

# **HDAC4** Antibody

Rabbit mAb Catalog # AP90913

### **Product Information**

ApplicationWB, IF, ICCPrimary AccessionP56524ReactivityHumanClonalityMonoclonal

Other Names Histone deacetylase 4; HD4; HDAC4; AHO3; BDMR;

IsotypeRabbit IgGHostRabbitCalculated MW119040

# **Additional Information**

**Dilution** WB 1:1000~1:2000 ICC/IF 1:50~1:200

**Purification** Affinity-chromatography

**Immunogen** A synthesized peptide derived from human HDAC4

**Description** 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.

Storage Condition and Buffer Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium

azide and 50% glycerol. Store at +4°C short term. Store at -20°C long term.

Avoid freeze / thaw cycle.

# **Protein Information**

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:<u>27708256</u>).

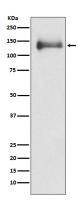
#### **Cellular Location**

Nucleus. Cytoplasm. Note=Shuttles between the nucleus and the cytoplasm. 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.

# **Images**



Western blot analysis of HDAC4 expression in Hela lysate.

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