

HDAC1 Antibody (aa466-482) Rabbit Polyclonal Antibody Catalog # ALS11219

Specification

HDAC1 Antibody (aa466-482) - Product Information

Application	IHC, WB, IF
Primary Accession	<u>013547</u>
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Calculated MW	55kDa KDa

HDAC1 Antibody (aa466-482) - Additional Information

Gene ID 3065

Other Names Histone deacetylase 1, HD1, 3.5.1.98, HDAC1, RPD3L1

Target/Specificity Amino acids 466-482 of Human HDAC-1

Reconstitution & Storage +4°C or -20°C, Avoid repeated freezing and thawing.

Precautions HDAC1 Antibody (aa466-482) is for research use only and not for use in diagnostic or therapeutic procedures.

HDAC1 Antibody (aa466-482) - Protein Information

Name HDAC1 {ECO:0000303|PubMed:10846170, ECO:0000312|HGNC:HGNC:4852}

Function

Histone deacetylase that catalyzes the deacetylation of lysine residues on the N-terminal part of the core histones (H2A, H2B, H3 and H4) (PubMed:16762839, PubMed:17704056, PubMed:28497810). Histone deacetylation gives a tag for epigenetic repression and plays an important role in transcriptional regulation, cell cycle progression and developmental events (PubMed:16762839, PubMed:17704056). Histone deacetylases act via the formation of large multiprotein complexes (PubMed:16762839). Histone deacetylases act via the formation of large multiprotein complexes (PubMed:16762839). Acts as a

component of the histone deacetylase NuRD complex which participates in the remodeling of



chromatin (PubMed:16428440, PubMed:28977666). As part of the SIN3B complex is recruited downstream of the constitutively active genes transcriptional start sites through interaction with histones and mitigates histone acetylation and RNA polymerase II progression within transcribed regions contributing to the regulation of transcription (PubMed:21041482). Also functions as a deacetylase for non-histone targets, such as NR1D2, RELA, SP1, SP3, STAT3 and

TSHZ3 (PubMed:12837748, PubMed:16285960, PubMed:16478997, PubMed:17996965, PubMed:19343227). Deacetylates SP proteins, SP1 and SP3, and regulates their function (PubMed:<a href="http://www.uniprot.org/citations/12837748"

target="_blank">12837748, PubMed:16478997). Component of the BRG1-RB1-HDAC1 complex, which negatively regulates the CREST-mediated transcription in resting neurons (PubMed:19081374). Upon calcium stimulation, HDAC1 is released from the complex and CREBBP is recruited, which facilitates transcriptional activation (PubMed:19081374). Deacetylates TSHZ3 and regulates its transcriptional repressor activity (PubMed:<a href="http://www.uniprot.org/citations/19343227"

target="_blank">19343227). Deacetylates 'Lys-310' in RELA and thereby inhibits the transcriptional activity of NF-kappa-B (PubMed:17000776). Deacetylates NR1D2 and abrogates the effect of KAT5- mediated relieving of NR1D2 transcription repression activity (PubMed:<a href="http://www.uniprot.org/citations/17996965"

target="_blank">17996965). Component of a RCOR/GFI/KDM1A/HDAC complex that suppresses, via histone deacetylase (HDAC) recruitment, a number of genes implicated in multilineage blood cell development (By similarity). Involved in CIART-mediated transcriptional repression of the circadian transcriptional activator: CLOCK-BMAL1 heterodimer (By similarity). Required for the transcriptional repression of circadian target genes, such as PER1, mediated by the large PER complex or CRY1 through histone deacetylation (By similarity). In addition to protein deacetylase activity, also has protein-lysine deacylase activity: acts as a protein decrotonylase by mediating decrotonylation ((2E)-butenoyl) of histones (PubMed:28497810).

Cellular Location Nucleus

Tissue Location Ubiquitous, with higher levels in heart, pancreas and testis, and lower levels in kidney and brain

Volume 50 μl

HDAC1 Antibody (aa466-482) - Protocols

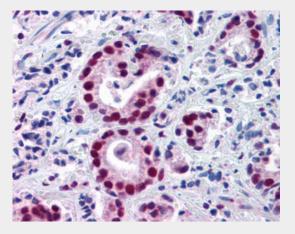
Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- <u>Blocking Peptides</u>
- <u>Dot Blot</u>
- Immunohistochemistry



- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

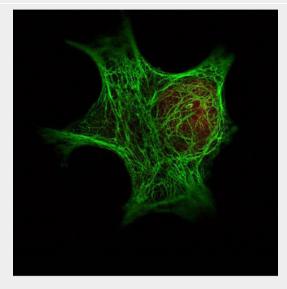
HDAC1 Antibody (aa466-482) - Images



Anti-HDAC1 antibody IHC of human prostate carcinoma.

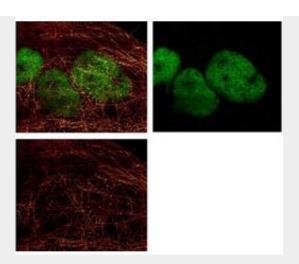


Anti-HDAC-1 Antibody - Western Blot.



Anti-HDAC-1 polyclonal antibody-Immunofluorescence Microscopy.





Immunofluorescence Microscopy - HDAC Antibody. HDAC1 Antibody (aa466-482) - Background

Responsible for the deacetylation of lysine residues on the N-terminal part of the core histones (H2A, H2B, H3 and H4). Histone deacetylation gives a tag for epigenetic repression and plays an important role in transcriptional regulation, cell cycle progression and developmental events. Histone deacetylases act via the formation of large multiprotein complexes. Deacetylates SP proteins, SP1 and SP3, and regulates their function. Component of the BRG1-RB1-HDAC1 complex, which negatively regulates the CREST- mediated transcription in resting neurons. Upon calcium stimulation, HDAC1 is released from the complex and CREBBP is recruited, which facilitates transcriptional activation. Deacetylates TSHZ3 and regulates its transcriptional repressor activity. Deacetylates 'Lys-310' in RELA and thereby inhibits the transcriptional activity of NF-kappa-B. Deacetylates NR1D2 and abrogates the effect of KAT5-mediated relieving of NR1D2 transcription repression activity. Component of a RCOR/GFI/KDM1A/HDAC complex that suppresses, via histone deacetylase (HDAC) recruitment, a number of genes implicated in multilineage blood cell development. Involved in CIART-mediated transcriptional repression of the circadian transcriptional activator: CLOCK-ARNTL/BMAL1 heterodimer. Required for the transcriptional repression of circadian target genes, such as PER1, mediated by the large PER complex or CRY1 through histone deacetylation.

HDAC1 Antibody (aa466-482) - References

Taunton J., et al. Science 272:408-411(1996). Furukawa Y., et al. Cytogenet. Cell Genet. 73:130-133(1996). Sparrow D.B., et al. EMBO J. 18:5085-5098(1999). Huynh K.D., et al. Genes Dev. 14:1810-1823(2000). Cai R.L., et al.J. Biol. Chem. 275:27909-27916(2000).