


Anti-HIF-1 alpha antibody [H1alpha67] ab1

★★★★★ [44 Abreviews](#) [340 References](#) [5 图像](#)

概述

产品名称	Anti-HIF-1 alpha抗体[H1alpha67]
描述	小鼠单克隆抗体[H1alpha67] to HIF-1 alpha
宿主	Mouse
经测试应用	适用于: IP, ICC/IF, WB, Flow Cyt (Intra) 不适用于: IHC-Fr or IHC-P
种属反应性	与反应: Human 预测可用于: Mouse, Rat 
免疫原	Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.
阳性对照	WB: HeLa (DFO treated 0.5mM, 24h) nuclear lysate (ab180880); Human whole cell lysate (human lung adenocarcinoma cell line ADLC-5M2) (DFO treated 100uM, 16h). IP: HeLa (DFO treated 0.5 mg) nuclear lysate. ICC/F: MCF7 cells. Flow Cyt (Intra): HeLa (Human epithelial cell line from cervix adenocarcinoma) cells treated with 1mM Deferoxamine (ab120727) for 24 hours.
常规说明	<p>For WB, we recommend using positive control samples such as DFO or CoCl₂ treated nuclear cell lysates such as ab180880. Ensure cell lysis occurs quickly (within 2 mins) if removed from hypoxia. Loading a high amount of sample (>50 µg) and addition of protease inhibitors (e.g. ab65621) may also enhance detection.</p> <p>This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or conjugation for your experiments, please contact orders@abcam.com.</p> <p>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.</p> <p>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</p>

性能

形式	Liquid
存放说明	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C. Avoid freeze / thaw cycle.

of common human cancers and their metastases, due to the presence of intratumoral hypoxia and as a result of mutations in genes encoding oncoproteins and tumor suppressors.

序列相似性

Contains 1 basic helix-loop-helix (bHLH) domain.

Contains 1 PAC (PAS-associated C-terminal) domain.

Contains 2 PAS (PER-ARNT-SIM) domains.

结构域

Contains two independent C-terminal transactivation domains, NTAD and CTAD, which function synergistically. Their transcriptional activity is repressed by an intervening inhibitory domain (ID).

翻译后修饰

In normoxia, is hydroxylated on Pro-402 and Pro-564 in the oxygen-dependent degradation domain (ODD) by EGLN1/PHD1 and EGLN2/PHD2. EGLN3/PHD3 has also been shown to hydroxylate Pro-564. The hydroxylated prolines promote interaction with VHL, initiating rapid ubiquitination and subsequent proteasomal degradation. Deubiquitinated by USP20. Under hypoxia, proline hydroxylation is impaired and ubiquitination is attenuated, resulting in stabilization.

In normoxia, is hydroxylated on Asn-803 by HIF1AN, thus abrogating interaction with CREBBP and EP300 and preventing transcriptional activation. This hydroxylation is inhibited by the Cu/Zn-chelator, Clioquinol.

S-nitrosylation of Cys-800 may be responsible for increased recruitment of p300 coactivator necessary for transcriptional activity of HIF-1 complex.

Requires phosphorylation for DNA-binding.

Sumoylated; by SUMO1 under hypoxia. Sumoylation is enhanced through interaction with RWDD3. Desumoylation by SENP1 leads to increased HIF1A stability and transcriptional activity. Ubiquitinated; in normoxia, following hydroxylation and interaction with VHL. Lys-532 appears to be the principal site of ubiquitination. Clioquinol, the Cu/Zn-chelator, inhibits ubiquitination through preventing hydroxylation at Asn-803.

