Performance guarenteed



Product Details

Size	400 µL		
Species Reactivity	Human, Mouse		
Published Species	Human		
Host/Isotype	Rabbit / IgG		
Class	Polyclonal		
Туре	Antibody		
Conjugate	Unconjugated		
Immunogen	KLH conjugated synthetic peptide between 2-32 amino acids from the N-terminal region of human HDAC9		
Form	Liquid		
Concentration	2 mg/mL		
Purification	Ammonium sulfate precipitation, Size-exclusion - Dialysis		
Storage buffer	PBS		
Contains	0.09% sodium azide		
Storage conditions	-20° C, Avoid Freeze/Thaw Cycles		
RRID	AB_2117071		

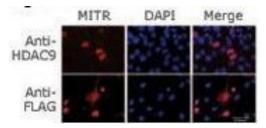
Applications	Tested Dilution	Publications
Western Blot (WB)	1:1,000	1 Publication
Immunohistochemistry (Paraffin) (IHC (P))	1:50-1:100	-
Immunocytochemistry (ICC/IF)	1:1,000	-
Immunoprecipitation (IP)	1:100	-
ChIP assay (ChIP)	1-3 μL	-

Product Specific Information

This antibody is predicted to react with chicken and mouse based on sequence homology.

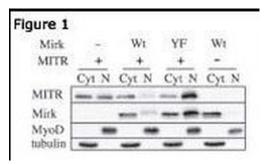
1

Product Images For HDAC9 Polyclonal Antibody



HDAC9 Antibody (PA5-11245) in ICC/IF

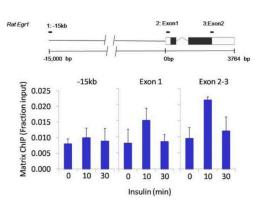
Immunofluorescent analysis of MITR for a compartmentalization study in undifferentiated C2C12 myoblasts transfected with a MITR-expressing plasmid. MITR is detected by using an anti-HDAC9 polyclonal antibody (Product # PA5-11245) (top panel) or a FLAG antibody (bottom panel) detecting a FLAG epitope fused at the N-term end of the MITR construct.



HDAC9 Antibody (PA5-11245) in WB

Western blot analysis of HDAC9 in C2C12 cells using a HDAC9 polyclonal antibody (Product # PA5-11245). Cells were transfected with plasmids coding for Mirk (Wt), kinase-inactive Mirk (YF) or MITR. Mirk, MyoD and tubulin proteins are shown for cytoplasmic (Cyt) and nuclear (N) extracts from undifferentiated C2C12 myoblasts.





Chromatin immunoprecipitation analysis of HDAC9 was performed using crosslinked chromatin from 1x10⁶ HTC-IR rat hepatoma cells treated with insulin for 0, 10, and 30 minutes. Immunoprecipitation was performed using a multiplex microplate Matrix ChIP assay (see reference for Matrix ChIP protocol: http://www. ncbi.nlm.nih.gov/pubmed/22098709) with 1.0 µL/100 µL well volume of an HDAC9 polyclonal antibody (Product # PA5-11245). Chromatin aliquots from ~1x10^5 cells were used per ChIP pull-down. Quantitative PCR data were done in guadruplicate using 1 µL of eluted DNA in 2 µL SYBR real-time PCR reactions containing primers to amplify -15kb upstream of the Eqr1 gene or exon-1 or exon-2-3 of Egr1. PCR calibration curves were generated for each primer pair from a dilution series of sheared total genomic DNA. Quantitation of immunoprecipitated chromatin is presented as signal relative to the total amount of input chromatin. Results represent the mean +/- SEM for three experiments. A schematic representation of the rat Egr-1 locus is shown above the data where boxes represent exons (black boxes = translated regions, white boxes = untranslated regions), the zigzag line represents an intron, and the straight line represents upstream sequence. Regions amplified by Egr-1 primers are represented by black bars. Data courtesy of the Innovators Program.

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□ 1 Reference

Western Blot (1)

BioMed research international

Histone Deacetylase9 Represents the Epigenetic Promotion of M1 Macrophage Polarization and Inflammatory Response via TLR4 Regulation.

"PA5-11245 was used in Western Blotting to indicate that toll-like receptor 4 (TLR4) was not only positively correlated to the HDAC9 gene, but was also upregulated in atherosclerosis, where it was also significantly upregulated in the atherosclerosis cell model of oxidized low-density lipoprotein-induced macrophages."

Authors: Cao X, Zhang M, Li H, Chen K, Wang Y, Yang J

Year 2022

Species Human

Dilution 1:1000

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