Product Datasheet

SLC31A1/CTR1 Antibody - BSA Free NB100-402

Unit Size: 0.1 ml

Store at 4C short term. Store at -20C long term. Avoid freeze-thaw cycles.

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NB100-402

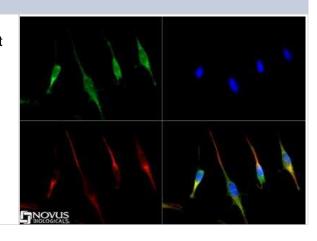
SLC31A1/CTR1 Antibody - BSA Free

SECSTATION Antibody - BSA Free	
Product Information	
Unit Size	0.1 ml
Concentration	1 mg/ml
Storage	Store at 4C short term. Store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.02% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	PBS
Product Description	
Host	Rabbit
Gene ID	1317
Gene Symbol	SLC31A1
Species	Human, Mouse, Rat, Porcine, Canine, Xenopus, Zebrafish
Reactivity Notes	Canine reactivity reported in scientific literature (PMID: 29901089).
Immunogen	A synthetic peptide derived from a C-terminal sequence of human SLC31A1/CTR1 [Uniprot: O15431]
Product Application Details	
Applications	Western Blot, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Frozen, Immunohistochemistry-Paraffin
Recommended Dilutions	Western Blot, Immunohistochemistry 1:250, Immunocytochemistry/ Immunofluorescence 1:500, Immunohistochemistry-Paraffin 1:250,

Immunohistochemistry-Frozen

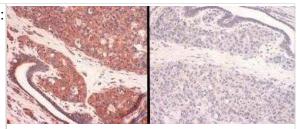
Images

Immunocytochemistry/Immunofluorescence: SLC31A1/CTR1 Antibody [NB100-402] - CTR1 antibody was tested in NIH-3T3 cells with DyLight 488 (green). Nuclei and alpha-tubulin were counterstained with DAPI (blue) and DyLight 550 (red).

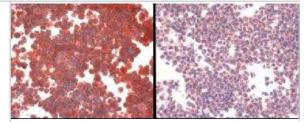




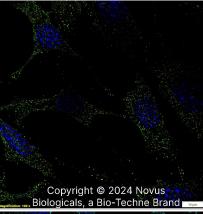
Immunohistochemistry: SLC31A1/CTR1 Antibody [NB100-402] - Panel 1: human CTR1 staining of breast cancer tissue. Panel 2: human CTR1-antigen competition in breast cancer tissue.



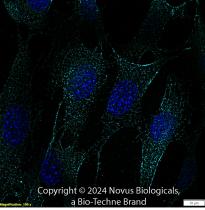
Immunohistochemistry: SLC31A1/CTR1 Antibody [NB100-402] - Left panel: Antibody staining of hCTR1 overexpressing cell line. Right panel: Antibody plus peptide competition staining.



SLC31A1/CRT1 was detected in immersion fixed NIH3T3 Mouse fibroblast cell line using Rabbit anti-SLC31A1/CRT1 Antigen Affinity-purified Polyclonal Antibody (Catalog # NB100-402) at 2 µg/mL overnight at 4C. Cells were stained using DyLight 488-conjugated Anti-Rabbit IgG (H+L) Cross-Absorbed Secondary Antibody (green) and counterstained with DAPI (blue). Cells were imaged using a 100X objective and digitally deconvolved.



SLC31A1/CRT1 was detected in immersion fixed NIH3T3 Mouse fibroblast cell line using Rabbit anti-SLC31A1/CRT1 Antigen Affinity-purified Polyclonal Antibody conjugated to DyLight 650 (Catalog # NB100-402C) (light blue) at 10 µg/mL overnight at 4C. Cells were counterstained with DAPI (blue). Cells were imaged using a 100X objective and digitally deconvolved.



Publications

Federica Iannelli, Andrea Ilaria Zotti, Maria Serena Roca, Laura Grumetti, Rita Lombardi, Tania Moccia, Carlo Vitagliano, Maria Rita Milone, Chiara Ciardiello, Francesca Bruzzese, Alessandra Leone, Ernesta Cavalcanti, Rossella De Cecio, Giuseppina Iachetta, Salvatore Valiante, Franco Ionna, Francesco Caponigro, Elena Di Gennaro, Alfredo Budillon Valproic Acid Synergizes With Cisplatin and Cetuximab in vitro and in vivo in Head and Neck Cancer by Targeting the Mechanisms of Resistance Frontiers in Cell and Developmental Biology 2020-08-17 [PMID: 33015030]

Elodie A. Pérès, Jérôme Toutain, Louis-Paul Paty, Didier Divoux, Méziane Ibazizène, Stéphane Guillouet, Louisa Barré, Aurélien Vidal, Michel Cherel, Mickaël Bourgeois, Myriam Bernaudin, Samuel Valable 64 Cu-ATSM/ 64 Cu-Cl 2 and their relationship to hypoxia in glioblastoma: a preclinical study EJNMMI Research 2019-12-19 [PMID: 31858290]

Fantin J, Toutain J, Pérès EA et al. Assessment of hypoxia and oxidative-related changes in a lung-derived brain metastasis model by [64Cu][Cu(ATSM)] PET and proteomic studies EJNMMI research 2023-11-25 [PMID: 38006431] (IHC-Fr, Human)

Schoonover KE, Queern SL, Lapi SE, Roberts RC. Impaired copper transport in schizophrenia results in a copperdeficient brain state: A new side to the dysbindin story The World Journal of Biological Psychiatry 2020-01-02 [PMID: 30230404] (Western Blot)

Choi H, Oh D, Kim M et al. Copper deficiency affects the developmental competence of porcine oocytes matured in vitro Frontiers in Cell and Developmental Biology 2022-09-07 [PMID: 36158185] (Immunocytochemistry/Immunofluorescence)

