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[KD Validated] Calreticulin Rabbit mAb

Catalog No.: A20986 Recombinant 3 Publications

Basic Information

Observed MW

55kDa

Calculated MW

47kDa

Category

Primary antibody

Applications

ELISA,WB,IHC-P,IF/ICC

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC51269

Background

Calreticulin is a highly conserved chaperone protein which resides primarily in the endoplasmic reticulum, and is involved in a variety of cellular processes, among them, cell adhesion. Additionally, it functions in protein folding quality control and calcium homeostasis. Calreticulin is also found in the nucleus, suggesting that it may have a role in transcription regulation. Systemic lupus erythematosus is associated with increased autoantibody titers against calreticulin. Recurrent mutations in calreticulin have been linked to various neoplasms, including the myeloproliferative type.

Recommended Dilutions

WB	1:500 - 1:1000
IHC-P	1:1000 - 1:5000
IF/ICC	1:50 - 1:200

Immunogen Information

Gene ID	Swiss Prot
811	P27797

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 18-203 of human Calreticulin (NP_004334.1).

Synonyms

RO; CRT; SSA; cC1qR; HEL-S-99n; [KD Validated] Calreticulin

Contact

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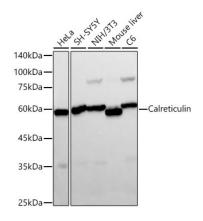
Product Information

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

Storage

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

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



Western blot analysis of various lysates using [KD Validated] Calreticulin Rabbit mAb (A20986) at 1:1000 dilution incubated overnight at 4°C.

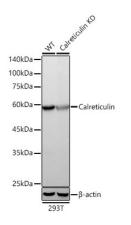
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.

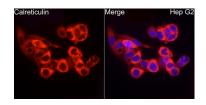


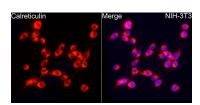
Western blot analysis of lysates from wild type (WT) and Calreticulin knockdown (KD) 293T cells using [KD Validated] Calreticulin Rabbit mAb (A20986) at 1:1000 dilution incubated overnight at 4° C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 µg per lane.

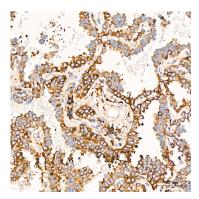
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



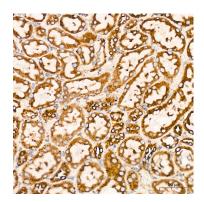




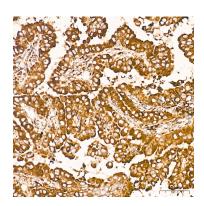
Confocal imaging of Hep G2 cells using [KD Validated] Calreticulin Rabbit mAb (A20986,dilution 1:50)(Red) . DAPI was used for nuclear staining (blue). Objective: 40x.

Confocal imaging of NIH/3T3 cells using [KD Validated] Calreticulin Rabbit mAb (A20986,dilution 1:50) (Red). DAPI was used for nuclear staining (blue). Objective: 40x.

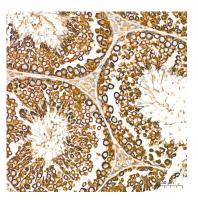
Immunohistochemistry analysis of Calreticulin in paraffin-embedded Human lung adenocarcinoma tissue using [KD Validated] Calreticulin Rabbit mAb (A20986) at a dilution of 1:2000 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



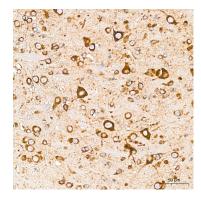
Immunohistochemistry analysis of Calreticulin in paraffin-embedded rat kidney tissue using [KD Validated] Calreticulin Rabbit mAb (A20986) at a dilution of 1:2000 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of Calreticulin in paraffin-embedded human thyroid cancer tissue using [KD Validated] Calreticulin Rabbit mAb (A20986) at a dilution of 1:2000 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of Calreticulin in paraffin-embedded mouse testis tissue using [KD Validated] Calreticulin Rabbit mAb (A20986) at a dilution of 1:2000 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of Calreticulin in paraffin-embedded mouse brain tissue using [KD Validated] Calreticulin Rabbit mAb (A20986) at a dilution of 1:2000 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.