

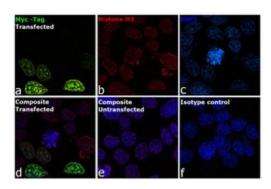


c-Myc Monoclonal Antibody (9E10), Alexa Fluor 488

Product Details	
Size	50 μL
Species Reactivity	Human
Published Species	Human
Host/Isotype	Mouse / IgG1
Class	Monoclonal
Туре	Antibody
Clone	9E10
Conjugate	Alexa Fluor® 488
Immunogen	Synthetic peptide A(408) E E Q K L I S E E D L L R K R R E Q L K H K L E Q L R N S C A(438) of human c-Myc
Form	Liquid
Concentration	1 mg/mL
Purification	Protein A
Storage buffer	PBS with 1mg/mL BSA
Contains	0.05% sodium azide
Storage conditions	4° C, do not freeze
RRID	AB_2609819

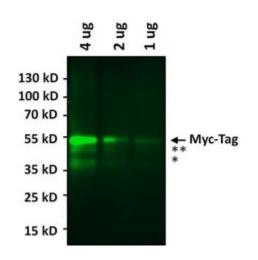
Applications	Tested Dilution	Publications
Western Blot (WB)	1:500	-
Immunocytochemistry (ICC/IF)	1:100	1 Publication

Product Images For c-Myc Monoclonal Antibody (9E10), Alexa Fluor 488



c-Myc Antibody (MA1-980-A488) in ICC/IF

Immunofluorescence analysis of Myc proto-oncogene protein was performed using 70% confluent log phase HEK-293 cells. The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 15 minutes, and blocked with 2% BSA for 45 minutes at room temperature. The cells were labeled with Myc Tag Polyclonal Antibody (Product # MA1-980-A488) at 1:100 in 0.1% BSA and Histone H3 antibody (Product # 711055) at 1:200 dilution in 0.1% BSA, incubated at 4 degree Celsius overnight, and then labeled with Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 647 (Product # A32733) at a dilution of 1:2000 for 45 minutes at room temperature. Panel a (Nuclei: Green) represents the Myc tag. Panel b (Nuclei: Red) represents Histone H3. Panel c (Nuclei: Blue) represents ProLong™ Diamond Antifade Mountant with DAPI (Product # P36962). Panel d represents the merged image showing the colocalization of nuclear signals in transfected cells. Panel e represents un-transfected HEK-293 cells. Panel f represents isotype control cells to assess background. The images were captured at 60X magnification.



c-Myc Antibody (MA1-980-A488) in WB

Western blot analysis of Myc Epitope Tag was performed by loading various amounts of E. coli lysate containing a multi-epitope tagged protein per well onto a 4-20% Tris-HCl polyacrylamide gel. Proteins were transferred to a low fluorescence PVDF membrane and blocked with Sea Block blocking buffer for at least 1 hour. The membrane was probed with a AlexaFluor 488-conjugated Myc Epitope Tag monoclonal antibody (Product # MA1-980-A488) at a dilution of 1:500 for 1 hour at room temperature on a rocking platform and washed in TBS-0.1% Tween-20. Detection was performed using a fluorescence imaging system.

□ 1 Reference

Immunocytochemistry (1)

Molecular biology of the cell

N-cadherin association with lipid rafts regulates its dynamic assembly at cell-cell junctions in C2C12 myoblasts.

"MA1-980-A488 was used in Immunocytochemistry-immunoflourescence to show that lipid rafts, as homophilic interaction and F-actin association, stabilize cadherin-dependent adhesive complexes."

Authors: Causeret M, Taulet N, Comunale F, Favard C, Gauthier-Rouvière C

Species Human

Dilution 1:2,000

Year 2005

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