. Integrin alpha-M/beta-2 is also a receptor for fibrinogen, factor X and ICAM1. It recognizes P1 and P2 peptides of fibrinogen gamma chain.
Tissue specificity	Predominantly expressed in monocytes and granulocytes.
Involvement in disease	Genetic variations in ITGAM has been associated with susceptibility to systemic lupus erythematosus type 6 (SLEB6) [MIM:609939]. Systemic lupus erythematosus (SLE) is a chronic, inflammatory and often febrile multisystemic disorder of connective tissue. It affects principally the skin, joints, kidneys and serosal membranes. It is thought to represent a failure of the regulatory mechanisms of the autoimmune system.
Sequence similarities	Belongs to the integrin alpha chain family. Contains 7 FG-GAP repeats. Contains 1 VWFA domain.
Domain	The integrin I-domain (insert) is a VWFA domain. Integrins with I-domains do not undergo

#### protease cleavage.

Membrane.

#### **Cellular localization**

#### Images



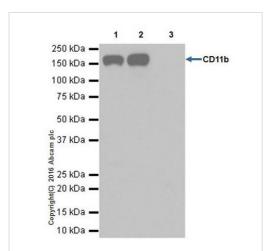
Flow Cytometry - Anti-CD11b antibody [EPR19387] -

BSA and Azide free (ab232427)

line transformed with Abelson murine leukemia virus) cells labeling CD11b with <u>ab184308</u> at 1/70 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A] -lsotype control (<u>ab172730</u>) (black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (blue). Goat anti Rabbit IgG (Alexa Fluor<sup>®</sup> 488) at 1/2000 dilution was used as the secondary antibody.

Flow cytometric analysis of RAW 264.7 (Mouse macrophage cell

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab184308**).



Immunoprecipitation - Anti-CD11b antibody [EPR19387] - BSA and Azide free (ab232427) CD11b was immunoprecipitated from 0.35 mg of RAW 264.7 (Mouse macrophage cell line transformed with Abelson murine leukemia virus) whole cell lysate with <u>ab184308</u> at 1/40 dilution. Western blot was performed from the immunoprecipitate using <u>ab184308</u> at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>), was used for detection at 1/10000 dilution.

Lane 1: RAW 264.7 whole cell lysate, 10µg (Input).

Lane 2: ab184308 IP in RAW 264.7 whole cell lysate.

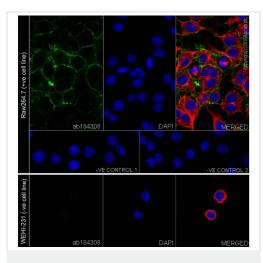
Lane 3: Rabbit lgG,monoclonal [EPR25A]- lsotype Control (**ab172730**) instead of **ab184308** in RAW 264.7 whole cell

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 8 seconds.

lysate.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab184308</u>).



Immunocytochemistry/ Immunofluorescence - Anti-CD11b antibody [EPR19387] - BSA and Azide free (ab232427) Immunofluorescent analysis of 100% methanol-fixed RAW 264.7 (Mouse macrophage cell line transformed with Abelson murine leukemia virus) and WEHI-231 (Mouse B cell lymphoma cell line) cells labeling CD11b with <u>ab184308</u> at 1/500 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor<sup>®</sup> 488) (<u>ab150077</u>) secondary antibody at 1/1,000 dilution (green).

Confocal image showing membrane staining on RAW 264.7 cells and no staining on WEHI-231 cells (negative cell line).

The nuclear counter stain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] -Loading Control (**ab7291**) at 1/1,000 dilution and Goat Anti-Mouse IgG H&L (Alexa Fluor<sup>®</sup> 594) preadsorbed (**ab150120**) at 1/1,000 dilution (red).

The negative controls are as follows:

-ve control 1: <u>ab184308</u> at 1/500 dilution followed by <u>ab150120</u> at 1/1,000 dilution.

-ve control 2: <u>ab7291</u> at 1/1000 dilution followed by <u>ab150077</u> at 1/1,000 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab184308**).



free (ab232427)

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