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# HDAC4 (N-terminus) Antibody

Purified Mouse Monoclonal Antibody (Mab)
Catalog # AP53277

## Product Information

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| **Application** | WB, IP |
| **Primary Accession** | P56524 |
| **Reactivity** | Human |
| **Host** | Mouse |
| **Clonality** | Monoclonal |
| **Isotype** | IgG2a |
| **Calculated MW** | 119040 |

## Additional Information

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| **Gene ID** | 9759 |
| **Other Names** | EC 3.5.1.98;HA6116;HD 4;HD4;HDAC 4;HDAC A;HDAC4;HDAC4_HUMAN;HDACA;Histone Deacetylase 4; Histone Deacetylase A;KIAA0288. |
| **Dilution** | WB~~1:1000 IP~~1:500 |
| **Format** | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3. |
| **Storage** | Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles. |

## Protein Information

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| **Name** | HDAC4 ( HGNC:14063) |
| **Synonyms** | KIAA0288 |
| **Function** | 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. Involved in muscle maturation via its interaction with the myocyte enhancer factors such as MEF2A, MEF2C and MEF2D. Involved in the MTA1-mediated epigenetic regulation of ESR1 expression in breast cancer. Deacetylates HSPA1A and HSPA1B at 'Lys-77' leading to their preferential binding to co-chaperone STUB1 (PubMed:27708256).<br><br>Nucleus. Cytoplasm. Note=Shuttles between the nucleus and the cytoplasm. |

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| **Cellular Location** | Upon muscle cells differentiation, it accumulates in the nuclei of myotubes, suggesting a positive role of nuclear HDAC4 in muscle differentiation. The export to cytoplasm depends on the interaction with a 14-3-3 chaperone protein and is due to its phosphorylation at Ser-246, Ser-467 and Ser-632 by CaMK4 and SIK1. The nuclear localization probably depends on sumoylation Interaction with SIK3 leads to HDAC4 retention in the cytoplasm (By similarity). {ECO:0000250\|UniProtKB:Q6NZM9} |
| **Tissue Location** | Ubiquitous. |

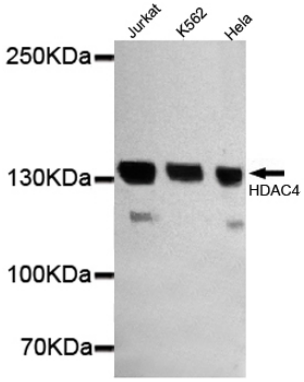
## Background

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. Involved in muscle maturation via its interaction with the myocyte enhancer factors such as MEF2A, MEF2C and MEF2D. Involved in the MTA1-mediated epigenetic regulation of ESR1 expression in breast cancer.

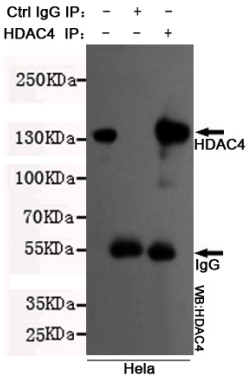
## References


Grozinger C.M.,et al.Proc. Natl. Acad. Sci. U.S.A. 96:4868-4873(1999).
Ohara O.,et al.DNA Res. 4:53-59(1997).
Ohara O.,et al.Submitted (DEC-1999) to the EMBL/GenBank/DDBJ databases.
Hillier L.W.,et al.Nature 434:724-731(2005).
Mural R.J.,et al.Submitted (SEP-2005) to the EMBL/GenBank/DDBJ databases.


## Images




Western blot detection of HDAC4 in Jurkat,Hela and K562 cell lysates using HDAC4 mouse mAb (1:1000 diluted).Predicted band size: 140KDa.Observed band size: 140KDa.




Immunoprecipitation analysis of Hela cell lysates using HDAC4 mouse mAb.

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