

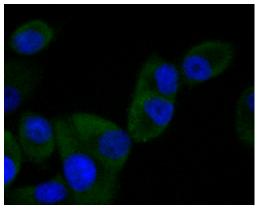


SV2B Monoclonal Antibody (A4C1)

Product Details		
Size	100 μL	
Species Reactivity	Human, Rat	
Host/Isotype	Mouse / IgG1	
Class	Monoclonal	
Туре	Antibody	
Clone	A4C1	
Conjugate	Unconjugated	
Immunogen	Recombinant protein within human SV2B aa 400-600	
Form	Liquid	
Concentration	2 mg/mL	
Purification	Protein G	
Storage buffer	TBS, pH 7.4, with 50% glycerol, 0.2% BSA	
Contains	0.05% sodium azide	
Storage conditions	Store at 4°C short term. For long term storage, store at -20°C, avoiding freeze/thaw cycles.	
RRID	AB_2898837	

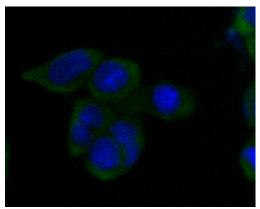
Applications	Tested Dilution	Publications
Western Blot (WB)	1:500-1:1,000	-
Immunohistochemistry (Paraffin) (IHC (P))	1:50-1:200	-
Immunocytochemistry (ICC/IF)	1:50-1:100	-
Flow Cytometry (Flow)	1:50-1:100	-

Product Images For SV2B Monoclonal Antibody (A4C1)



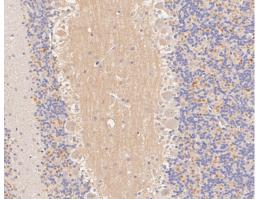
SV2B Antibody (MA5-41083) in ICC/IF

Immunocytochemistry/Immunofluorescence analysis of SV2B in PANC-1 cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with SV2B monoclonal Antibody (Product # MA5-41083) at a dilution of 1:50 for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Mouse IgG was used as the secondary antibody at 1:1,000 dilution. The nuclear counter stain is DAPI (blue).



SV2B Antibody (MA5-41083) in ICC/IF

Immunocytochemistry/Immunofluorescence analysis of SV2B in F9 cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with SV2B monoclonal Antibody (Product # MA5-41083) at a dilution of 1:50 for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Mouse IgG was used as the secondary antibody at 1:1,000 dilution. The nuclear counter stain is DAPI (blue).



SV2B Antibody (MA5-41083) in IHC (P)

Immunohistochemistry (Paraffin) analysis of SV2B in paraffin-embedded rat cerebellum tissue. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with SV2B monoclonal Antibody (Product # MA5-41083) at a dilution of 1:100 for 30 minutes at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with

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