# abcam

# Product datasheet

# Anti-GATA2 antibody [EPR2822(2)] ab109241





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#### Overview

**Product name** Anti-GATA2 antibody [EPR2822(2)]

**Description** Rabbit monoclonal [EPR2822(2)] to GATA2

**Host species** Rabbit

**Tested applications** Suitable for: ChIP, WB, IP

Species reactivity Reacts with: Mouse, Rat, Human

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

Positive control WB: bEnd.3, PC-12, HEK-293, and K562 whole cell lysates. Mouse placenta lysate; IP: K-562

whole cell lysate.

**General notes** 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® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

#### **Properties**

**Form** Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

**Purity** Protein A purified

Clonality Monoclonal Clone number EPR2822(2)

ΙgG Isotype

### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab109241 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
ChIP		Use at an assay dependent concentration.
WB		1/1000. Predicted molecular weight: 51 kDa.  For unpurified use at 1/1000 - 1/10000.
IP		1/30. For unpurified use at 1/10 - 1/100.

#### **Target**

Function Transcriptional activator which regulates endothelin-1 gene expression in endothelial cells. Binds

to the consensus sequence 5'-AGATAG-3'.

Tissue specificity Endothelial cells.

**Sequence similarities**Contains 2 GATA-type zinc fingers.

Cellular localization Nucleus.

# **Images**



Immunoprecipitation - Anti-GATA2 antibody [EPR2822(2)] (ab109241)

GATA2 was immunoprecipitated from 0.35 mg K-562 (Human chronic myelogenous leukemia lymphoblast) cell lysate 10  $\mu$ g with ab109241 at 1/30 dilution (2 $\mu$ g in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab109241 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366) was used at 1/5000 dilution.

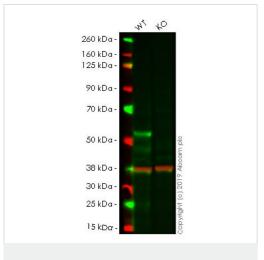
Lane 1: K-562 (Human chronic myelogenous leukemia lymphoblast) cell lysate 10  $\mu g$ 

Lane 2: ab109241 IP in K-562 cell lysate

Lane 3: Rabbit monoclonal lgG ( $\underline{ab172730}$ ) instead of ab109241 in K562 cell lysate

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

Exposure time: 8 seconds



Western blot - Anti-GATA2 antibody [EPR2822(2)] (ab109241)

All lanes: Anti-GATA2 antibody [EPR2822(2)] (ab109241) at 1 µg/ml

Lane 1: Wild-type HEK-293 whole cell lysate

**Lane 2**: GATA2 knockout HEK-293 (Human epithelial cell line from embryonic kidney) whole cell lysate

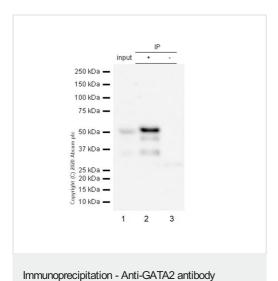
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 51 kDa

**Lanes 1 - 2:** Merged signal (red and green). Green - ab109241 observed at 50 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

ab109241 was shown to recognize GATA2 in wild-type HEK-293 cells as signal was lost at the expected MW in GATA2 knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and GATA2 knockout samples were subjected to SDS-PAGE. The membrane was blocked with 3% Milk. Ab109241 and <a href="mailto:ab8245">ab8245</a> (Mouse anti-GAPDH loading control) were incubated overnight at 4°C at 1 µg/ml and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preabsorbed <a href="mailto:ab216773">ab216773</a> and Goat anti-Mouse lgG H&L (IRDye® 680RD) preabsorbed <a href="mailto:ab216776">ab216776</a> secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



[EPR2822(2)] (ab109241)

Purified ab109241 at 1/30 dilution (2  $\mu$ g) immunoprecipitating GATA2 in K-562 whole cell lysate.

Lane 1 (input): K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate 10  $\mu g$ 

Lane 2 (+): ab109241 + K-562 whole cell lysate.

Lane 3 (-): Rabbit monoclonal IgG (ab172730) instead of ab109241 in K-562 whole cell lysate.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) (1/1000 dilution) was used for Western blotting.

Blocking Buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM/TBST.

Observed band size: 51 kDa



Western blot - Anti-GATA2 antibody [EPR2822(2)] (ab109241)

**All lanes :** Anti-GATA2 antibody [EPR2822(2)] (ab109241) at 1/1000 dilution (purified)

Lane 1: bEnd.3 (Mouse brain endothelioma) whole cell lysate

Lane 2: Mouse placenta lysate

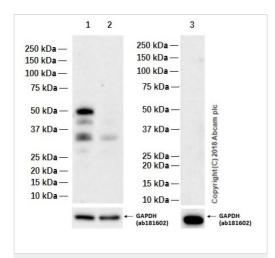
Lane 3 : PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysate

Lysates/proteins at 15 µg per lane.

#### **Secondary**

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

**Predicted band size:** 51 kDa **Observed band size:** 51 kDa



Western blot - Anti-GATA2 antibody [EPR2822(2)] (ab109241)

**All lanes :** Anti-GATA2 antibody [EPR2822(2)] (ab109241) at 1/1000 dilution

**Lane 1 :** K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate prepared using 1% SDS hot lysis method

Lane 2 : K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate prepared using RIPA lysis method Lane 3 : U-937 (Human histiocytic lymphoma monocyte) whole cell

lysate

Lysates/proteins at 20 µg per lane.

#### **Secondary**

**All lanes :** Goat Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/2000 dilution

Predicted band size: 51 kDa

Exposure time: 3 minutes

Blocking and diluting buffer: 5% NFDM/TBST.

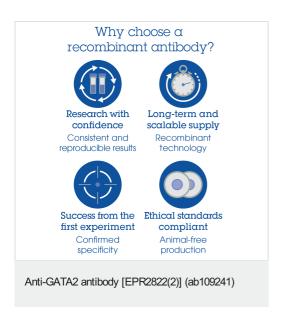
The different result in K-562 is due to the lysates preparation method.

For Lysate preparation protocol, please refer to the protocol book in the protocol section and/or <a href="https://example.com/here/downloadable.com/">here (downloadable copy)</a>.

The expression profile observed in U-937 is consistent with the

literature (PMID: 19212333).

Negative control: U-937 (PMID: 19212333)



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