ABflo® 488 Rabbit anti-Human CD31/PECAM1 mAb

Background

Catalog No.: A22508 1 Publications



Basic Information

Observed MW Refer to figures

Calculated MW 83kDa

Category Primary antibody

Applications FC

Cross-Reactivity Human

CloneNo number ARC54799

Conjugate

ABflo® 488. Ex:491nm. Em:516nm.

Recommended Dilutions

5 µl per 10^6 cells in

100 µl volume

FC

Immunogen Information

Gene ID 5175

Swiss Prot P16284

Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

The protein encoded by this gene is found on the surface of platelets, monocytes, neutrophils, and some types of T-cells, and makes up a large portion of endothelial cell

and is likely involved in leukocyte migration, angiogenesis, and integrin activation.

intercellular junctions. The encoded protein is a member of the immunoglobulin superfamily

Synonyms

CD31; PECA1; GPIIA'; PECAM-1; endoCAM; CD31/EndoCAM

Contact

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Product Information

Source	
Rabbit	

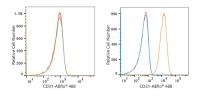
lsotype IgG **Purification** Affinity purification

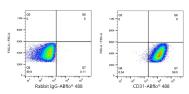
Storage

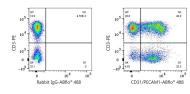
Store at 2-8°C. Avoid freeze.

Buffer: PBS containing 0.2% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

Validation Data







Flow cytometry:1X10^6 SK-BR-3 cells (negative control,left) and HEL cells (right) were surface-stained with ABflo® 488 Rabbit anti-Human CD31/PECAM1 mAb(A22508,5 µl/Test,orange line) or ABflo® 488 Rabbit IgG isotype control (A22069,5 µl/Test,blue line). Non-fluorescently stained cells were used as blank control (red line). Flow cytometry:1X10^6 HEL cells were surface-stained with ABflo® 488 Rabbit IgG isotype control (A22069,5 µl/Test,left) or ABflo® 488 Rabbit anti-Human CD31 mAb(A22508,5 µl/Test,right). Flow cytometry: 1X10^6 Human PBMC were surface-stained with PE Rabbit anti-Human CD3 mAb (A27110,5 μ /Test) and ABflo® 488 Rabbit lgG isotype control (A22069,5 μ /Test,left) or ABflo® 488 Rabbit anti-Human CD31/PECAM1 mAb (A22508,5 μ /Test,right). Cells in the Lymphocytes gate were used for analysis.