

Basic Information

Product Name	Anti-CD56/NCAM1 Antibody		
Gene Name	NCAM1		
Source	Rabbit		
Clonality	Polyclonal		
Isotype	IgG		
Species Reactivity	human, mouse, rat		
Tested Application	WB, ICC/IF, FCM, ELISA		
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.		
Immunogen	E.coli-derived human NCAM1 recombinant protein (Position: T80-T328).		
Concentration	500 ug/ml		
Purification	Immunogen affinity purified.		
Observed MW	120-150 kDa		
Dilution Ratios	Western blot (WB):	1:500-2000	
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400	
	Flow Cytometry (Fixed):	1:50-200	
	Enzyme linked immunosorbent assay (ELISA):	1:100-1000	

Storage

12 months from date of receipt, -20°C as supplied.

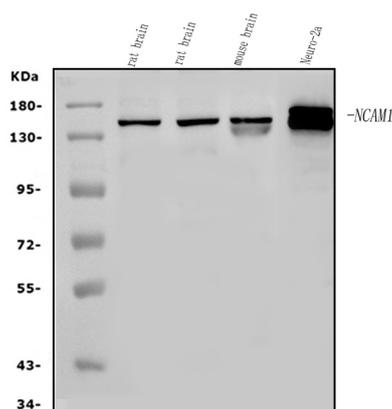
Background Information

NCAM is a membrane-bound glycoprotein that plays a role in cell-cell and cell-matrix adhesion through both its homophilic and heterophilic binding activity. The neural cell adhesion molecule appears on early embryonic cells and is important in the formation of cell collectives and their boundaries at sites of morphogenesis. Later in development it is found on various differentiated tissues and is a major CAM mediating adhesion among neurons and between neurons and muscle. NCAM gene is mapped to 11q23. The neural cell adhesion molecule (NCAM) can influence a number of diverse intercellular events, including junctional communication, the association of axons with pathways and targets, and signals that alter levels of neurotransmitter enzymes.

Reference

Anti-CD56/NCAM1 Antibody被引用在3文献中。

Selected Validation Data



Western blot analysis of CD56/NCAM1 using anti-CD56/NCAM1 antibody (A00184-4). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: rat brain tissue lysates,

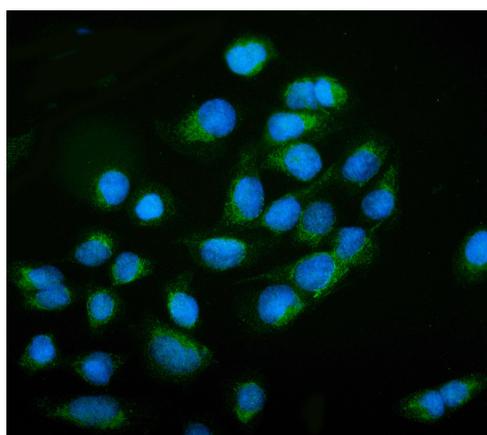
Lane 2: rat brain tissue lysates,

Lane 3: mouse brain tissue lysates,

Lane 4: Mouse Neuro-2a whole cell lysates.

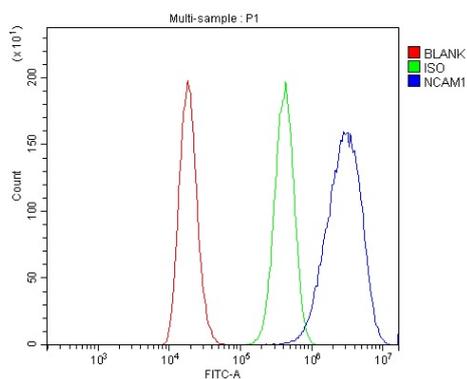
After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-CD56/NCAM1 antigen affinity purified polyclonal antibody (A00184-4) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for CD56/NCAM1 at approximately 120-150 kDa. The expected band size for CD56/NCAM1 is at 95 kDa.



ICC/IF analysis of CD56/NCAM1 using anti-CD56/NCAM1 antibody (A00184-4).

CD56/NCAM1 was detected in an immunocytochemical section of U2OS cells. The section was incubated with rabbit anti-CD56/NCAM1 Antibody (A00184-4) at a dilution of 1:100. Fluoro488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of 293T cells using anti-CD56/NCAM1 antibody (A00184-4).

Overlay histogram showing 293T cells stained with A00184-4 (Blue line). The cells were fixed with 4% paraformaldehyde and blocked with 10% normal goat serum. And then incubated with rabbit anti-CD56/NCAM1 Antibody (A00184-4) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.