

# Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free ab247712

Recombinant RabMAb

6 Images

### Overview

<b>Product name</b>	Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free
<b>Description</b>	Rabbit monoclonal [EPR4116] to Glycophorin C/GPC - BSA and Azide free
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> IHC-P, IP, Flow Cyt (Intra), WB
<b>Species reactivity</b>	<b>Reacts with:</b> Human
<b>Immunogen</b>	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.
<b>Positive control</b>	Flow Cyt: K562 cell lysate; IP: TF-1 lysate.
<b>General notes</b>	<p>ab247712 is the carrier-free version of <a href="#">ab108925</a>.</p> <p>Our <b>carrier-free</b> 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.</p> <p>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.</p> <p>Use our <b>conjugation kits</b> for antibody conjugates that are ready-to-use in as little as 20 minutes with &lt;1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.</p> <p>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.</p> <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"><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></ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a>.</p> <p>Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with</p>

these species. Please contact us for more information.

## Properties

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<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C. Do Not Freeze.
<b>Storage buffer</b>	pH: 7.2 Constituent: PBS
<b>Carrier free</b>	Yes
<b>Purity</b>	Affinity purified
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	EPR4116
<b>Isotype</b>	IgG

## Applications

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**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab247712 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
<b>IHC-P</b>		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
<b>IP</b>		Use at an assay dependent concentration.
<b>Flow Cyt (Intra)</b>		Use at an assay dependent concentration.
<b>WB</b>		Use at an assay dependent concentration. Predicted molecular weight: 14 kDa.

## Target

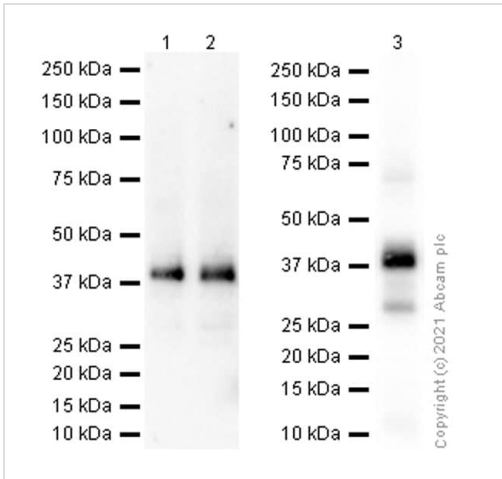
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<b>Function</b>	This protein is a minor sialoglycoprotein in human erythrocyte membranes. The blood group Gerbich antigens and receptors for Plasmodium falciparum merozoites are most likely located within the extracellular domain. Glycophorin-C plays an important role in regulating the stability of red cells.
<b>Tissue specificity</b>	Glycophorin-C is expressed in erythrocytes. Glycophorin-D is ubiquitous.
<b>Sequence similarities</b>	Belongs to the glycophorin-C family.
<b>Cellular localization</b>	Cell membrane. Linked to the membrane via band 4.1.

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## Images

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Western blot - Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free (ab247712)

**All lanes** : Anti-Glycophorin C/GPC antibody [EPR4116] (**ab108925**) at 1/1000 dilution (Purified)

**Lane 1** : TF-1 (Human Erythroleukemia erythroblast) whole cell lysate

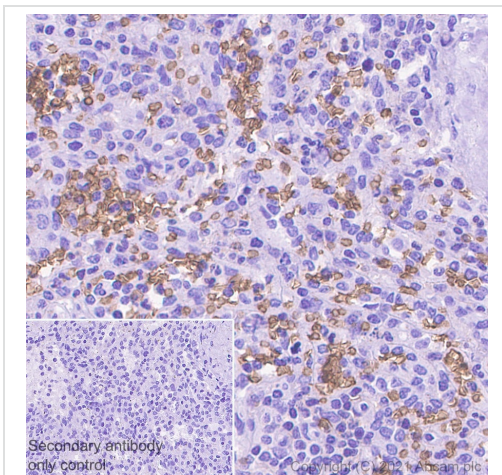
**Lane 2** : K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate

**Lane 3** : Human red blood lysate

### Secondary

**All lanes** : Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

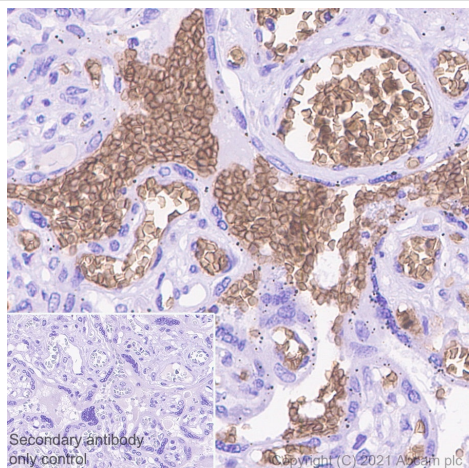
**Predicted band size:** 14 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free (ab247712)

This data was developed using **ab108925**, the same antibody clone in a different buffer formulation.

