

## Product datasheet

### Anti-SUZ12 antibody [EPR26230-82] ab307891

KO VALIDATED

Recombinant

RabMAb

13 Images

#### Overview

|                     |   |
|---------------------|---|
| Product name        | Anti-SUZ12 antibody [EPR26230-82]   |
| Description         | Rabbit monoclonal [EPR26230-82] to SUZ12  |
| Host species        | Rabbit  |
| Specificity         | Unsuitable for human FC-Intra.  |
| Tested applications | <b>Suitable for:</b> ICC/IF, Flow Cyt (Intra), IHC-P, WB<br><b>Unsuitable for:</b> ChIP or IP   |
| Species reactivity  | <b>Reacts with:</b> Mouse, Rat, Human   |
| Immunogen           | Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.  |
| Positive control    | WB: Wild-type HAP1 whole cell lysate. HeLa, 293T, NIH/3T3, U-87 MG, SH-SY5Y, NCCIT, Neuro-2a, C6, F9 and Caco-2 whole cell lysate. IHC-P: Human tonsil tissue. Human diffuse large B-cell lymphoma. Mouse and rat testis tissue. Mouse large B-cell lymphoma. ICC/IF: Wild-type HAP1 cells. HeLa and F9 cells. Flow Cyt (Intra): F9 cells.  |
| General notes       | <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> |

#### 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. Avoid freeze / thaw cycle. |
| Storage buffer       | pH: 7.2<br>Preservative: 0.01% Sodium azide<br>Constituents: 0.05% BSA, 40% Glycerol (glycerin, glycerine), 59% PBS     |
| Purity               | Protein A purified  |

|                     |             |
|---------------------|-------------|
| <b>Clonality</b>    | Monoclonal  |
| <b>Clone number</b> | EPR26230-82 |
| <b>Isotype</b>      | IgG         |

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab307891 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

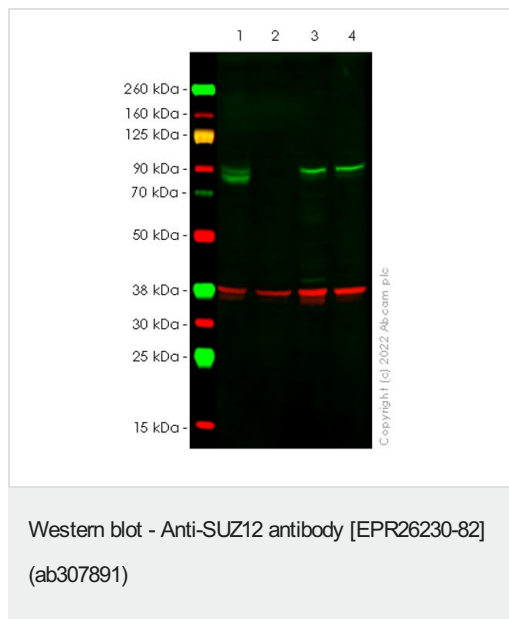
| Application      | Abreviews | Notes  |
|------------------|-----------|--|
| ICC/IF           |           | 1/50.  |
| Flow Cyt (Intra) |           | 1/50.  |
| IHC-P            |           | 1/200 - 1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. |
| WB               |           | 1/1000. Predicted molecular weight: 83 kDa.  |

**Application notes** Is unsuitable for ChIP or IP.

## Target

|                               |   |
|-------------------------------|---|
| <b>Function</b>               | Polycomb group (PcG) protein. Component of the PRC2/EED-EZH2 complex, which methylates 'Lys-9' (H3K9me) and 'Lys-27' (H3K27me) of histone H3, leading to transcriptional repression of the affected target gene. The PRC2/EED-EZH2 complex may also serve as a recruiting platform for DNA methyltransferases, thereby linking two epigenetic repression systems. Genes repressed by the PRC2/EED-EZH2 complex include HOXC8, HOXA9, MYT1 and CDKN2A. |
| <b>Tissue specificity</b>     | Overexpressed in breast and colon cancer.   |
| <b>Involvement in disease</b> | Note=A chromosomal aberration involving SUZ12 may be a cause of endometrial stromal tumors. Translocation t(7;17)(p15;q21) with JAZF1. The translocation generates the JAZF1-SUZ12 oncogene consisting of the N-terminus part of JAZF1 and the C-terminus part of SUZ12. It is frequently found in all cases of endometrial stromal tumors, except in endometrial stromal sarcomas, where it is rarer.  |
| <b>Sequence similarities</b>  | Belongs to the VEFS (VRN2-EMF2-FIS2-SU(Z)12) family.<br>Contains 1 C2H2-type zinc finger.   |
| <b>Developmental stage</b>    | Expressed at low levels in quiescent cells. Expression rises at the G1/S phase transition.  |
| <b>Cellular localization</b>  | Nucleus.  |

