

APC/Fire™ 750 anti-human CD49d

Analyte Specific Reagent. Analytical and performance characteristics are not established.

Catalog# / Size 987702 / 500 μL

 Clone
 9F10

 Workshop
 V S215

Other Names VLA-4 α chain, α4 integrin, Integrin α4 chain, ITGA4

Isotype Mouse IgG1, κ

Description CD49d is a 150 kD α integrin chain known as α4 integrin or VLA-4 α chain. It forms a

heterodimer with either integrin $\beta1$ ($\alpha4\beta1$, VLA-4) or $\beta7$ ($\alpha4\beta7$). CD49d is expressed broadly on T lymphocytes, B lymphocytes, monocytes, thymocytes, eosinophils, basophils, mast cells, NK cells, dendritic cells, and some non-hematopoietic cells, but not on normal red blood cells, platelets or neutrophils. VLA-4 binds to VCAM-1 (CD106) and fibronectin. $\alpha4\beta7$ is the receptor for VCAM-1 and MAdCAM-1. CD49d participates in mononuclear cell trafficking to endothelial sites of inflammation and has roles in cell-cell interactions and cell adhesion to extracellular matrices. CD49d is involved in lymphocyte migration, T cell activation, and hematopoietic stem

cell differentiation.

Product Details

Reactivity Human

Formulation Phosphate-buffered solution, pH7.2, containing True-Stain Monocyte Blocker™, 0.09% sodium

azide, 0.2% (w/v) BSA (origin USA), and a stabilizer.

Preparation The antibody was purified by affinity chromatography and conjugated with APC/Fire™ 750 under

optimal conditions.

Concentration 200 µg/mL

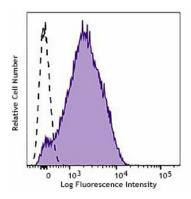
Storage & Handling The antibody solution should be stored undiluted between 2°C and 8°C, and protected from

prolonged exposure to light. Do not freeze.

Application Suggested for Flow Cytometry

Disclaimer WARNINGS AND PRECAUTIONS

- 1. Use appropriate personal protective equipment and safety practices per universal precautions when working with this reagent. Refer to the reagent safety data sheet.
- This antibody contains sodium azide. Follow federal, state and local regulations to dispose of this reagent. Sodium azide build-up in metal wastepipes may lead to explosive conditions; if disposing of reagent down wastepipes, flush with water after disposal.
- 3. All specimens, samples and any material coming in contact with them should be considered potentially infectious and should be disposed of with proper precautions and in accordance with federal, state and local regulations.
- 4. Do not use this reagent beyond the expiration date stated on the label.
- 5. Do not use this reagent if it appears cloudy or if there is any change in the appearance of the reagent as these may be an indication of possible deterioration.
- 6. Avoid prolonged exposure of the reagent or stained cells to light.



Typical results from human peripheral blood lymphocytes stained either with 9F10 APC/Fire™ 750 used at 5 µL/test (filled histogram) or with an isotype control (open histogram).

Symbols Glossary*

Symbol	Meaning	Symbol Title	Symbol No.	Symbol	Meaning	Symbol Title	Symbol No.
REF	Catalog number	Catalogue number	5.1.6	$\bigcap_{\mathbf{i}}$	Indicates the need for the user to consult the instructions for use.	Consult instructions for use	5.4.3
1	Indicates the temperature limits to which the medical device can be safely exposed.	Temperature limit	5.3.7	类	Indicates a medical device that needs protection from light sources.	Keep away from sunlight	5.3.2
K	Indicates the upper limit of temperature to which the medical device can be safely exposed.	Upper limit of temperature	5.3.6	Ω	Indicates the date after which the medical device is not to be used.	Use-by date	5.1.4
	Indicates the medical device manufacturer.	Manufacturer	5.1.1	EC REP	Indicates the authorized representative in the European Community.	Authorized representative in the European Community	
LOT	Indicates the manufacturer's batch code so that the batch or lot can be identified.	Batch code	5.1.5	IVD	Indicates a medical device that is intended to be used as an in vitro diagnostic medical device.	In vitro diagnostic medical device	5.5.1

^{*} Symbol information is from EN ISO 15223-1:2016 Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements

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