# abcam

# Product datasheet

# Anti-SIKE1 antibody [EPR14692] ab183509





# 6 Images

#### Overview

**Product name** Anti-SIKE1 antibody [EPR14692]

**Description** Rabbit monoclonal [EPR14692] to SIKE1

**Host species** Rabbit

**Tested applications** Suitable for: IP, IHC-P, WB Species reactivity Reacts with: Mouse. Human

Predicted to work with: Rat

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

Positive control SH-SY5Y, HeLa and HEK-293T cell lysates; Human kidney tissue.

**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: 7.2

Preservative: 0.01% Sodium azide

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

**Purity** Protein A purified

Clonality Monoclonal Clone number EPR14692

Isotype lqG

#### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab183509 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
IP		1/30 - 1/50.
IHC-P		1/50. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/10000 - 1/50000. Detects a band of approximately 24 kDa (predicted molecular weight: 24 kDa).

#### **Target**

and TLR3-
and TLR

triggered IRF3. Inhibits TLR3-mediated activation of interferon-stimulated response elements (ISRE) and the IFN-beta promoter. May act by disrupting the interactions of IKBKE or TBK1 with

TICAM1/TRIF, IRF3 and DDX58/RIG-I. Does not inhibit NF-kappa-B activation pathways.

**Tissue specificity** Widely expressed. Expressed in brain, heart, skeletal muscle, colon, thymus, spleen, kidney, liver,

small intestine, placenta, lung and leukocytes. Present in all cell lines tested (at protein level).

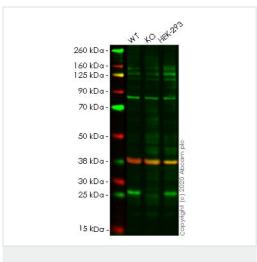
**Sequence similarities** Belongs to the SIKE family.

**Post-translational** Phosphorylated upon DNA damage, probably by ATM or ATR.

modifications

Cellular localization Cytoplasm.

## **Images**



Western blot - Anti-SIKE1 antibody [EPR14692] (ab183509)

All lanes: Anti-SIKE1 antibody [EPR14692] (ab183509) at

1/10000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: SIKE1 knockout HeLa cell lysate

Lane 3: HEK-293 cell lysate

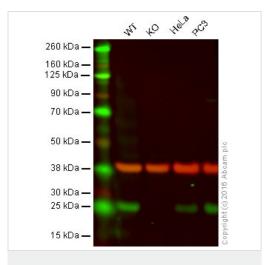
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 24 kDa

**Lanes 1-3:** Merged signal (red and green). Green - ab183509 observed at 25 kDa. Red - loading control **ab8245** observed at 37 kDa.

ab183509 Rabbit monoclonal [EPR14692] to SIKE1 was shown to specifically react with Suppressor of IKBKE 1 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265933 (knockout cell lysate ab258188) was used. Wild-type and Suppressor of IKBKE 1 knockout samples were subjected to SDS-PAGE. ab183509 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 10000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-SIKE1 antibody [EPR14692] (ab183509)

**All lanes :** Anti-SIKE1 antibody [EPR14692] (ab183509) at 1/10000 dilution

Lane 1: Wild-type HAP1 cell lysate

Lane 2: SIKE1 knockout HAP1 cell lysate

Lane 3 : HeLa cell lysate

Lane 4 : PC3 cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 24 kDa

**Lanes 1 - 4**: Merged signal (red and green). Green - ab183509 observed at 25 kDa. Red - loading control, **ab8245**, observed at 37 kDa.

ab183509 was shown to specifically react with SIKE1 when SIKE1 knockout samples were used. Wild-type and SIKE1 knockout samples were subjected to SDS-PAGE. ab183509 and <u>ab8245</u> (loading control to GAPDH) were both diluted 1/10 000 and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed <u>ab216773</u> and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed <u>ab216776</u> secondary antibodies at 1/10000 dilution for 1 hour at

KDa 1 2 3

250 —
150 —
100 —
75 —
50 —
37 —
25 —
20 —
15 —

Western blot - Anti-SIKE1 antibody [EPR14692] (ab183509)

room temperature before imaging.

**All lanes :** Anti-SIKE1 antibody [EPR14692] (ab183509) at 1/20000 dilution

Lane 1: SH-SY5Y cell lysate

Lane 2: Hela cell lysate

Lane 3: 293 cell lysate

Lysates/proteins at 20 µg per lane.

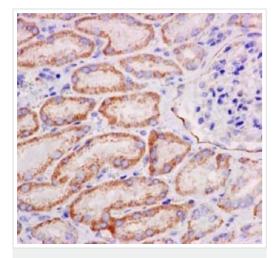
# **Secondary**

**All lanes :** Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 24 kDa

Additional bands at: 24 kDa. We are unsure as to the identity of

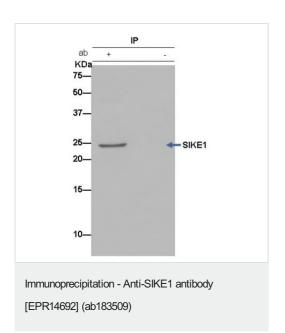
these extra bands.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SIKE1 antibody
[EPR14692] (ab183509)

Immunohistochemical analysis of paraffin embedded Human kidney tissue labeling SIKE1 with ab183509 at 1/50.

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.



Lysate from 293 cells was immunoprecipitated with ab183509 at a 1/50 dilution. For the subsequent blot, ab183509 used at a 1/10000 dilution with an anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at a 1/1500 dilution for the secondary. Blocking/ Dilution buffer: 5% NFDM/TBST. Lane 2: TBS instead of lysate.



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

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