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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

Mouse anti Human CD56

exalpha.com/products/mouse-anti-human-cd56/0561

Catalog number: **0561**

Clone	C5.9
Isotype	IgG2b
Product Type	Monoclonal Antibody
Units	250 µg
Host	Mouse
Species Reactivity	Human
Application	Flow Cytometry Immunocytochemistry Immunohistochemistry (paraffin)

Background

NCAM / CD56, as a member of the immunoglobulin superfamily of adhesion molecules is characterized by several immunoglobulin (Ig)-like domains. The extracellular part of NCAM consists of five of these Ig domains and two fibronectin type III homology regions. NCAM is encoded by a single copy gene composed of 26 exons. However, at least 20-30 distinct isoforms can be generated by alternative splicing and by posttranslational modifications, such as sialylation. During sialylation, polysialic acid (PSA) carbohydrates are attached to the extracellular part of NCAM. Through its extracellular region, NCAM mediates homophilic interactions. In addition, NCAM can also undergo heterophilic interactions by binding extracellular matrix components, such as laminin, or other cell adhesion molecules, such as integrins. NCAM can be found in central and peripheral nerve cells, neuroendocrine tissues and at the surface of NK-cells. Also, NCAM is present in malignancies derived from these tissues and cells.

Synonyms: CD56, NCAM, Neural Cell Adhesion Molecule

Source

Derived from the hybridization of mouse Sp2/0 myeloma cells with spleen cells from BALB/c F1 mice immunized with an extract of the KG1a cell line.

Immunogen: KG1a cell line extract.

Product

Purified monoclonal antibody clone C5.9, filtered, in PBS containing 0.08% azide

Product Form: Unconjugated

Purification Method: ProteinG Chromatography

Concentration: See vial for concentration

Applications

Antibody can be used for Immunohistochemistry (2-5 µg/ml, formalin-fixed, paraffin embedded tissues). Optimal concentration should be evaluated by serial dilutions.

Functional Analysis: Flow Cytometry

Storage

Product should be stored at -20°C. Aliquot to avoid freeze/thaw cycles

Product Stability: Reagents are stable for the period shown on the vial label when stored properly

Shipping Conditions: Ship at ambient temperature, freeze upon arrival

Caution

This product is intended FOR RESEARCH USE ONLY, and FOR TESTS IN VITRO, not for use in diagnostic or therapeutic procedures involving humans or animals. It may contain hazardous ingredients. Please refer to the Safety Data Sheets (SDS) for additional information and proper handling procedures. Dispose product remainders according to local regulations. This datasheet is as accurate as reasonably achievable, but Exalpha Biologicals accepts no liability for any inaccuracies or omissions in this information.

References

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6. Muench, Marcus O., et al. Isolation of Definitive Zone and Chromaffin Cells Based upon Expression of CD56 (Neural Cell Adhesion Molecule) in the Human Fetal Adrenal Gland,

J Clinical Endocrinology & Metabolism. 2003, 88: 3921-3930, PRODUCT SPECIFIC REFERENCES 1. Horikawa, M., et al. Abnormal Natural Killer Cell Function in Systemic Sclerosis: Altered Cytokine Production and Defective Killing Activity. Journal of Investigative Dermatology 2005, 125: 731-737 2. Ishimoto, H., et al. Midkine, a heparin-binding growth factor, selectively stimulates proliferation of definitive zone cells of the human fetal adrenal gland. Journal of Endocrinol. Metab 2006, 91: 4050-4056 3. Muench, M., et al. Isolation of Definitive Zone and Chromaffin Cells Based upon Expression of CD56 (Neural Cell Adhesion Molecule) in the Human Fetal Adrenal Gland. Journal of Clinical Endocrinol. Metab 2003, 88: 3921-3930 4. Miles, L.A., et al. Cell-Surface Actin Binds Plasminogen and Modulates Neurotransmitter Release from Catecholaminergic Cells. Journal of Neuroscience 2006, 26: 13017-13024

Protein Reference(s)

Database Name: UniProt

Accession Number: P13591

IHC: Formalin-fixed, paraffin-embedded human neuroblastoma stained with Exalpha's CD56, clone C5.9 at 2 ug/ml using peroxidase-conjugate and DAB chromogen. Cell membrane staining of tumor cells is clearly evident. High-temperature heating was used as the retrieval protocol, pressure cooker with citrate buffer, pH 6.5, for 10 min. Exalpha's anti-CD56, clone C5.9, was added at 2 ug/ml for 1 hour at room temperature followed by anti-Mouse HRP for 30 minutes at room temperature. Staining was visualized using DAB and methyl green counter stain. Flow Cytometry: Flow cytometry data using Exalpha's Mouse anti Human CD56 antibody 0561 at 8 µg/ml on human lymphocytes, with Rat anti Mouse PE IgG2a & 2b as a second step. The negative control was generated without the CD56 antibody and without the Rat anti Mouse PE IgG2a & 2b. Western Blot: Western blot analysis using Exalpha's Mouse anti Human CD56 antibody 0561 at 10 µg/ml (left lane) and 1 µg/ml (right lane) showing a strong reaction on recombinant human NCAM-1/CD56 protein.