

Product Datasheet

Anti-NKCC1/SLC12A2 Antibody (monoclonal, 6G7D2) (orb1184753)

Catalog Number orb1184753

Description Anti-NKCC1/SLC12A2 Antibody (monoclonal, 6G7D2). Tested in IF, IHC, WB

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

Species/Host Mouse

Reactivity Human, Mouse, Rat

Conjugation Unconjugated

Tested Applications IF, IHC, WB

Immunogen E.coli-derived human NKCC1/SLC12A2 recombinant protein (Position: K889-

K943).

Form/Appearance Lyophilized

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

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

Application notes Western blot, 0.25-0.5 μg/ml, Human Immunohistochemistry(Paraffin-embedded

Section), 2-5 μ g/ml, Human, Mouse, Rat Immunofluorescence, 5 μ g/ml, Human, Mouse, Rat. Adding 0.2 ml of distilled water will yield a concentration of 500

μg/ml

Isotype Mouse IgG2a

Clonality Monoclonal

Clone Number 6G7D2

Biorbyt Ltd.

7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: info@biorbyt.com Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558 **Biorbyt LLC.**

68 TW Alexander Drive, Durham, NC, 27713, United States

Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u>
Phone: +1 (415) 906-5211 | Fax: +1 (415) 651-8558



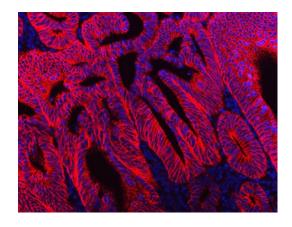


Antibody Type Primary Antibody

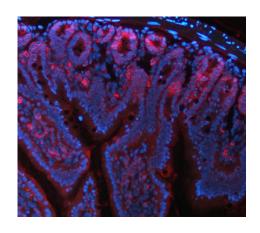
MW 200 kDa

Uniprot ID P55011

Expiration Date 12 months from date of receipt.



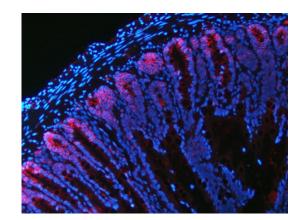
IF analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 antibody. NKCC1/SLC12A2 was detected in a paraffin-embedded section of human colorectal adenocarcinoma tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 5 µg/mL mouse anti-NKCC1/SLC12A2 Antibody overnight at 4°C. Biotin conjugated goat anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using DyLight®550 Conjugated Avidin. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.



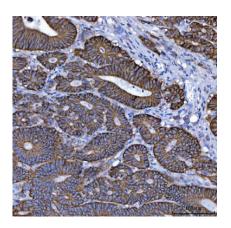
IF analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 antibody. NKCC1/SLC12A2 was detected in a paraffin-embedded section of mouse colon tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 5 µg/mL mouse anti-NKCC1/SLC12A2 Antibody overnight at 4°C. Biotin conjugated goat anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using DyLight®550 Conjugated Avidin. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.



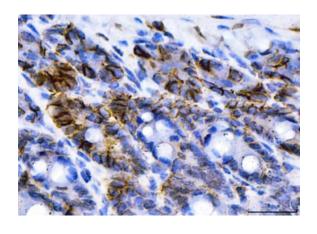




IF analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 antibody. NKCC1/SLC12A2 was detected in a paraffin-embedded section of rat colon tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 5 μg/mL mouse anti-NKCC1/SLC12A2 Antibody overnight at 4°C. Biotin conjugated goat anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using DyLight®550 Conjugated Avidin. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.



IHC analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 antibody. NKCC1/SLC12A2 was detected in a paraffin-embedded section of human colorectal 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 mouse anti-NKCC1/SLC12A2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Mouse IgG Super Vision Assay Kit with DAB as the chromogen.



IHC analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 antibody. NKCC1/SLC12A2 was detected in a paraffin-embedded section of mouse colon 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 mouse anti-NKCC1/SLC12A2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Mouse IgG Super Vision Assay Kit with DAB as the chromogen.

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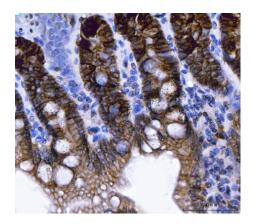
Biorbyt LLC.

68 TW Alexander Drive,
Durham, NC, 27713, United States
Email: info@biorbyt.com, support@biorbyt.com

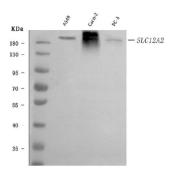
Phone: <u>+1 (415) 906-5211</u> | Fax: <u>+1 (415) 651-8558</u>







IHC analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 antibody. NKCC1/SLC12A2 was detected in a paraffin-embedded section of rat colon 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 mouse anti-NKCC1/SLC12A2 Antibody overnight at 4°C. Peroxidase Conjugated Goat Anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Mouse IgG Super Vision Assay Kit with DAB as the chromogen.



Western blot analysis of NKCC1/SLC12A2 using anti-NKCC1/SLC12A2 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 A549 whole cell lysates, Lane 2: human CACO-2 whole cell lysates, Lane 3: human PC-3 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 mouse anti-NKCC1/SLC12A2 antigen affinity purified monoclonal 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-mouse IgG-HRP secondary antibody at a dilution of 1:10000 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 NKCC1/SLC12A2 at approximately 200 kDa. The expected band size for NKCC1/SLC12A2 is at 131 kDa.

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