

#### **SIRT7 Antibody**

Purified Rabbit Polyclonal Antibody Catalog # ABV11620

### **Specification**

## **SIRT7 Antibody - Product Information**

Application
Primary Accession
Reactivity
Host
Clonality
Isotype
Calculated MW

WB
O9NRC8
Human
Rabbit
Polyclonal
Rabbit IgG
44898

# **SIRT7 Antibody - Additional Information**

**Gene ID 51547** 

**Other Names** 

SIR2; Sirtuin 7; Silent Information Regulator 7

Target/Specificity

SIRT7

#### **Formulation**

50 μg of antibody in 100 μl PBS containing 0.2% gelatin and 0.05% sodium azide

#### Handling

The antibody solution should be gently mixed before use.

## **Background Descriptions**

### **Precautions**

SIRT7 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

# **SIRT7 Antibody - Protein Information**

Name SIRT7 {ECO:0000303|PubMed:22722849, ECO:0000312|HGNC:HGNC:14935}

#### **Function**

NAD-dependent protein-lysine deacylase that can act both as a deacetylase or deacylase (desuccinylase, depropionylase, deglutarylase and dedecanoylase), depending on the context (PubMed:<a href="http://www.uniprot.org/citations/22722849" target="\_blank">22722849</a>, PubMed:<a href="http://www.uniprot.org/citations/26907567" target="\_blank">26907567</a>, PubMed:<a href="http://www.uniprot.org/citations/30653310" target="\_blank">30653310</a>, PubMed:<a href="http://www.uniprot.org/citations/31542297" target="\_blank">31542297</a>, PubMed:<a href="http://www.uniprot.org/citations/35939806" target="\_blank">35939806</a>).



