abcam

Product datasheet

Anti-HDAC4 antibody ab79521

8 References 1 Image

Overview

Product name Anti-HDAC4 antibody

Description Rabbit polyclonal to HDAC4

Host species Rabbit

Tested applications Suitable for: IHC-P

Species reactivity Reacts with: Human

Predicted to work with: Rat, Chicken, Zebrafish, Orangutan

Immunogen Synthetic peptide:

CISSDPRYWYGKTQHS

, corresponding to amino acids 194-209 of Human HDAC4

Run BLAST with
Run BLAST with

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer Preservative: 0.02% Sodium azide

Constituent: PBS

Purity Protein G purified

Clonality Polyclonal

Isotype IgG

Applications

1

The Abpromise guarantee

Our Abpromise quarantee covers the use of ab79521 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		Use a concentration of 5 µg/ml.

Target

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.

Tissue specificity

Ubiquitous.

Involvement in disease

Defects in HDAC4 are the cause of brachydactyly-mental retardation syndrome (BDMR) [MIM:600430]. A syndrome resembling the physical anomalies found in Albright hereditary osteodystrophy. Common features are mild facial dysmorphism, congenital heart defects, distinct brachydactyly type E, mental retardation, developmental delay, seizures, autism spectrum disorder, and stocky build. Soft tissue ossification is absent, and there are no abnormalities in parathyroid hormone or calcium metabolism.

Sequence similarities

Belongs to the histone deacetylase family. HD type 2 subfamily.

Domain

The nuclear export sequence mediates the shuttling between the nucleus and the cytoplasm.

Post-translational modifications

 $Phosphory lated \ by \ CaMK4 \ at \ Ser-246, \ Ser-467 \ and \ Ser-632. \ Phosphory lation \ at \ other \ residues$

is required for the interaction with 14-3-3.

Sumoylation on Lys-559 is promoted by the E3 SUMO-protein ligase RANBP2, and prevented by

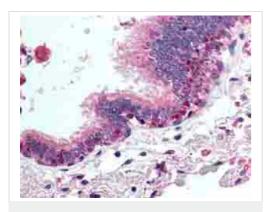
phosphorylation by CaMK4.

Cellular localization

Nucleus. Cytoplasm. 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. The nuclear localization probably depends on sumovlation.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HDAC4 antibody (ab79521)

ab79521 at 5µg/ml staining human HDAC4 by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded lung, respiratory epithelium sections).

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