

Anti-Calnexin/CANX Antibody Picoband™

Catalog Number: A03372-2

About CANX

Calnexin (CNX) is a 67 kDa integral protein of the endoplasmic reticulum. This gene encodes a member of the calnexin family of molecular chaperones. The encoded protein is a calcium-binding, endoplasmic reticulum (ER)-associated protein that interacts transiently with newly synthesized N-linked glycoproteins, facilitating protein folding and assembly. It may also play a central role in the quality control of protein folding by retaining incorrectly folded protein subunits within the ER for degradation. Alternatively spliced transcript variants encoding different isoforms have been described.

Overview

Product Name	Anti-Calnexin/CANX Antibody Picoband™
Reactive Species	Human, Mouse
Description	Boster Bio Anti-Calnexin/CANX Antibody Picoband™ catalog # A03372-2. Tested in ELISA, Flow Cytometry, IF, IHC, ICC, WB applications. This antibody reacts with Human, Mouse.
Application	ELISA, Flow Cytometry, IF, IHC, ICC, WB
Clonality	Polyclonal
Formulation	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na ₂ HPO ₄ .
Storage Instructions	At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and thawing.
Host	Rabbit
Uniprot ID	P27824

Technical Details

Immunogen	E.coli-derived human Calnexin/CANX recombinant protein (Position: E68-R582).
Recommended Detection Systems	Boster recommends Enhanced Chemiluminescent Kit with anti-Rabbit IgG (EK1002) for Western blot, and HRP Conjugated anti-Rabbit IgG Super Vision Assay Kit (SV0002-1) for IHC(P) and ICC.
Cross Reactivity	No cross-reactivity with other proteins.
Isotype	Rabbit IgG
Form	Lyophilized
Concentration	Adding 0.2 ml of distilled water will yield a concentration of 500 ug/ml.
Purification	Immunogen affinity purified.

Suggested Dilutions

Dilute the sample so that the expected range of concentrations fall within the detection range of this kit.

If the expected range of concentration is unknown, a pilot test should be conducted to decide the optimal dilution ratio for your samples.

Some PubMed article(s) citing the expression level of this target are as follows:

Boster Bio's internal QC testing used:

Western blot, 0.1-0.25 ug/ml, Human, Mouse

Immunohistochemistry(Paraffin-embedded Section), 2-5 ug/ml, Human

Immunocytochemistry/Immunofluorescence, 5 ug/ml, Human

Flow Cytometry, 1-3 ug/1x10⁶ cells, Human

Direct ELISA, 0.1-0.5 ug/ml, Human

Anti-Calnexin/CANX Antibody Picoband™ (A03372-2) Images

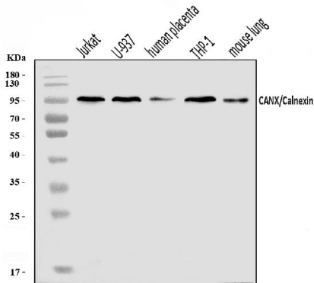


Figure 1. Western blot analysis of Calnexin/CANX using anti-Calnexin/CANX antibody (A03372-2).

Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human Jurkat whole cell lysates,

Lane 2: human U-937 whole cell lysates,

Lane 3: human placenta tissue lysates,

Lane 4: human THP-1 whole cell lysates,

Lane 5: mouse lung tissue lysates.

After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes.

Blocked the membrane with 5% non-fat milk/TBS for 1.5

hour at RT. The membrane was incubated with rabbit anti-

Calnexin/CANX antigen affinity purified polyclonal antibody

(Catalog # A03372-2) at 0.25 ug/mL overnight at 4°C, then

washed with TBS-0.1%Tween 3 times with 5 minutes each

and probed with a goat anti-rabbit IgG-HRP secondary

antibody at a dilution of 1:5000 for 1.5 hour at RT. The

signal is developed using an Enhanced Chemiluminescent

detection (ECL) kit (Catalog # EK1002) with Tanon 5200

system. A specific band was detected for Calnexin/CANX at

approximately 97 kDa. The expected band size for

Calnexin/CANX is at 97 kDa.

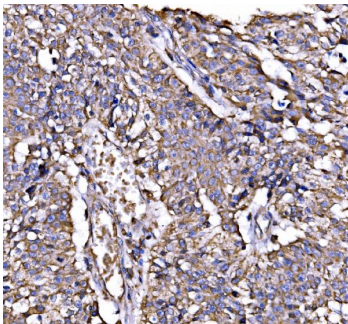


Figure 2. IHC analysis of Calnexin/CANX using anti-Calnexin/CANX antibody (A03372-2).

Calnexin/CANX was detected in a paraffin-embedded section

of human laryngeal carcinoma tissue. Heat mediated

antigen retrieval was performed in EDTA buffer (pH 8.0,

epitope retrieval solution). The tissue section was blocked

with 10% goat serum. The tissue section was then incubated

with 2 ug/ml rabbit anti-Calnexin/CANX Antibody (A03372-2)

overnight at 4°C. Biotinylated goat anti-rabbit IgG was used

as secondary antibody and incubated for 30 minutes at

37°C. The tissue section was developed using Streptavidin-

Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the

chromogen.