Specifically mediates deacetylation of histone H3 at 'Lys-18' (H3K18Ac) (PubMed:<a href="http://www.uniprot.org/citations/22722849" target=" blank">22722849</a>, PubMed:<a href="http://www.uniprot.org/citations/30420520" target="blank">30420520</a>, PubMed:<a href="http://www.uniprot.org/citations/35939806" target="\_blank">35939806</a>). In contrast to other histone deacetylases, displays strong preference for a specific histone mark, H3K18Ac, directly linked to control of gene expression (PubMed:<a href="http://www.uniprot.org/citations/22722849" target=" blank">22722849</a>, PubMed:<a href="http://www.uniprot.org/citations/30653310" target=" blank">30653310</a>). H3K18Ac is mainly present around the transcription start site of genes and has been linked to activation of nuclear hormone receptors; SIRT7 thereby acts as a transcription repressor (PubMed: <a href="http://www.uniprot.org/citations/22722849" target=" blank">22722849</a>). Moreover, H3K18 hypoacetylation has been reported as a marker of malignancy in various cancers and seems to maintain the transformed phenotype of cancer cells (PubMed: <a href="http://www.uniprot.org/citations/22722849" target=" blank">22722849</a>). Also able to mediate deacetylation of histone H3 at 'Lys-36' (H3K36Ac) in the context of nucleosomes (PubMed:<a href="http://www.uniprot.org/citations/30653310" target=" blank">30653310</a>). Also mediates deacetylation of non-histone proteins, such as ATM, CDK9, DDX21, DDB1, FBL, FKBP5/FKBP51, GABPB1, RAN, RRP9/U3-55K and POLR1E/PAF53 (PubMed: <a href="http://www.uniprot.org/citations/24207024" target=" blank">24207024</a>, PubMed:<a href="http://www.uniprot.org/citations/26867678" target=" blank">26867678</a>, PubMed:<a href="http://www.uniprot.org/citations/28147277" target="blank">28147277</a>, PubMed:<a href="http://www.uniprot.org/citations/28886238" target="blank">28886238</a>, PubMed:<a href="http://www.uniprot.org/citations/28426094" target="blank">28426094</a>, PubMed:<a href="http://www.uniprot.org/citations/30540930" target="blank">30540930</a>, PubMed:<a href="http://www.uniprot.org/citations/31075303" target="\_blank">31075303</a>, PubMed:<a href="http://www.uniprot.org/citations/30944854" target="blank">30944854</a>, PubMed:<a href="http://www.uniprot.org/citations/28790157" target="blank">28790157</a>). Enriched in nucleolus where it stimulates transcription activity of the RNA polymerase I complex (PubMed: <a href="http://www.uniprot.org/citations/16618798" target=" blank">16618798</a>, PubMed:<a href="http://www.uniprot.org/citations/19174463" target="\_blank">19174463</a>, PubMed:<a href="http://www.uniprot.org/citations/24207024" target="blank">24207024</a>). Acts by mediating the deacetylation of the RNA polymerase I subunit POLR1E/PAF53, thereby promoting the association of RNA polymerase I with the rDNA promoter region and coding region (PubMed: <a href="http://www.uniprot.org/citations/16618798" target=" blank">16618798</a>, PubMed:<a href="http://www.uniprot.org/citations/19174463" target="blank">19174463</a>, PubMed:<a href="http://www.uniprot.org/citations/24207024" target="blank">24207024</a>). In response to metabolic stress, SIRT7 is released from nucleoli leading to hyperacetylation of POLR1E/PAF53 and decreased RNA polymerase I transcription (PubMed:<a href="http://www.uniprot.org/citations/24207024" target="\_blank">24207024</a>). Required to restore the transcription of ribosomal RNA (rRNA) at the exit from mitosis (PubMed: <a href="http://www.uniprot.org/citations/19174463" target=" blank">19174463</a>). Promotes pre-ribosomal RNA (pre-rRNA) cleavage at the 5'-terminal processing site by mediating deacetylation of RRP9/U3-55K, a core subunit of the U3 snoRNP complex (PubMed:<a href="http://www.uniprot.org/citations/26867678" target=" blank">26867678</a>). Mediates 'Lys-37' deacetylation of Ran, thereby regulating the nuclear export of NF-kappa-B subunit RELA/p65 (PubMed: <a href="http://www.uniprot.org/citations/31075303" target=" blank">31075303</a>). Acts as a regulator of DNA damage repair by mediating deacetylation of ATM during the late stages of DNA damage response, promoting ATM dephosphorylation and deactivation (PubMed:<a href="http://www.uniprot.org/citations/30944854" target=" blank">30944854</a>). Suppresses the activity of the DCX (DDB1-CUL4-X-box) E3 ubiquitin-protein ligase complexes by mediating deacetylation of DDB1, which prevents the interaction between DDB1 and CUL4 (CUL4A or CUL4B) (PubMed:<a href="http://www.uniprot.org/citations/28886238" target=" blank">28886238</a>). Activates RNA polymerase II transcription by mediating deacetylation of CDK9, thereby promoting 'Ser-2' phosphorylation of the C-terminal domain (CTD) of RNA polymerase II (PubMed:<a href="http://www.uniprot.org/citations/28426094" target=" blank">28426094</a>). Deacetylates FBL, promoting histone- glutamine methyltransferase activity of FBL (PubMed: <a



