HEXIM1 Rabbit mAb

Catalog No.: A24208 Recombinant



Basic Information

Observed MW

65kDa

Calculated MW

41kDa

Category

Primary antibody

Applications

ELISA,WB,IHC-P,IF/ICC,IP

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC65105

IHC-P

Background

Expression of this gene is induced by hexamethylene-bis-acetamide in vascular smooth muscle cells. This gene has no introns.

Recommended Dilutions

WB 1:500 - 1:1000

IF/ICC 1:50 - 1:200

IP 0.5μg-4μg antibody for 200μg-400μg extracts of

whole cells

1:50 - 1:200

Immunogen Information

Gene ID Swiss Prot 10614 094992

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 255-359 of human HEXIM1(NP_006451.1).

Synonyms

CLP1; EDG1; HIS1; MAQ1; HEXIM1

Contact

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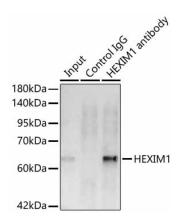
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

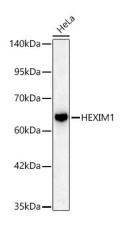
Storage

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

Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.



Immunoprecipitation of HEXIM1 in 300 μg extracts from 293T cells using 3 μg HEXIM1 Rabbit mAb (A24208). Western blot analysis was performed using HEXIM1 Rabbit mAb (A24208) at 1:1000 dilution.



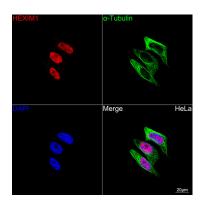
Western blot analysis of lysates from HeLa cells using HEXIM1 Rabbit mAb(A24208) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

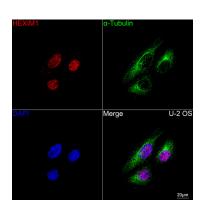
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

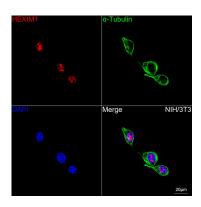
Exposure time: 25s.



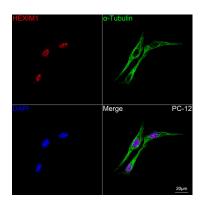
Confocal imaging of HeLa cells using HEXIM1 Rabbit mAb (A24208, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500)(Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



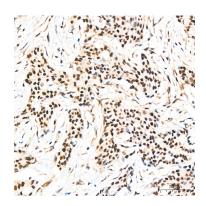
Confocal imaging of U-2 OS cells using HEXIM1 Rabbit mAb (A24208,dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007,dilution 1:500)(Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



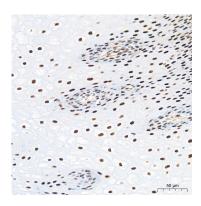
Confocal imaging of NIH/3T3 cells using HEXIM1 Rabbit mAb (A24208,dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007,dilution 1:500)(Red). The cells were counterstained with α-Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green).DAPI was used for nuclear staining (Blue). Objective: 100x.



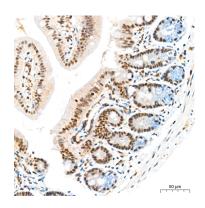
Confocal imaging of PC-12 cells using HEXIM1 Rabbit mAb (A24208,dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007,dilution 1:500)(Red). The cells were counterstained with α-Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



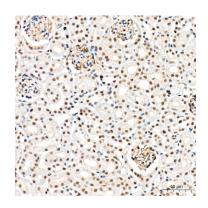
Immunohistochemistry analysis of HEXIM1 in paraffin-embedded human breast cancer tissue using HEXIM1 Rabbit mAb (A24208) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of HEXIM1 in paraffin-embedded human esophagus tissue using HEXIM1 Rabbit mAb (A24208) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



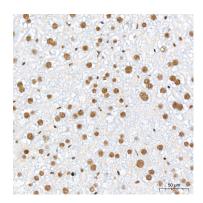
Immunohistochemistry analysis of HEXIM1 in paraffin-embedded mouse intestin tissue using HEXIM1 Rabbit mAb (A24208) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of HEXIM1 in paraffin-embedded mouse kidney tissue using HEXIM1 Rabbit mAb (A24208) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of HEXIM1 in paraffin-embedded rat colon tissue using HEXIM1 Rabbit mAb (A24208) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of HEXIM1 in paraffin-embedded rat liver tissue using HEXIM1 Rabbit mAb (A24208) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.