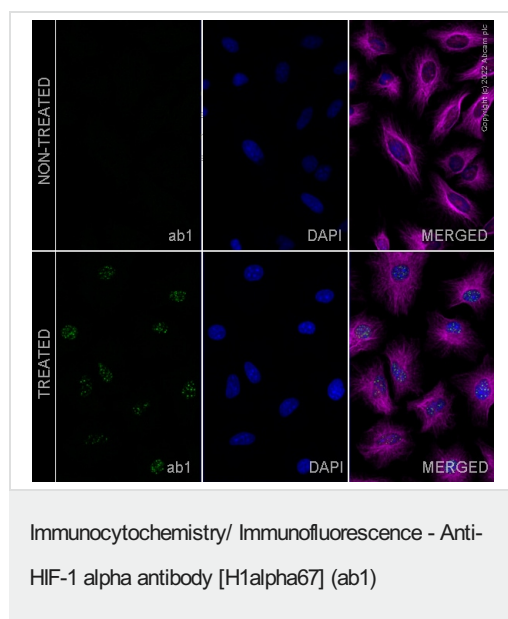
The iron and 2-oxoglutarate dependent 3-hydroxylation of asparagine is (S) stereospecific within HIF CTAD domains.

细胞定位

Cytoplasm. Nucleus. Cytoplasmic in normoxia, nuclear translocation in response to hypoxia.

Colocalizes with SUMO1 in the nucleus, under hypoxia.

图片

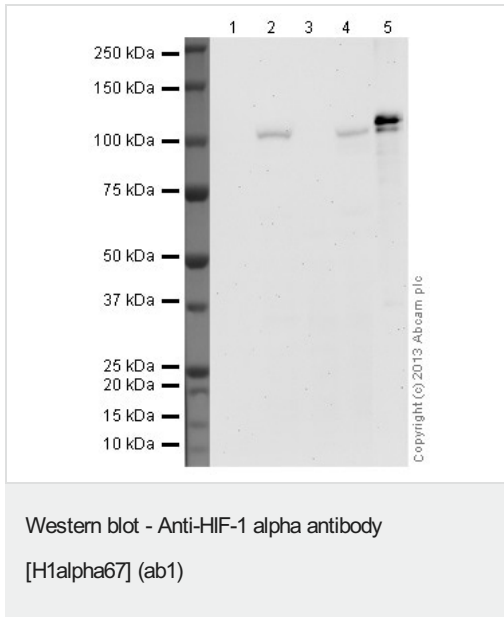


ab1 staining HIF-1 alpha in HeLa DFO cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab1 at 10µg/ml and [ab6046](#), Rabbit polyclonal to beta Tubulin - Loading Control. Cells were then incubated with [ab150117](#), Goat polyclonal Secondary Antibody to Mouse IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (shown in green) and [ab150080](#), Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 594) at 1/1000 dilution (shown in pseudocolour magenta). Nuclear DNA was labelled with DAPI (shown in blue).

Also suitable in cells fixed with 4% paraformaldehyde (10 min).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal

sections is shown.



All lanes : Anti-HIF-1 alpha antibody [H1alpha67] (ab1) at 5 µg/ml

Lane 1 : HeLa nuclear extract lysate (**ab150036**) at 40 µg

Lane 2 : HeLa-DFO treated (0.5mM, 24h) Nuclear Lysate (**ab180880**) at 40 µg

Lane 3 : HeLa nuclear control at 40 µg

Lane 4 : HeLa nuclear DFO treated at 40 µg

Lane 5 : Recombinant Human HIF-1 alpha protein (**ab154478**) at 0.001 µg

Secondary

All lanes : Goat Anti-Mouse IgG H&L (HRP) preadsorbed (**ab97040**) at 1/10000 dilution

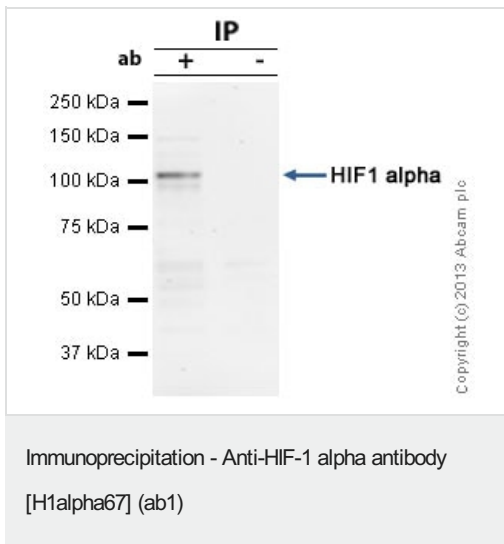
Performed under reducing conditions.