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href="http://www.uniprot.org/citations/30540930" target=" blank">30540930</a>). Acts as a regulator of mitochondrial function by catalyzing deacetylation of GABPB1 (By similarity). Regulates Akt/AKT1 activity by mediating deacetylation of FKBP5/FKBP51 (PubMed:<a href="http://www.uniprot.org/citations/28147277" target="\_blank">28147277</a>). Required to prevent R- loop-associated DNA damage and transcription-associated genomic instability by mediating deacetylation and subsequent activation of DDX21, thereby overcoming R-loop-mediated stalling of RNA polymerases (PubMed:<a href="http://www.uniprot.org/citations/28790157" target=" blank">28790157</a>). In addition to protein deacetylase activity, also acts as a protein-lysine deacylase (PubMed:<a href="http://www.uniprot.org/citations/27436229" target="\_blank">27436229</a>, PubMed:<a href="http://www.uniprot.org/citations/27997115" target="\_blank">27436229</a>, PubMed:<a href="http://www.uniprot.org/citations/31542297" target="blank">31542297</a>). Acts as a protein depropionylase by mediating depropionylation of Osterix (SP7), thereby regulating bone formation by osteoblasts (By similarity). Acts as a histone deglutarylase by mediating deglutarylation of histone H4 on 'Lys-91' (H4K91glu); a mark that destabilizes nucleosomes by promoting dissociation of the H2A-H2B dimers from nucleosomes (PubMed: <a href="http://www.uniprot.org/citations/31542297" target=" blank">31542297</a>). Acts as a histone desuccinylase: in response to DNA damage, recruited to DNA double- strand breaks (DSBs) and catalyzes desuccinylation of histone H3 on 'Lys-122' (H3K122succ), thereby promoting chromatin condensation and DSB repair (PubMed:<a href="http://www.uniprot.org/citations/27436229" target=" blank">27436229</a>). Also promotes DSB repair by promoting H3K18Ac deacetylation, regulating non-homologous end joining (NHEJ) (By similarity). Along with its role in DNA repair, required for chromosome synapsis during prophase I of female meiosis by catalyzing H3K18Ac deacetylation (By similarity). Involved in

their association with the nuclear lamina (PubMed:<a href="http://www.uniprot.org/citations/31226208" target=" blank">31226208</a>). Required to stabilize ribosomal DNA (rDNA) heterochromatin and prevent cellular senescence induced by rDNA instability (PubMed: <a href="http://www.uniprot.org/citations/29728458" target=" blank">29728458</a>). Acts as a negative regulator of SIRT1 by preventing autodeacetylation of SIRT1, restricting SIRT1 deacetylase activity (By similarity).

transcriptional repression of LINE-1 retrotransposon via H3K18Ac deacetylation, and promotes

# **Cellular Location**

Nucleus, nucleolus. Nucleus, nucleoplasm. Chromosome. Cytoplasm. Note=Mainly localizes in the nucleolus and nucleoplasm (PubMed:24207024, PubMed:28886238, PubMed:28790157, PubMed:31075303). Associated with rDNA promoter and transcribed region (PubMed:16079181, PubMed:19174463). Associated with nucleolar organizer regions during mitosis (PubMed:16079181, PubMed:19174463). In response to stress, released from nucleolus to nucleoplasm (PubMed:24207024) Associated with chromatin (PubMed:22722849). In response to DNA damage, recruited to DNA double-strand breaks (DSBs) sites (PubMed:27436229) (Probable). Located close to the nuclear membrane when in the cytoplasm (PubMed:11953824).

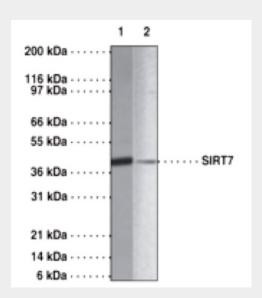
### SIRT7 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

### SIRT7 Antibody - Images





WB using SIRT7 pAb. Lane1. Human liver homogenate; Lane2. PBMC lysate.

# **SIRT7 Antibody - Background**

The sirtuins represent a distinct class of trichostatin A-insensitive lysyl-deacetylases (class III HDACs) and have been shown to catalyze a reaction that couples lysine deacetylation to the formation of nicotinamide and O-acetyl-ADP-ribose from NAD+ and the abstracted acetyl group. SIRT7 is a member of this family of proteins and is present in prokaryotes and eukaryotes. All SIR2-like proteins have a sirtuin core domain, which contains a series of sequence motifs conserved in organisms ranging from bacteria to humans. Bacterial, yeast, and mammalian sirtuins are able to metabolize NAD+ and several act as mono-ADP-ribosyltransferases. The enzymatic function of sirtuins is not yet completely understood but as mentioned above, recent reports of histone-activated SIR2-mediated NAD+ metabolism and NAD+-activated SIR2-mediated histone deacetylation s μggest a possible coupled reciprocal activation mechanism involving interactions of SIR2 with NAD+ and the N-ε-acetyl-lysine groups of acetylated histone.