# abcam

## Product datasheet

# HRP Anti-HDAC2 antibody [Y461] ab195851

KO VALIDATED

Recombinant RabMAb

2 References 3 Images

Overview

**Product name** HRP Anti-HDAC2 antibody [Y461]

**Description** HRP Rabbit monoclonal [Y461] to HDAC2

**Host species** Rabbit HRP Conjugation

**Tested applications** Suitable for: WB Species reactivity

Reacts with: Human

Predicted to work with: Mouse, Rat

Synthetic peptide. This information is proprietary to Abcam and/or its suppliers. **Immunogen** 

Positive control WB: K562 whole cell lysate.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit **General notes** 

monoclonal antibodies. For details on our patents, please refer to **RabMAb® patents**.

**Properties** 

**Form** Liquid

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C. Storage instructions

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.1% Proclin 300 Solution

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

**Purity** Protein A purified

Clonality Monoclonal

Clone number Y461 Isotype ΙgG

**Applications** 

Our Abpromise guarantee covers the use of ab195851 in the following tested applications. The Abpromise guarantee

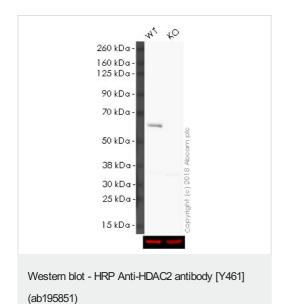
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/10000. Detects a band of approximately 60 kDa (predicted molecular weight: 55 kDa).

#### **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. Forms transcriptional repressor complexes by associating with MAD, SIN3, YY1 and N-COR. Interacts in the late S-phase of DNA-replication with DNMT1 in the other transcriptional repressor complex composed of DNMT1, DMAP1, PCNA, CAF1. Deacetylates TSHZ3 and regulates its transcriptional repressor activity.	
Tissue specificity	Widely expressed; lower levels in brain and lung.	
Sequence similarities	Belongs to the histone deacetylase family. HD type 1 subfamily.	
Post-translational modifications	S-nitrosylated by GAPDH. In neurons, S-Nitrosylation at Cys-262 and Cys-274 does not affect the enzyme activity but abolishes chromatin-binding, leading to increases acetylation of histones and activate genes that are associated with neuronal development. In embryonic cortical neurons, S-Nitrosylation regulates dendritic growth and branching.	
Cellular localization	Nucleus.	

### **Images**



**All lanes :** HRP Anti-HDAC2 antibody [Y461] (ab195851) at 1/10000 dilution

Lane 1 : Wild-type HAP1 whole cell lysate

Lane 2: HDAC2 knockout HAP1 whole cell lysate

Lysates/proteins at 20 µg per lane.

**Predicted band size:** 55 kDa **Observed band size:** 55 kDa

Exposure time: 30 seconds

ab195851 was shown to specifically react with HDAC2 in wild-type HAP1 cells as signal was lost in HDAC2 knockout cells. Wild-type and HDAC2 knockout samples were subjected to SDS-PAGE.

Ab195851 and <u>ab9484</u> (Mouse anti-GAPDH loading control) were incubated overnight at 4°C at 1/10000 dilution and 1/20000 dilution respectively. The loading control was imaged using the Licor Odyssey CLx prior to blots being developed with ECL technique.

1
250 kDa —
150 kDa —
100 kDa —
75 kDa —
37 kDa —
37 kDa —
25 kDa —
20 kDa —
20 kDa —
15 kDa —
10 kDa —

(ab195851)

HRP Anti-HDAC2 antibody [Y461] (ab195851) at 1/10000 dilution + K562 (Human erythromyeloblastoid leukemia cell line) Whole Cell Lysate at 10  $\mu g$ 

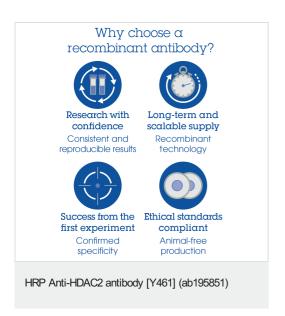
Developed using the ECL technique.

Performed under reducing conditions.

**Predicted band size:** 55 kDa **Observed band size:** 60 kDa

Exposure time: 30 seconds

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 3% milk before being incubated with ab195851 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406** 



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