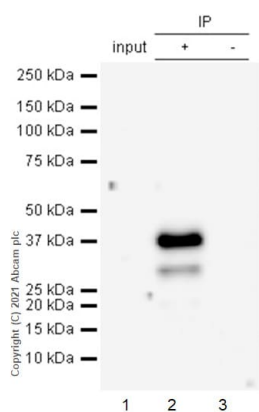
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human spleen tissue sections labeling Glycophorin C/GPC with Purified **ab108925** at 1:1500 dilution (0.06 µg/ml). Heat mediated antigen retrieval was performed using Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0) . Tissue was counterstained with Hematoxylin. Rabbit specific IHC polymer detection kit HRP/DAB (**ab209101**) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free (ab247712)

This data was developed using **ab108925**, the same antibody clone in a different buffer formulation.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human placenta tissue sections labeling Glycophorin C/GPC with Purified **ab108925** at 1:1500 dilution (0.06 µg/ml). Heat mediated antigen retrieval was performed using Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0) . Tissue was counterstained with Hematoxylin. Rabbit specific IHC polymer detection kit HRP/DAB (**ab209101**) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control. M3



Immunoprecipitation - Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free (ab247712)

This data was developed using **ab108925**, the same antibody clone in a different buffer formulation.

Glycophorin C/GPC was immunoprecipitated from 0.35 mg TF-1 (Human Erythroleukemia erythroblast) whole cell lysate 10 µg with **ab108925** at 1/50 dilution (2µg). VeriBlot for IP Detection Reagent (HRP)(**ab131366**) was used at 1/5000 dilution.

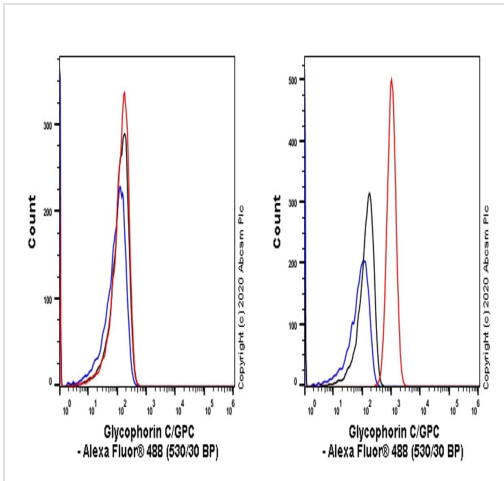
Lane 1: TF-1 (Human Erythroleukemia erythroblast) whole cell lysate 10 µg

Lane 2: abab108925 IP in TF-1 whole cell lysate

Lane 3: Rabbit monoclonal IgG (**ab172730**) instead of **ab108925** in TF-1 whole cell lysate

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

Lower band could be N-Deglycosylated GPC. (PMID: 16461900)



Flow Cytometry (Intracellular) - Anti-Glycophorin C/GPC antibody [EPR4116] - BSA and Azide free (ab247712)

This data was developed using **ab108925**, the same antibody clone in a different buffer formulation.

Flow Cytometry analysis of A549 (Human lung carcinoma epithelial cell, Left) / K-562 (Human chronic myelogenous leukemia lymphoblast, Right) cells labeling Glycophorin C/GPC with purified **ab108925** at 1/10000 dilution (0.1 µg/mL) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit IgG (Alexa Fluor® 488, **ab150077**) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal IgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).

Negative control: A549.

Why choose a recombinant antibody?

 <p><b>Research with confidence</b> Consistent and reproducible results</p>	 <p><b>Long-term and scalable supply</b> Recombinant technology</p>
 <p><b>Success from the first experiment</b> Confirmed specificity</p>	 <p><b>Ethical standards compliant</b> Animal-free production</p>

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

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