## Images



**All lanes :** Anti-SUZ12 antibody [EPR26230-82] (ab307891) at 1/1000 dilution

**Lane 1 :** Wild-type HAP1 (human chronic myelogenous leukemia near-haploid cell line) whole cell lysate

**Lane 2 :** SUZ12 knockout HAP1 whole cell lysate

**Lane 3 :** HeLa (human cervix adenocarcinoma epithelial cell) whole cell lysate

**Lane 4 :** Caco-2 (human colorectal adenocarcinoma epithelial cell) whole cell lysate

Lysates/proteins at 20 µg per lane.

### Secondary

**All lanes :** Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ([ab216773](#)) at 1/10000 dilution

**Predicted band size:** 83 kDa

Lysates at 20 µg per lane.

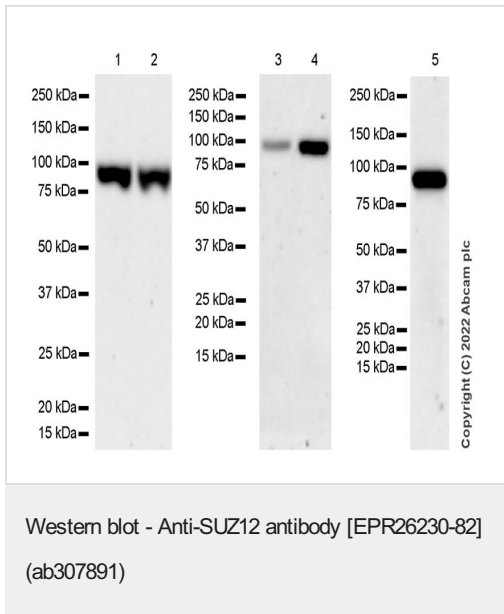
The samples were run on a Bis-Tris gel.

Performed under reducing conditions.

False colour image of Western blot: Anti-SUZ12 antibody (ab307891) staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody 6C5 ([ab8245](#)) loading control staining at 1/20000 dilution, shown in red.

In Western blot, ab307891 was shown to bind specifically to SUZ12. A band was observed at 95 kDa in wild-type HAP1 cell lysates with no signal observed at this size in SUZ12 knockout cell line. To generate this image, wild-type and SUZ12 knockout HAP1 cell lysates were analyzed. First, samples were run on an SDS-PAGE gel then transferred onto an immobilon-FL PVDF membrane. Membranes were blocked in Intercept® (TBS) Blocking Buffer diluted with an equal volume of 0.1% TBS before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L

(IRDye® 800CW) preabsorbed (**ab216773**) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (**ab216776**) at 1/20000 dilution.



**All lanes :** Anti-SUZ12 antibody [EPR26230-82] (ab307891) at 1/1000 dilution

**Lane 1 :** 293T (human embryonic kidney epithelial cell) whole cell lysate

**Lane 2 :** NIH/3T3 (mouse embryonic fibroblast), whole cell lysate

**Lane 3 :** U-87 MG (human glioblastoma-astrocytoma epithelial cell) whole cell lysate

**Lane 4 :** SH-SY5Y (human neuroblastoma epithelial cell) whole cell lysate

**Lane 5 :** NCCIT (human pluripotent embryonic carcinoma epithelial cell) whole cell lysate

Lysates/proteins at 20 µg per lane.

