

Product Datasheet

Stanniocalcin 2/STC2 Antibody (orb1939920)

Catalog Number orb1939920

Category Antibodies

Description Anti-Stanniocalcin 2/STC2 Antibody. Tested in WB, IHC, ICC/IF, IF, Flow

Cytometry, ELISA applications. This antibody reacts with Human, Mouse, Rat.

Clonality Polyclonal

Species/Host Rabbit

Isotype IgG

Conjugation Unconjugated

Reactivity Human, Mouse, Rat

Form/Appearance Lyophilized

Concentration Adding 0.2 ml of distilled water will yield a concentration of 500 μg/ml.

Purification Immunogen affinity purified.

Immunogen E.coli-derived human Stanniocalcin 2/STC2 recombinant protein (Position: A51-

Q226). Human STC2 shares 90.3% and 89.8% amino acid (aa) sequence identity

with mouse and rat STC2, respectively.

UniProt ID 076061

MW 33 kDa

Tested applications ELISA, FC, ICC, IF, IHC, WB

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Application notes

Western blot, 0.25- $0.5~\mu g/ml$, Human, Mouse, Rat Immunohistochemistry, 2- $5~\mu g/ml$, Human Immunocytochemistry/Immunofluorescence, $5~\mu g/ml$, Human Immunofluorescence, $5~\mu g/ml$, Human Flow Cytometry (Fixed), 1- $3~\mu g$ /1x106 cells, Human ELISA, 0.1- $0.5~\mu g/ml$, -. Adding 0.2~ml of distilled water will yield a concentration of $500~\mu g/ml$

concentration of 50

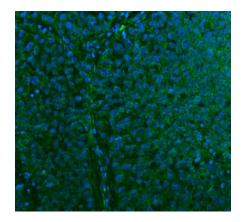
Cross Reactivity No cross reactivity with other proteins.

Antibody Type Primary Antibody

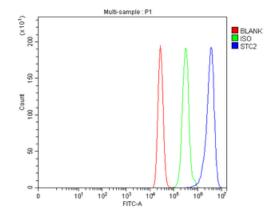
Storage Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -

20°C in small aliquots to prevent freeze-thaw cycles.

Note For research use only



IF analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human lung cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 5 μ g/mL rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. DyLight®488 Conjugated Goat Anti-Rabbit IgG was used as secondary antibody at 1:500 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.



Flow Cytometry analysis of MCF-7 cells using anti-Stanniocalcin 2/STC2 antibody. Overlay histogram showing MCF-7 cells (Blue line). The cells were fixed with 4% paraformaldehyde and blocked with 10% normal goat serum. And then incubated with rabbit anti-Stanniocalcin 2/STC2 Antibody (1 μ g/1x10^6 cells) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (5-10 μ g/1x10^6 cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1 μ g/1x10^6) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

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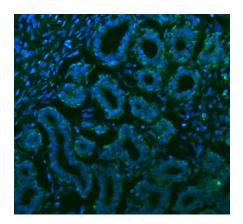
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IF analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human breast cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 5 μ g/mL rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. DyLight®488 Conjugated Goat Anti-Rabbit IgG was used as secondary antibody at 1:500 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.



IF analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in an immunocytochemical section of PC-3 cells. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent for 15 mins. The cells were blocked with 10% goat serum. And then incubated with 5 μg/mL rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. Cy3 Conjugated Goat Anti-Rabbit IgG was used as secondary antibody at 1:500 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.

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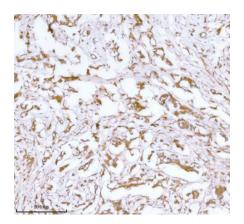
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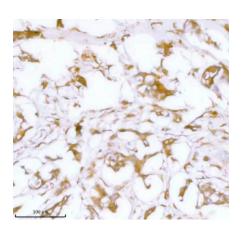
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IHC analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human breast cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Antirabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



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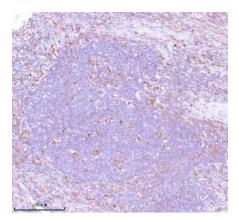
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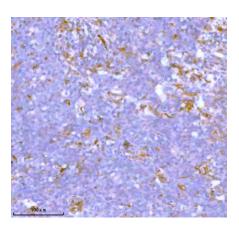
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IHC analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human lung squamous cell carcinoma tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



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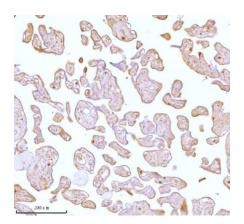
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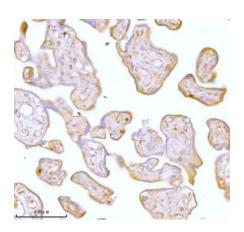
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IHC analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human placenta tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Antirabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



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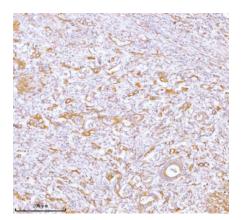
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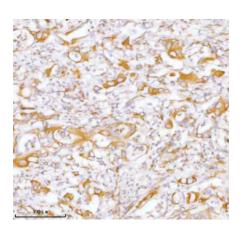
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IHC analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human rectum adenocarcinoma tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Antirabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



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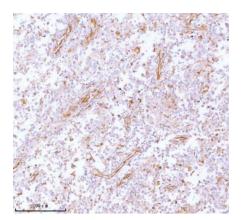
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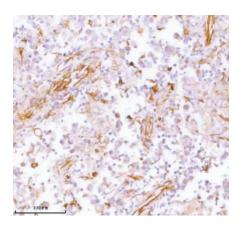
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IHC analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Stanniocalcin 2/STC2 was detected in a paraffin-embedded section of human testicular seminoma tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-Stanniocalcin 2/STC2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Antirabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit with DAB as the chromogen.



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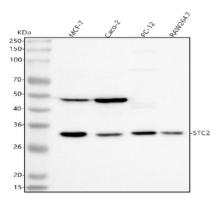
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Western blot analysis of Stanniocalcin 2/STC2 using anti-Stanniocalcin 2/STC2 antibody. Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: human MCF-7 whole cell lysates, Lane 2: human Caco-2 whole cell lysates, Lane 3: rat PC-12 whole cell lysates, Lane 4: mouse Raw264.7 whole cell 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-Stanniocalcin 2/STC2 antigen affinity purified polyclonal antibody at 0.5 μg/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 with Tanon 5200 system. A specific band was detected for Stanniocalcin 2/STC2 at approximately 33 kDa. The expected band size for Stanniocalcin 2/STC2 is at 33 kDa.

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