

Basic Information

Product Name	Anti-CD2AP Antibody (Clone#5F8)	
Gene Name	CD2AP	
Source	Mouse	
Clonality	Monoclonal	
Isotype	IgG1	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, ICC/IF, FCM	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	E. coli-derived human CD2AP recombinant protein (Position: K253-K337).	
Concentration	500 ug/ml	
Purification	protein G purified.	
Observed MW	80 kDa	
Dilution Ratios	Western blot (WB):	1:500-2000
	Immunohistochemistry (IHC):	1:50-400
	Flow Cytometry (Fixed):	1:50-200
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400
	(Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 25 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

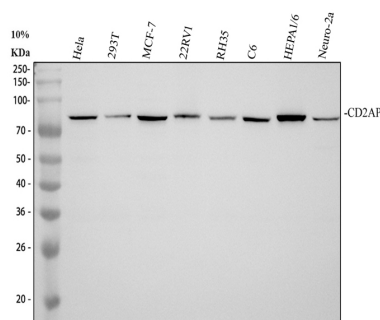
Storage

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

Background Information

CD2-associated protein is a protein that in humans is encoded by the CD2AP gene. This gene encodes a scaffolding molecule that regulates the actin cytoskeleton. The protein directly interacts with filamentous actin and a variety of cell membrane proteins through multiple actin binding sites, SH3 domains, and a proline-rich region containing binding sites for SH3 domains. The cytoplasmic protein localizes to membrane ruffles, lipid rafts, and the leading edges of cells. It is implicated in dynamic actin remodeling and membrane trafficking that occurs during receptor endocytosis and cytokinesis. Haploinsufficiency of this gene is implicated in susceptibility to glomerular disease.

Selected Validation Data



Western blot analysis of CD2AP using anti-CD2AP antibody

(M01756). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human Hela whole cell lysates,

Lane 2: human 293T whole cell lysates,

Lane 3: human MCF-7 whole cell lysates,

Lane 4: human 22RV1 whole cell lysates,

Lane 5: rat RH35 whole cell lysates,

Lane 6: rat C6 whole cell lysates,

Lane 7: mouse Hepa1/6 whole cell lysates,

Lane 8: mouse Neuro-2a whole cell lysates.

After electrophoresis, proteins were transferred to a membrane.

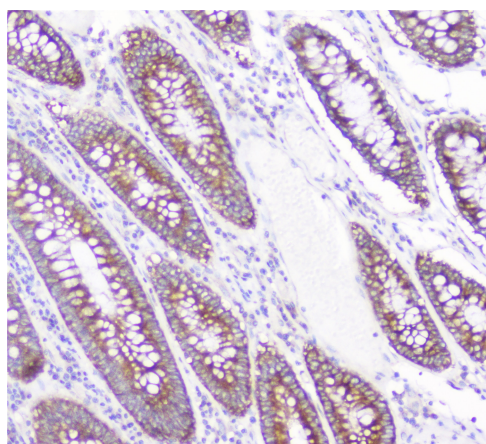
Then the membrane was incubated with mouse anti-CD2AP antigen

affinity purified monoclonal antibody (M01756) at a dilution of

1:1000 and probed with a goat anti-mouse IgG-HRP secondary antibody (Catalog # BA1050). The signal is developed using ECL Plus

Western Blotting Substrate (Catalog # AR1197). A specific band was detected for CD2AP at approximately 80 kDa. The expected band

size for CD2AP is at 71 kDa.



IHC analysis of CD2AP using anti-CD2AP antibody (M01756).

CD2AP was detected in a paraffin-embedded section of human colon

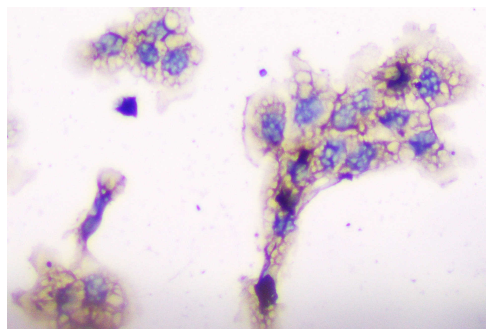
cancer tissue. Biotinylated goat anti-mouse IgG was used as

secondary antibody. The tissue section was incubated with mouse

anti-CD2AP Antibody (M01756) at a dilution of 1:200 and developed

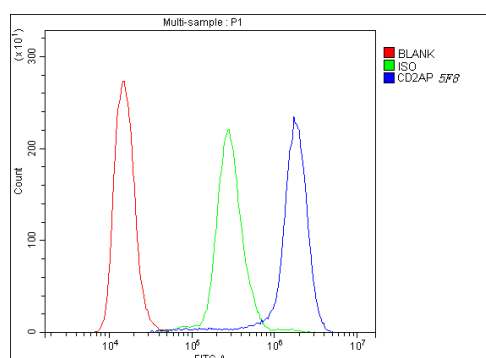
using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1021) with

DAB (Catalog # AR1027) as the chromogen.



ICC analysis of CD2AP using anti- CD2AP antibody (M01756).

CD2AP was detected in an immunocytochemical section of A431 cells. The section was incubated with mouse anti-CD2AP Antibody (M01756) at a dilution of 1:100. Biotinylated goat anti-mouse IgG was used as secondary antibody. The section was developed using Streptavidin-Biotin-Complex (SABC)(Catalog # SA1021) with DAB (Catalog # AR1027) as the chromogen.



Flow Cytometry analysis of K562 cells using anti-D2AP antibody (M01756). Overlay histogram showing K562 cells stained with M01756 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with mouse anti-D2AP Antibody (M01756, 1:100) for 30 min at 20°C. Fluoro®488 conjugated goat anti-mouse IgG (BA1126, 1:100) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was mouse IgG (1:100) used under the same conditions. Unlabelled sample (Red line) was also used as a control.