### Secondary

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

**Predicted band size:** 83 kDa

**Observed band size:** 95 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

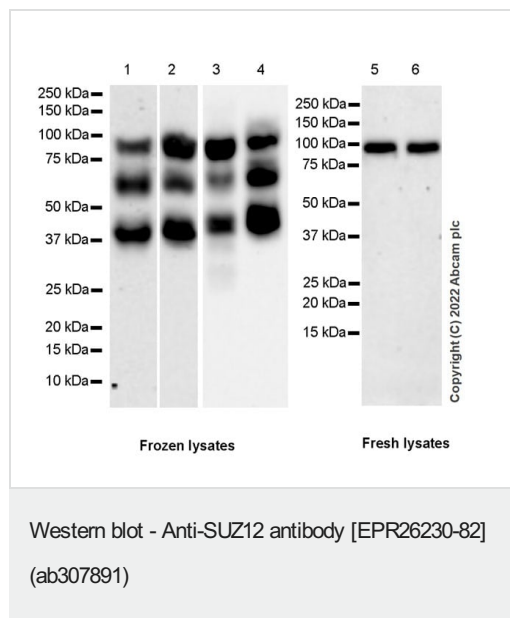
Exposure times:

Lane1 and 2: 81 seconds

Lane 3-5: 3 minutes

Lysates were freshly made and used immediately to minimize protein degradation.

The blot of lane 3-5 were developed using a high sensitivity ECL substrate.



**All lanes :** Anti-SUZ12 antibody [EPR26230-82] (ab307891) at 1/1000 dilution

**Lane 1 :** HeLa (human cervical adenocarcinoma epithelial cell) whole cell lysate

**Lane 2 :** Neuro-2a (mouse neuroblastoma neuroblast) whole cell lysate

**Lane 3 :** C6 (rat glial tumor glial cell) whole cell lysate

**Lane 4 :** F9 (mouse embryonal carcinoma epithelial cell) whole cell lysate

**Lane 5 :** C6 whole cell lysate

**Lane 6 :** Neuro-2a whole cell lysate

Lysates/proteins at 20 µg per lane.

## Secondary

**All lanes :** Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/20000 dilution

**Predicted band size:** 83 kDa

**Observed band size:** 95 kDa

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure times:

Lane 1 and 2: 3 minutes

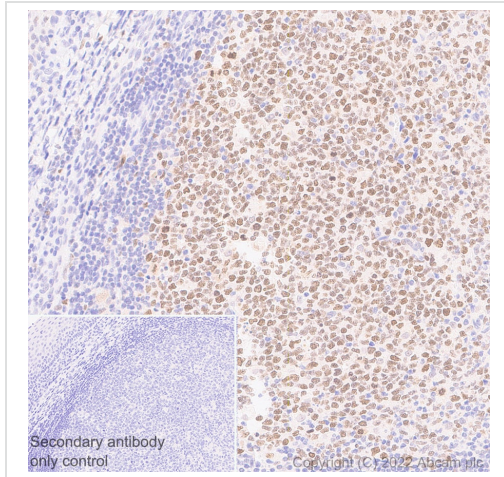
Lane 3 and 4: 114 seconds

Lane 5-6: 3 minutes

Lysates of lane 5 and 6 were freshly made and used immediately to minimize protein degradation.

The blot of lane 5-6 were developed using a high sensitivity ECL substrate.

The lower bands in lane 1-4 may be caused by degradation.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

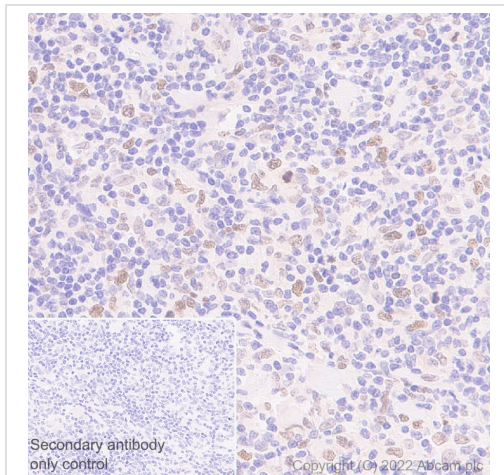
Immunohistochemical analysis of paraffin-embedded human tonsil tissue labeling SUZ12 with ab307891 at 1/200 dilution (2.465 µg/ml) followed by ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Nuclear staining in human tonsil germinal center (PMID: 20558579).