Predicted band size: 92 kDa

Exposure time: 20 minutes

We recommend using 5% milk in TBST as the blocking agent, decreasing to 2% milk in TBST during primary and secondary antibody incubation.

Blots were developed with **Goat Anti-Mouse IgG H&L (HRP) preadsorbed (ab97040) secondary antibody**



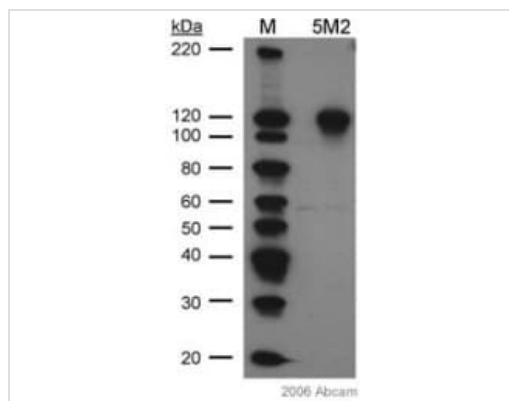
HIF-1 alpha was immunoprecipitated using 0.5 mg HeLa Nuclear DFO treated whole cell extract (**ab180880**), 5 µg of Mouse monoclonal to HIF-1 alpha and 50 µl of protein G magnetic beads (+). No antibody was added to the control (-).

The antibody was incubated under agitation with Protein G beads for 10 minutes, HeLa DFO treated whole cell extract lysate diluted in RIPA buffer was added to each sample and incubated for a further 10 minutes under agitation.

Proteins were eluted by addition of 40 µl SDS loading buffer and incubated for 10 minutes at 70°C; 10 µl of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with ab1.

Secondary: Goat polyclonal to mouse IgG light chain specific (HRP)
at 1:20,000 dilution.

Band: 110 kDa; HIF1 alpha



Western blot - Anti-HIF-1 alpha antibody

[H1alpha67] (ab1)

This image is taken from an Abreview submitted by Mike Campa, no further information is known about this image

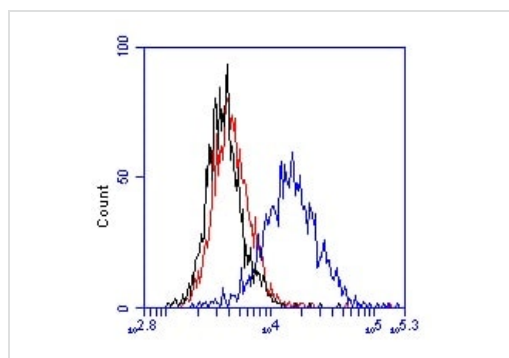
Anti-HIF-1 alpha antibody [H1alpha67] (ab1) at 1/400 dilution +
Human whole cell lysate (human lung adenocarcinoma cell line
ADLC-5M2) treated for 16 hours with 100 micromolar
deferroxamine (DFO) at 20 µg

Performed under reducing conditions.

Predicted band size: 92 kDa

Observed band size: 120 kDa

PVDF membrane was used and blocked for 16 hours in 5% milk.



Flow Cytometry (Intracellular) - Anti-HIF-1 alpha
antibody [H1alpha67] (ab1)

Flow cytometry using ab1. HeLa (Human epithelial cell line from
cervix adenocarcinoma) cells were cultured untreated or with 1mM
Deferoxamine (**ab120727**) for 24 hours to induce HIF-1-alpha
protein levels. Cells were then trypsinized, fixed with
paraformaldehyde and stained with ab1 (0.5 µg/mL). 1% BSA in
PBS was used as the blocking buffer throughout. ab1 was labeled
with and anti-mouse Alexa-Fluor® 488 dye. Unstained (black),
untreated (red) and DFO treated (blue) cell traces are shown.

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