

Datasheet for ABIN6160049

anti-CD56 antibody (CF®488A)



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Quantity:	100 μL
Target:	CD56 (NCAM1)
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This CD56 antibody is conjugated to CF®488A
Application:	Flow Cytometry (FACS), Western Blotting (WB), Immunofluorescence (IF), Immunohistochemistry (Formalin-fixed Sections) (IHC (f))

Product Details

Purpose:	Mouse Monoclonal anti-CD56 / NCAM (NCAM1/795), CF488A Conjugate		
Immunogen:	Recombinant human NCAM1 protein		
Clone:	NCAM1-795		
Isotype:	IgG1 kappa		
Characteristics:	This MAb reacts with an extracellular domain (close to transmembrane) of CD56/NCAM. Three		

isoforms of neural cell adhesion molecule (NCAM) are produced by differential splicing of the RNA transcript from a single gene. The 135 kDa isoform is the basic molecule, which is glycosylated or sialylated to produce the mature species. Anti-CD56 recognizes two proteins of the neural cell adhesion molecule, the basic molecule expressed on most neuroectodermally derived tissues and neoplasms (e.g. retinoblastoma, medulloblastomas, astrocytomas, neuroblastomas, and small cell carcinomas). It is also expressed on some mesodermally

derived tumors (rhabdomyosarcoma). Anti-CD56 plays an important role in the diagnosis of nodal and nasal NK/T-cell lymphomas. Primary antibodies are available purified, or with a selection of fluorescent CF® dyes and other labels. CF® dyes offer exceptional brightness and photostability. Note: Conjugates of blue fluorescent dyes like CF®405S and CF®405M are not recommended for detecting low abundance targets, because blue dyes have lower fluorescence and can give higher non-specific background than other dye colors.

Target Details

Target:	CD56 (NCAM1)
Alternative Name:	CD56 / NCAM (NCAM1 Products)
Background:	NCAM, Leu-19, NKH1, MSK39, NCAM120, NCAM140, NCAM180, Neural Cell Adhesion Molecule
Molecular Weight:	180, 145 and 125 kDa
Gene ID:	4684, 503878

Application Details

Application N	lotes:
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Immunohistology formalin-fixed 0.5-1 µg/mL

- Staining of formalin-fixed tissues requires boiling tissue sections in 10 mM citrate buffer, pH
 6.0, for 10-20 min followed by cooling at RT for 20 minutes
- Flow Cytometry 0.5-1 μg/million cells/0.1 mL
- Immunofluorescence 1-2 μg/mL
- Western blotting 0.5-1 μg/mL
- · Optimal dilution for a specific application should be determined by user

Comment:

Cerebellum, Pancreas, Neuroblastoma

Restrictions:

For Research Use only

Handling

Format:	Liquid
Concentration:	100 μg/mL
Buffer:	PBS/0.1 % BSA/0.05 % azide
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

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Handling Advice:

Protect from light