# Anti-ICAM1 Antibody [J0-F7]

## M1511-6



Mouse monoclonal IgG2c, primary antibodies
Human
WB, IF-Cell, FC
58 kDa
J0-F7
This gene encodes a cell surface glycoprotein which is typically expressed on endothelial cells and cells of the immune system. It binds to integrins of type CD11a / CD18, or CD11b / CD18 and is also exploited by Rhinovirus as a receptor.
Recombinant protein within human ICAM1 aa 30-300.
Raji cell lysates, Daudi cell lysates, Hela, HUVEC, Jurkat.
Membrane.
SwissProt: P05362 Human
1:500-1:2,000
1:50-1:100
1:50-1:100
1*PBS (pH7.4), 0.2% BSA, 40% Glycerol. Preservative: 0.05% Sodium Azide.
Store at +4 $^\circ\!{\rm C}$ after thawing. Aliquot store at -20 $^\circ\!{\rm C}$ or -80 $^\circ\!{\rm C}$ . Avoid repeated freeze / thaw cycles.
Protein A affinity purified.

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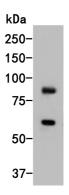
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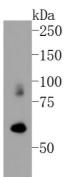
Applications:WB=Western blot IHC-P=Immunohistochemistry (paraffin) IF-Cell=Immunofluorescence (Cell) IF-Tissue=Immunofluorescence (Tissue) FC=Flow cytometry IP=Immunoprecipitation

#### M1511-6 - Page 2

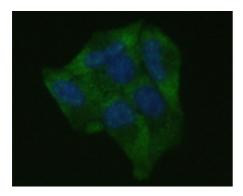
#### Images



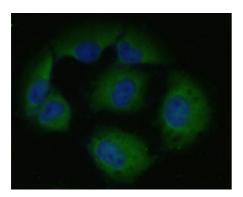
**Fig1:** Western blot analysis of ICAM1 on Raji cell lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (M1511-6, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Mouse IgG - HRP Secondary Antibody (HA1006) at 1:5,000 dilution was used for 1 hour at room temperature.



**Fig2:** Western blot analysis of ICAM1 on Daudi cell lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (M1511-6, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Mouse IgG - HRP Secondary Antibody (HA1006) at 1:5,000 dilution was used for 1 hour at room temperature.



**Fig3:** ICC staining of ICAM1 in Hela 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 the primary antibody (M1511-6, 1/100) 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).



**Fig4:** ICC staining of ICAM1 in HUVEC 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 the primary antibody (M1511-6, 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).

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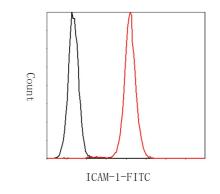
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**Fig5:** Flow cytometric analysis of ICAM1 was done on Jurkat cells. The cells were fixed, permeabilized and stained with the primary antibody (M1511-6, 1/100) (red). After incubation of the primary antibody at room temperature for an hour, the cells were stained with a FITC-conjugated Goat anti-Mouse IgG Secondary antibody at 1/200 dilution for 30 minutes.Unlabelled sample was used as a control (cells without incubation with primary antibody; black).

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

#### **Background References**

- 1. Marzolla V. et. al. Essential role of ICAM-1 in aldosterone-induced atherosclerosis. Int J Cardiol. 2017 Apr 1;232:233-242.
- 2. Othumpangat S. et. al. ICAM-1 regulates the survival of influenza virus in lung epithelial cells during the early stages of infection. Virology. 2016 Jan;487:85-94.

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