abcam

Product datasheet

Anti-Sumo 1 antibody [EP298] ab133352





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Overview

Product name Anti-Sumo 1 antibody [EP298]

Description Rabbit monoclonal [EP298] to Sumo 1

Host species Rabbit

Tested applications Suitable for: IHC-P, WB

Unsuitable for: Flow Cyt

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide within Human Sumo 1 aa 1-100. The exact sequence is proprietary.

Positive control WB- HeLa whole cell lysate (ab150035), MCF-7 whole cell lysate, A549 whole cell lysate,

> NIH/3T3 whole cell lysate, C6 whole cell lysate, HL-60 cell lysate and A431 cell lysate. IHC-Human papillary cell thyroid gland carcinoma tissue, Human Breast carcinoma tissue, Human Glioma tissue, Human Urinary bladder transitional carcinoma tissue and Human tonsil 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.

Stable for 12 months at -20°C.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 0.21% BSA, 59% PBS

Purity Protein A purified

Clonality Monoclonal

Clone number EP298

Applications

The Abpromise guarantee

Our Abpromise quarantee covers the use of ab133352 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
IHC-P	*****(1)	1/100 - 1/250. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		1/1000 - 1/10000. Predicted molecular weight: 11.5 kDa.

Application notes

Is unsuitable for Flow Cyt.

Target

Function

Ubiquitin-like protein that can be covalently attached to proteins as a monomer or a lysine-linked polymer. Covalent attachment via an isopeptide bond to its substrates requires prior activation by the E1 complex SAE1-SAE2 and linkage to the E2 enzyme UBE2I, and can be promoted by E3 ligases such as PIAS1-4, RANBP2 or CBX4. This post-translational modification on lysine residues of proteins plays a crucial role in a number of cellular processes such as nuclear transport, DNA replication and repair, mitosis and signal transduction. Involved for instance in targeting RANGAP1 to the nuclear pore complex protein RANBP2. Polymeric SUMO1 chains are also susceptible to polyubiquitination which functions as a signal for proteasomal degradation of modified proteins. May also regulate a network of genes involved in palate development.

Involvement in disease

Defects in SUMO1 are the cause of non-syndromic orofacial cleft type 10 (OFC10) [MIM:613705]; also called non-syndromic cleft lip with or without cleft palate 10. OFC10 is a birth defect consisting of cleft lips with or without cleft palate. Cleft lips are associated with cleft palate in two-third of cases. A cleft lip can occur on one or both sides and range in severity from a simple notch in the upper lip to a complete opening in the lip extending into the floor of the nostril and involving the upper gum. Note=A chromosomal aberation involving SUMO1 is the cause of OFC10. Translocation t(2;8)(q33.1;q24.3). The breakpoint occurred in the SUMO1 gene and resulted in haploinsufficiency confirmed by protein assays.

Sequence similarities

Belongs to the ubiquitin family. SUMO subfamily.

Contains 1 ubiquitin-like domain.

Post-translational modifications

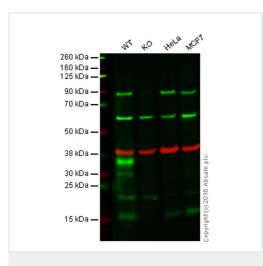
Cleavage of precursor form by SENP1 or SENP2 is necessary for function.

Polymeric SUMO1 chains undergo polyubiquitination by RNF4.

Cellular localization

Nucleus membrane. Nucleus speckle. Cytoplasm. Recruited by BCL11A into the nuclear body.

Images



Western blot - Anti-Sumo 1 antibody [EP298] (ab133352)



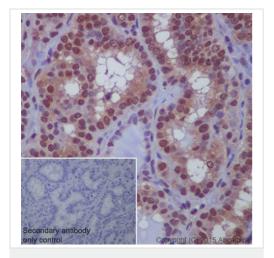
Lane 2: Sumo 1 knockout HAP1 cell lysate (20 µg)

Lane 3: HeLa cell lysate (20 µg)

Lane 4: MCF-7 cell lysate (20 µg)

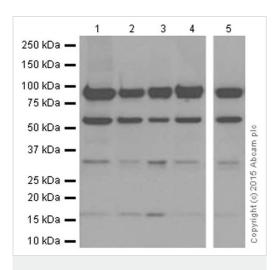
Lanes 1 - 4: Merged signal (red and green). Green - ab133352 observed at 16 kDa. Red - loading control, **ab8245**, observed at 37 kDa.

ab133352 was shown to recognize Sumo 1 when Sumo 1 knockout samples were used, along with additional cross-reactive bands. Wild-type and Sumo 1 knockout samples were subjected to SDS-PAGE. ab133352 and <u>ab8245</u> (loading control to GAPDH) were diluted 1/000 and 1/10000 respectively 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 room temperature before imaging.