Shin VY, Liu MX, Siu JM et al. Inhibition of EP2 receptor suppresses tumor growth and chemoresistance of gastric cancer American journal of cancer research 2022-10-15 [PMID: 36381319] (WB, Human)

Zhang X, Jiang Q, Su Y et al. AMPK phosphorylates and stabilises copper transporter 1 to synergise metformin and copper chelator for breast cancer therapy British journal of cancer 2023-02-17 [PMID: 36807336]

Saifi MA, Godugu C Copper chelation therapy inhibits renal fibrosis by modulating copper transport proteins BioFactors (Oxford, England) 2022-03-24 [PMID: 35322483]

Aloysius Dhivya M, Sulochana K, Devi S High glucose induced inflammation is inhibited by copper chelation via rescuing mitochondrial fusion protein 2 in retinal pigment epithelial cells Cellular Signalling 2022-01-01 [PMID: 34999205] (WB, Human)

Liu LL, Du D, Zheng W, Zhang Y Age-dependent Decline of Copper Clearance at the Blood-Cerebrospinal Fluid Barrier Neurotoxicology 2021-10-27 [PMID: 34718061] (IF/IHC, WB, Rat)

Guo J, Cheng J, Zheng N Et Al. Copper Promotes Tumorigenesis by Activating the PDK1-AKT Oncogenic Pathway in a Copper Transporter 1 Dependent Manner Advanced science (Weinheim, Baden-Wurttemberg, Germany) 2021-07-18 [PMID: 34278744]

Schoonover KE, Kennedy WM, Roberts RC Cortical copper transporter expression in schizophrenia: interactions of risk gene dysbindin-1 Journal of neural transmission (Vienna, Austria: 1996) 2021-04-22 [PMID: 33890175] (WB, Human)

More publications at http://www.novusbio.com/NB100-402



Procedures

Immunocytochemistry/Immunofluorescence Protocol for SLC31A1/CTR1 Antibody (NB100-402) Immunocytochemistry Protocol

Culture cells to appropriate density in 35 mm culture dishes or 6-well plates.

- 1. Remove culture medium and wash the cells briefly in PBS. Add 10% formalin to the dish and fix at room temperature for 10 minutes.
- 2. Remove the formalin and wash the cells in PBS.
- 3. Permeablize the cells with 0.1% Triton X100 or other suitable detergent for 10 min.
- 4. Remove the permeablization buffer and wash three times for 10 minutes each in PBS. Be sure to not let the specimen dry out.
- 5. To block nonspecific antibody binding, incubate in 10% normal goat serum from 1 hour to overnight at room temperature.
- 6. Add primary antibody at appropriate dilution and incubate overnight at 4C.
- 7. Remove primary antibody and replace with PBS. Wash three times for 10 minutes each.
- 8. Add secondary antibody at appropriate dilution. Incubate for 1 hour at room temperature.
- 9. Remove secondary antibody and replace with PBS. Wash three times for 10 minutes each.
- 10. Counter stain DNA with DAPi if required.

Western Blot Protocol for SLC31A1/CTR1 Antibody (NB100-402)

Western Blot Protocol

- 1. Perform SDS-PAGE on samples to be analyzed, loading 10-25 up of total protein per lane.
- 2. Transfer proteins to PVDF membrane according to the instructions provided by the manufacturer of the membrane and transfer apparatus.
- 3. Stain the membrane with Ponceau S (or similar product) to assess transfer success, and mark molecular weight standards where appropriate.
- 4. Rinse the blot TBS -0.05% Tween 20 (TBST).
- 5. Block the membrane in 5% Non-fat milk in TBST (blocking buffer) for at least 1 hour.
- 6. Wash the membrane in TBST three times for 10 minutes each.
- 7. Dilute primary antibody in blocking buffer and incubate overnight at 4C with gentle rocking.
- 8. Wash the membrane in TBST three times for 10 minutes each.
- 9. Incubate the membrane in diluted HRP conjugated secondary antibody in blocking buffer (as per manufacturer's instructions) for 1 hour at room temperature.
- 10. Wash the blot in TBST three times for 10 minutes each (this step can be repeated as required to reduce background).
- 11. Apply the detection reagent of choice in accordance with the manufacturer's instructions.



Immunohistochemistry-paraffin protocol specific for SLC31A1 / CTR1 Antibody (NB100-402) Immunohistochemistry-Paraffin Embedded Sections

Antigen Unmasking:

Bring slides to a boil in 10 mM sodium citrate buffer (pH 6.0) then maintain at a sub-boiling temperature for 10 minutes. Cool slides on bench-top for 30 minutes (keep slides in the sodium citrate buffer at all times).

Staining:

- 1. Wash sections in deionized water three times for 5 minutes each.
- 2. Wash sections in PBS for 5 minutes.
- 3. Block each section with 100-400 ul blocking solution (1% BSA in PBS) for 1 hour at room temperature.
- 4. Remove blocking solution and add 100-400 ul diluted primary antibody. Incubate overnight at 4 C.
- 5. Remove antibody solution and wash sections in wash buffer three times for 5 minutes each.
- 6. Add 100-400 ul HRP polymer conjugated secondary antibody. Incubate 30 minutes at room temperature.
- 7. Wash sections three times in wash buffer for 5 minutes each.
- 8. Add 100-400 ul DAB substrate to each section and monitor staining closely.
- 9. As soon as the sections develop, immerse slides in deionized water.
- 10. Counterstain sections in hematoxylin.
- 11. Wash sections in deionized water two times for 5 minutes each.
- 12. Dehydrate sections.
- 13. Mount coverslips.





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Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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