The section was incubated with ab307891 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

Immunohistochemical analysis of paraffin-embedded human diffuse large B-cell lymphoma tissue labeling SUZ12 with ab307891 at 1/200 dilution (2.465 µg/ml) followed by ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

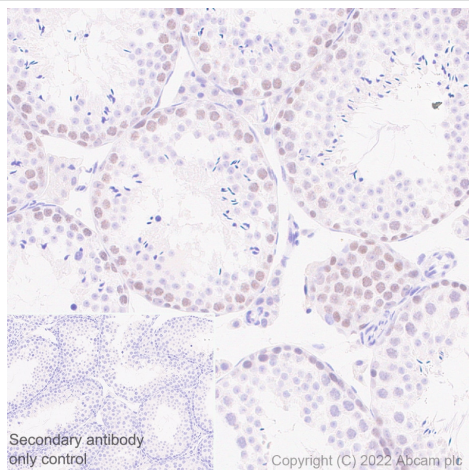
Nuclear staining in human diffuse large B-cell lymphoma (PMID: 20558579).

The section was incubated with ab307891 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins.





Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

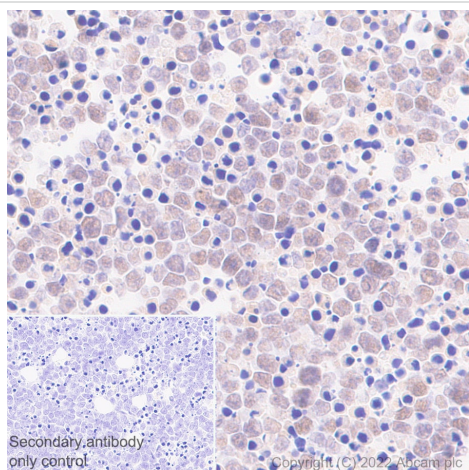
Immunohistochemical analysis of paraffin-embedded mouse testis tissue labeling SUZ12 with ab307891 at 1/1000 dilution (0.493 ug/ml) followed by ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Nuclear staining in mouse testis.

The section was incubated with ab307891 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

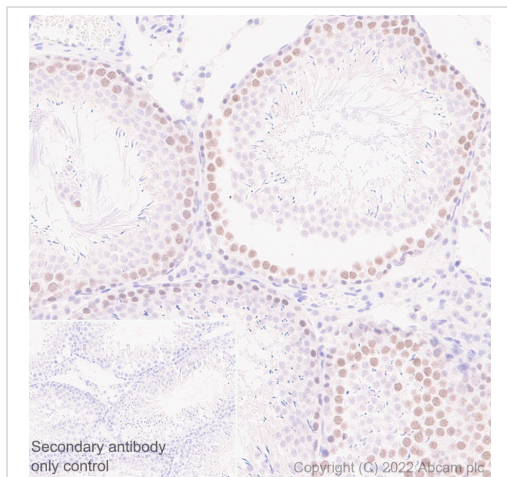
Immunohistochemical analysis of paraffin-embedded mouse large B-cell lymphoma tissue labeling SUZ12 with ab307891 at 1/1000 dilution (0.493 ug/ml) followed by ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Nuclear staining in mouse large B-cell lymphoma.

The section was incubated with ab307891 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

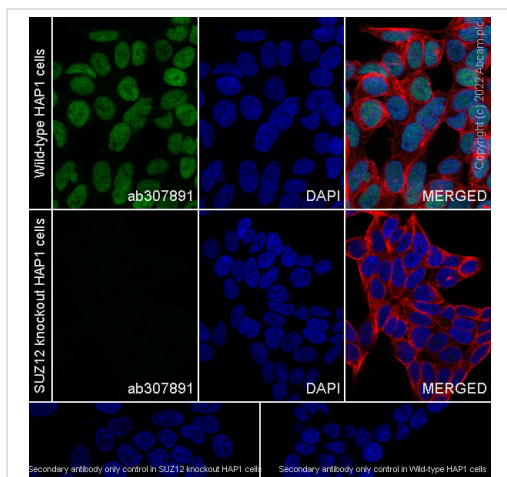
Immunohistochemical analysis of paraffin-embedded rat testis tissue labeling SUZ12 with ab307891 at 1/1000 dilution (0.493 ug/ml) followed by ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Nuclear staining in rat testis.

The section was incubated with ab307891 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond® Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins.



