

**SIRT7 Antibody**  
**Purified Rabbit Polyclonal Antibody**  
**Catalog # ABV11620****Specification**

---

**SIRT7 Antibody - Product Information**

Application	WB
Primary Accession	<a href="#">Q9NRC8</a>
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	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

[30540930](http://www.uniprot.org/citations/30540930)). 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:[28147277](http://www.uniprot.org/citations/28147277)). 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:[28790157](http://www.uniprot.org/citations/28790157)). In addition to protein deacetylase activity, also acts as a protein-lysine deacylase (PubMed:[27436229](http://www.uniprot.org/citations/27436229), PubMed:[27997115](http://www.uniprot.org/citations/27997115), PubMed:[31542297](http://www.uniprot.org/citations/31542297)). 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:[31542297](http://www.uniprot.org/citations/31542297)). 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:[27436229](http://www.uniprot.org/citations/27436229)). 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 transcriptional repression of LINE-1 retrotransposon via H3K18Ac deacetylation, and promotes their association with the nuclear lamina (PubMed:[31226208](http://www.uniprot.org/citations/31226208)). Required to stabilize ribosomal DNA (rDNA) heterochromatin and prevent cellular senescence induced by rDNA instability (PubMed:[29728458](http://www.uniprot.org/citations/29728458)). Acts as a negative regulator of SIRT1 by preventing autodeacetylation of SIRT1, restricting SIRT1 deacetylase activity (By similarity).

### Cellular Location

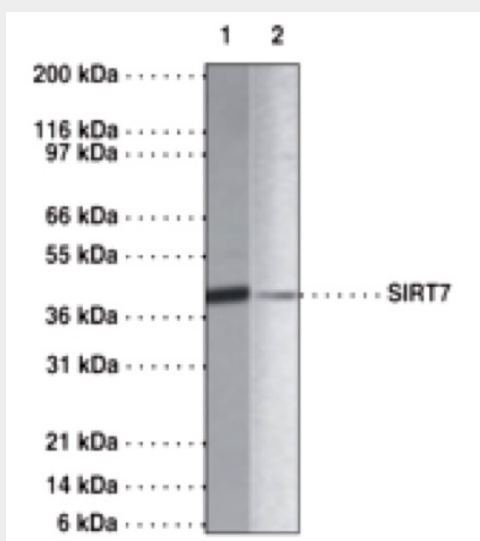
Nucleus, nucleolus. Nucleus, nucleoplasm. Chromosome. Cytoplasm. Note=Mainly localizes in the nucleolus and nucleoplasm (PubMed:24207024, PubMed:28886238, PubMed:28790157, PubMed:31075303). Associated with rDNA promoter and transcribed region (PubMed:16079181, PubMed:19174463). Associated with nucleolar 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](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [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<sup>+</sup> 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<sup>+</sup> 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<sup>+</sup> metabolism and NAD<sup>+</sup>-activated SIR2-mediated histone deacetylation suggest a possible coupled reciprocal activation mechanism involving interactions of SIR2 with NAD<sup>+</sup> and the N-ε-acetyl-lysine groups of acetylated histone.