Product datasheet

Anti-Integrin alpha V antibody [P2W7] ab11470

Overview

Product name: Anti-Integrin alpha V antibody [P2W7]
Description: Mouse monoclonal [P2W7] to Integrin alpha V
Host species: Mouse
Tested applications: Suitable for: IP, ICC/IF, WB, Flow Cyt
Species reactivity: Reacts with: Human
Immunogen: An ocular melanoma cell line expressing high levels of human integrin alpha V beta 1.
Positive control: Tested with squamous cell carcinoma cells. A375M melanoma cell line. A549 cell line IF/ICC
General notes: Human alpha v is about 150-160kDa under non-reducing conditions. Under reducing conditions it is cleaved into 2 bands of about 125-130kDa +20-25kDa.

Properties

Form: Liquid
Storage instructions: Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -20°C or -80°C. Avoid freeze / thaw cycle.
Storage buffer: Constituent: PBS
Purity: Protein A purified
Purification notes: The antibody is Protein A purified from tissue culture supernatant.
Clonality: Monoclonal
Clone number: P2W7
Isotype: IgG1

Applications

Our Abpromise guarantee covers the use of ab11470 in the following tested applications.
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

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<td>Use at an assay dependent concentration.</td>
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**Function**
The alpha-V integrins are receptors for vitronectin, cytotactin, fibronectin, fibrinogen, laminin, matrix metalloproteinase-2, osteopontin, osteomodulin, prothrombin, thrombospondin and vWF. They recognize the sequence R-G-D in a wide array of ligands. In case of HIV-1 infection, the interaction with extracellular viral Tat protein seems to enhance angiogenesis in Kaposi’s sarcoma lesions.

**Sequence similarities**
Belongs to the integrin alpha chain family.
Contains 7 FG-GAP repeats.

**Cellular localization**
Membrane.

**Images**

ICC/IF image of ab11470 stained A549 cells. The cells were 4% formaldehyde fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab11470, 5µg/ml) overnight at +4°C. The secondary antibody (green) was ab96879, a goat anti-mouse DyLight® 488 (IgG; H+L) used at a 1/250 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.
Overlay histogram showing A549 cells stained with ab11470 (red line). The cells were fixed with 80% methanol (5 min) and incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab11470, 1µg/1x10^6 cells) for 30 min at 22°C. The secondary antibody used was a goat anti-mouse DyLight® 488 (IgG; H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG1 [ICIGG1] (ab91353, 2µg/1x10^6 cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive result in 4% paraformaldehyde (10 min) fixed A549 cells used under the same conditions.

Please note that Abcam do not have any data for use of this antibody on non-fixed cells. We welcome any customer feedback.

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