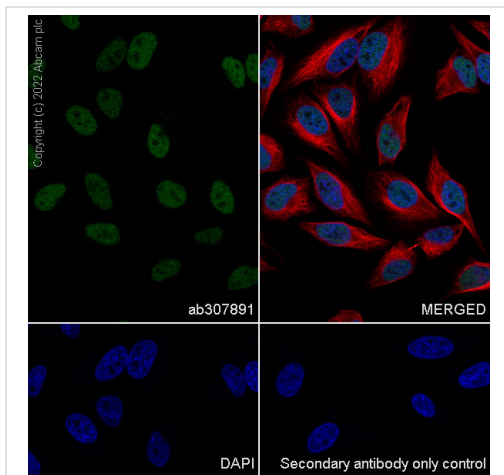
Immunocytochemistry/ Immunofluorescence - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized SUZ12 KO HAP1 (human chronic myelogenous leukemia near-haploid cell) cells labeling SUZ12 with ab307891 at 1/50 dilution (9.86 ug/ml) followed by **ab150081** Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 dilution (2 ug/mL) (Green).

Confocal image showing nuclear staining in parental HAP1 cells. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). **ab195889** Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 dilution (2.5 ug/ml) (Red). Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is **ab150081** Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (2 ug/mL).





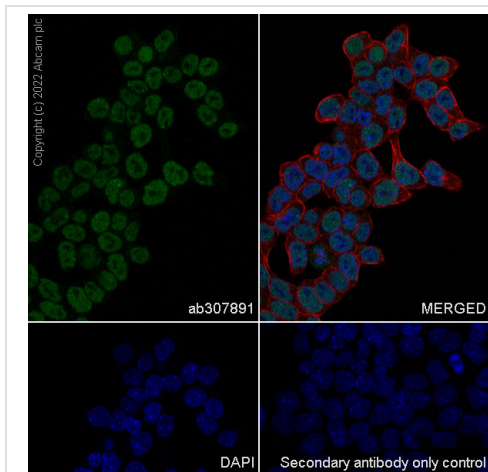
Immunocytochemistry/ Immunofluorescence - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (human cervix adenocarcinoma epithelial cell) cells labeling SUZ12 with ab307891 at 1/50 dilution (9.86 ug/ml) followed by **ab150081** Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 dilution (2 ug/mL) (Green).

Confocal image showing nuclear staining in HeLa cell line.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). **ab195889** Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 dilution (2.5 ug/ml) (Red). Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is **ab150081** Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (2 ug/mL).



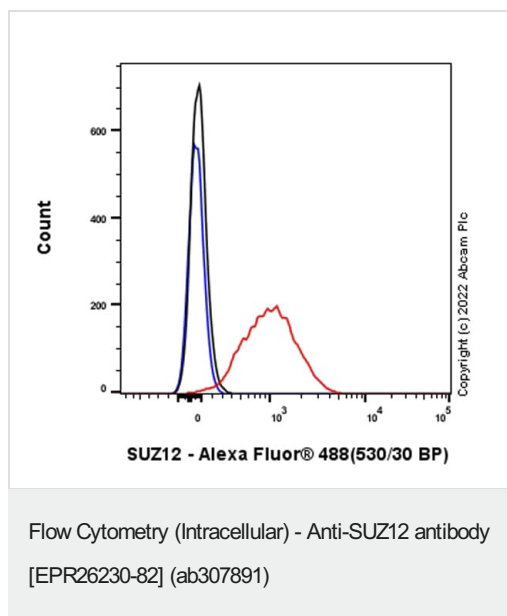
Immunocytochemistry/ Immunofluorescence - Anti-SUZ12 antibody [EPR26230-82] (ab307891)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized F9 (mouse embryonal carcinoma epithelial cell) cells labeling SUZ12 with ab307891 at 1/50 dilution (9.86 ug/ml) followed by **ab150081** Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 dilution (2 ug/mL) (Green).

Confocal image showing nuclear staining in F9 cell line.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). **ab195889** Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 dilution (2.5 ug/ml) (Red). Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is **ab150081** Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (2 ug/mL).



Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized F9 (mouse embryonal carcinoma epithelial cell) cells labeling SUZ12 with ab307891 at 1/50 dilution (1 ug) (Red) compared with a Rabbit monoclonal IgG (**ab172730**) (Black) isotype control and an unlabeled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor® 488, **ab150081**) at 1/5000 dilution was used as the secondary antibody.

Why choose a recombinant antibody?

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

Anti-SUZ12 antibody [EPR26230-82] (ab307891)

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

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- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

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