abcam

Product datasheet

HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker ab195897





4 Images

Overview

Product name HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker

HRP Rabbit monoclonal [EPR6825] to KDM1/LSD1 - Nuclear Marker **Description**

Host species Rabbit Conjugation HRP

Tested applications Suitable for: IHC-P, WB Species reactivity Reacts with: Rat. Human

Predicted to work with: Mouse

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

WB: HeLa, HAP1, Jurkat and PC12 whole cell lysates. IHC-P: FFPE Human Testis Normal Positive control **General notes**

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity - Long-term security of supply - Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb® patents**.

Properties

Form Liquid

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

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 EPR6825

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab195897 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		1/50. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		1/5000. Detects a band of approximately 110 kDa (predicted molecular weight: 92 kDa).

Target

Function

Histone demethylase that demethylates both 'Lys-4' (H3K4me) and 'Lys-9' (H3K9me) of histone H3, thereby acting as a coactivator or a corepressor, depending on the context. Acts by oxidizing the substrate by FAD to generate the corresponding imine that is subsequently hydrolyzed. Acts as a corepressor by mediating demethylation of H3K4me, a specific tag for epigenetic transcriptional activation. Demethylates both mono- (H3K4me1) and di-methylated (H3K4me2) H3K4me. May play a role in the repression of neuronal genes. Alone, it is unable to demethylate H3K4me on nucleosomes and requires the presence of RCOR1/CoREST to achieve such activity. Also acts as a coactivator of androgen receptor (ANDR)-dependent transcription, by being recruited to ANDR target genes and mediating demethylation of H3K9me, a specific tag for epigenetic transcriptional repression. The presence of PRKCB in ANDR-containing complexes, which mediates phosphorylation of 'Thr-6' of histone H3 (H3T6ph), a specific tag that prevents demethylation H3K4me, prevents H3K4me demethylase activity of KDM1A. Demethylates dimethylated 'Lys-370' of p53/TP53 which prevents interaction of p53/TP53 with TP53BP1 and represses p53/TP53-mediated transcriptional activation. Demethylates and stabilizes the DNA methylase DNMT1. Required for gastrulation during embryogenesis.

Tissue specificity Ubiquitously expressed.

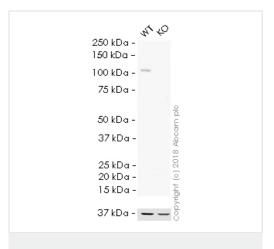
Sequence similarities Belongs to the flavin monoamine oxidase family.

Contains 1 SWIRM domain.

Domain The SWIRM domain may act as an anchor site for a histone tail.

Cellular localization Nucleus.

Images



Western blot - HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker (ab195897) **All lanes :** HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker (ab195897) at 1/5000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

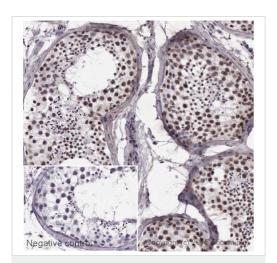
Lane 2: KDM1A (KDM1 / LSD1) knockout HAP1 whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 92 kDa **Observed band size:** 110 kDa

Exposure time: 20 minutes

ab195897 was shown to specifically react with KDM1 / LSD1 in wild-type HAP1 cells as signal was lost in KDM1A (KDM1 / LSD1) knockout cells. Wild-type and KDM1A (KDM1 / LSD1) knockout samples were subjected to SDS-PAGE. Ab195897 and **ab184095** (Mouse monoclonal [mAbcam 9484] to GAPDH - Loading Control (Alexa Fluor[®] 680) loading control) were incubated overnight at 4°C at 1/5000 dilution and 1/20000 dilution respectively. The loading control was imaged using the Licor Odyssey CLx prior to blots being developed with ECL technique.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker (ab195897)

IHC image of KDM1/LSD1 staining in a section of formalin-fixed paraffin-embedded normal human testis*. The section was pretreated using pressure cooker heat mediated antigen retrieval with sodium citrate buffer (pH6) for 30mins, and incubated overnight at +4°C with ab195897 at 1/50 dilution. DAB was used as the chromogen (ab103723), diluted 1/100 and incubated for 10min at room temperature. The section was counterstained with haematoxylin and mounted with DPX. The inset negative control image is taken from an identical assay without primary antibody.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

^{*}Tissue obtained from the Human Research Tissue Bank,

supported by the NIHR Cambridge Biomedical Research Centre



Western blot - HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker (ab195897) **All lanes :** HRP Anti-KDM1/LSD1 antibody [EPR6825] - Nuclear Marker (ab195897) at 1/5000 dilution

Lane 1: HeLa whole cell lysate (ab150035)

Lane 2 : Jurkat (Human T cell lymphoblast-like cell line) Whole Cell Lysate

Lane 3: PC12 (Rat adrenal pheochromocytoma cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

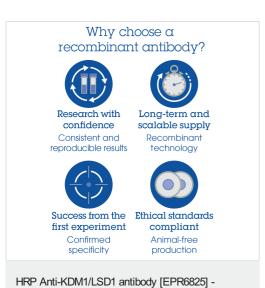
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 92 kDa **Observed band size:** 110 kDa

Exposure time: 4 minutes

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 ab195897 overnight at 4°C. Antibody binding was visualised using ECL development solution **ab133406**.



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Nuclear Marker (ab195897)

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