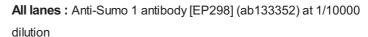


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sumo 1 antibody
[EP298] (ab133352)

Immunohistochemical staining of paraffin-embedded human thyroid carcinoma sections labelling Sumo 1 with purified ab133352 at dilution of 1:100. The secondary antibody used was ab97051; a goat anti-rabbit IgG H&L (HRP) at dilution of 1/500. The sample was counter-stained with hematoxylin. Antigen retrieval was performed using EDTA Buffer; pH 9.0. PBS was used instead of the primary antibody as the negative control and is shown in the inset.



Western blot - Anti-Sumo 1 antibody [EP298] (ab133352)



Lane 1: HeLa (human cervix adenocarcinoma) whole cell lysate

Lane 2: MCF-7 (human breast carcinoma) whole cell lysate

Lane 3: A549 (human lung carcinoma) whole cell lysate

Lane 4: NIH/3T3 (mouse embryo) whole cell lysate

Lane 5: C6 (rat glioma) whole cell lysate

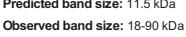
Lysates/proteins at 20 µg per lane.

Secondary

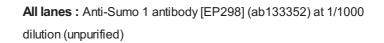
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/10000

dilution

Predicted band size: 11.5 kDa



Blocking and diluting buffer 5% NFDM/TBST



Lane 1: HeLa cell lysate Lane 2: HL60 cell lysate Lane 3: A431 cell lysate

Lane 4: NIH3T3 cell lysate

Lysates/proteins at 10 µg per lane.

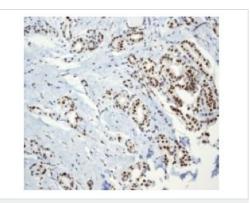
KDa 1 2 3 25 20. 15-10

Western blot - Anti-Sumo 1 antibody [EP298] (ab133352)

Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

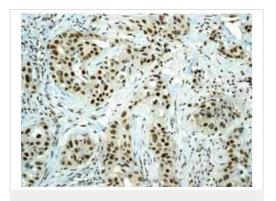
Predicted band size: 11.5 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sumo 1 antibody
[EP298] (ab133352)

Immunohistochemical analysis of paraffin-embedded Human papillary cell thyroid gland carcinoma tissue labelling Sumo 1 with un-purified ab133352 at 1/100 dilution.

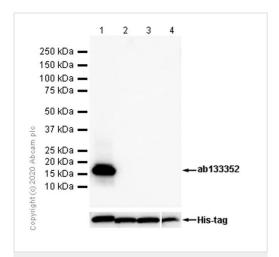
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sumo 1 antibody
[EP298] (ab1333352)

Immunohistochemical analysis of paraffin embedded Human Breast carcinoma tissue labelling Sumo 1 with un-purified ab133352 at dilution of 1/100-1/250.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Western blot - Anti-Sumo 1 antibody [EP298] (ab133352)

All lanes : Anti-Sumo 1 antibody [EP298] (ab133352) at 1/1000 dilution

Lane 1 : Recombinant Human Sumo 1 protein (ab140417) at 0.01

Lane 2 : Recombinant Human Sumo 2 protein (ab140420) at 0.01 µg

Lane 3: Recombinant Human Sumo 3 protein (ab140414) at 0.01

Lane 4 : Recombinant Human Sumo 4 protein (<u>ab157025</u>) at 0.1 μg

Secondary

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

Developed using the ECL technique.

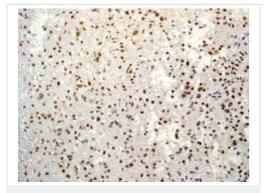
Predicted band size: 11.5 kDa **Observed band size:** 16 kDa

Exposure time: 10 seconds

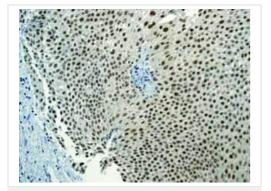
Blocking buffer: 5% NFDM/TBST

Immunohistochemical analysis of paraffin embedded Human Glioma tissue labelling Sumo 1 with un-purified ab133352 at dilution of 1/100-1/250.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



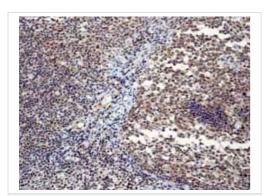
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sumo 1 antibody
[EP298] (ab133352)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sumo 1 antibody
[EP298] (ab133352)

Immunohistochemical analysis of paraffin embedded Human
Urinary bladder transitional carcinoma tissue labelling Sumo 1 with
un-purified ab133352 at dilution of 1/100-1/250.

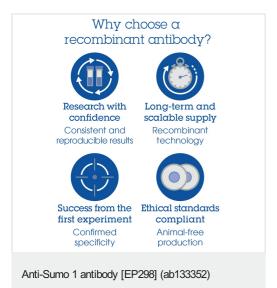
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Sumo 1 antibody
[EP298] (ab133352)

Immunohistochemical analysis of paraffin embedded normal Human tonsil tissue labelling Sumo 1 with un-purified ab133352 at dilution of 1/100-1/250.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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