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[KO Validated] HIF1AN/FIH-1 Rabbit mAb

Catalog No.: A27299 KO Validated Recombinant

Basic Information

Observed MW

40kDa

Calculated MW

41kDa

Category

Primary antibody

Applications

WB,IHC-P,IF/ICC,ELISA

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC73051

Background

Enables several functions, including 2-oxoglutarate-dependent dioxygenase activity; NF-kappaB binding activity; and transition metal ion binding activity. Involved in several processes, including negative regulation of Notch signaling pathway; negative regulation of transcription from RNA polymerase II promoter in response to hypoxia; and protein hydroxylation. Located in cytosol; nucleoplasm; and perinuclear region of cytoplasm. Colocalizes with nucleus.

Recommended Dilutions

WB 1:8000 - 1:32000

IHC-P 1:100 - 1:400

IF/ICC 1:100 - 1:400

ELISA Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

Immunogen Information

Gene ID55662

Swiss Prot
Q9NWT6

Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

Synonyms

FIH1

Contact

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\odot	Ī	www.abclonal.com.cn

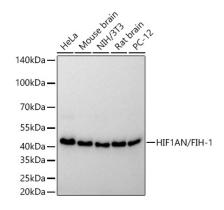
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Western blot analysis of various lysates using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at 1:16000 dilution incubated overnight at 4°C.

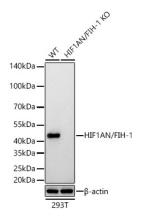
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 45s.

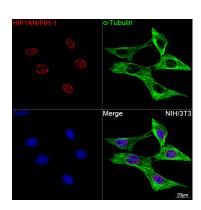


Western blot analysis of lysates from wild type (WT) and HIF1AN/FIH-1 knockout (KO) 293T cells using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at 1:16000 dilution incubated overnight at 4°C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

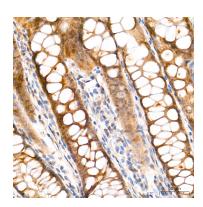
Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020).

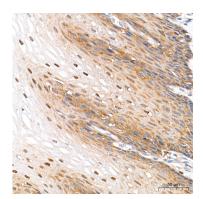
Exposure time: 45s.



Confocal imaging of NIH/3T3 cells using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit lgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo \$ 488-conjugated Goat Anti-Mouse lgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.

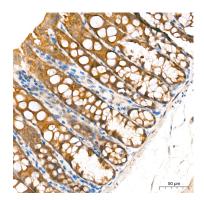


Immunohistochemistry analysis of paraffinembedded Human colon tissue using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Human esophagus tissue using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

Validation Data



Immunohistochemistry analysis of paraffinembedded Mouse colon tissue using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat testis tissue using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded 293T and 293T-HIF1AN-KO cells using [KO Validated] HIF1AN/FIH-1 Rabbit mAb (A27299) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.