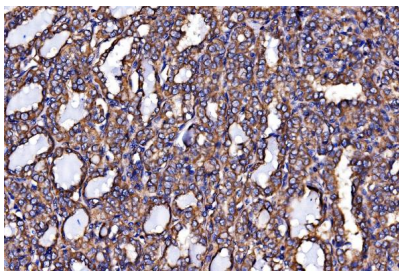


Figure 3. IHC analysis of Calnexin/CANX using anti-Calnexin/CANX antibody (A03372-2).

Calnexin/CANX was detected in a paraffin-embedded section

of human hashimoto thyroiditis tissue. Heat mediated

antigen retrieval was performed in EDTA buffer (pH 8.0,

epitope retrieval solution). The tissue section was blocked

with 10% goat serum. The tissue section was then incubated

with 2 ug/ml rabbit anti-Calnexin/CANX Antibody (A03372-2)

overnight at 4°C. Biotinylated goat anti-rabbit IgG was used

as secondary antibody and incubated for 30 minutes at

37°C. The tissue section was developed using Streptavidin-

Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the

chromogen.

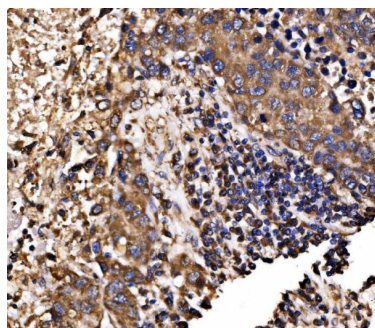


Figure 4. IHC analysis of Calnexin/CANX using anti-Calnexin/CANX antibody (A03372-2). Calnexin/CANX was detected in a paraffin-embedded section of human liver cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 ug/ml rabbit anti-Calnexin/CANX Antibody (A03372-2) overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

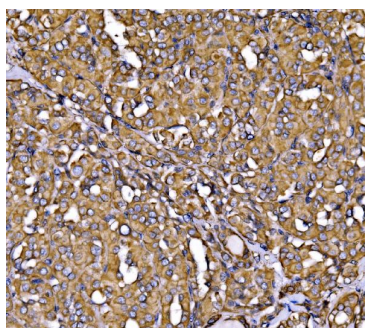


Figure 5. IHC analysis of Calnexin/CANX using anti-Calnexin/CANX antibody (A03372-2). Calnexin/CANX was detected in a paraffin-embedded section of human thyroid papillary carcinoma tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 ug/ml rabbit anti-Calnexin/CANX Antibody (A03372-2) overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

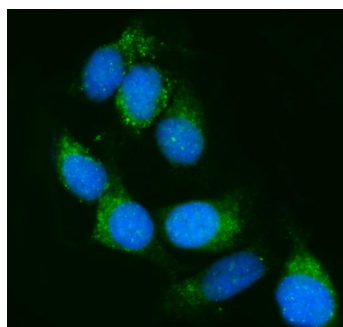


Figure 6. IF analysis of Calnexin/CANX using anti-Calnexin/CANX antibody (A03372-2). Calnexin/CANX was detected in an immunocytochemical section of Hela cells. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent (AR0022) for 15 mins. The cells were blocked with 10% goat serum. And then incubated with 5 ug/mL rabbit anti-Calnexin/CANX Antibody (A03372-2) overnight at 4°C. DyLight@488 Conjugated Goat Anti-Rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution and incubated for 30 minutes at 37°C. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.

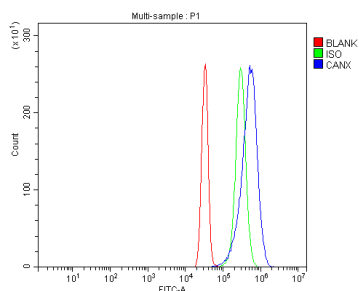


Figure 7. Flow Cytometry analysis of SiHa cells using anti-Calnexin/CANX antibody (A03372-2). Overlay histogram showing SiHa cells stained with A03372-2 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Calnexin/CANX Antibody (A03372-2, 1 ug/1x10⁶ cells) for 30 min at 20°C. DyLight@488 conjugated goat anti-rabbit IgG (BA1127, 5-10 ug/1x10⁶ cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1 ug/1x10⁶) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

1. PubMed ID: 10.1186/s13046-021-01979-7, Hypoxic tumor-derived exosomal miR-31-5p promotes lung adenocarcinoma metastasis by negatively regulating SATB2-reversed EMT and activating MEK/ERK signaling
2. PubMed ID: 10.3760/cma.j.issn.0366-6999.2011.20.023, Calcineurin is involved in cardioprotection induced by ischemic postconditioning through attenuating endoplasmic reticulum stress
3. PubMed ID: 10.1007/s10529-021-03148-4, Serum-derived exosomes accelerate scald wound healing in mice by optimizing cellular functions and promoting Akt phosphorylation

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