

Anti-Human CD49d Antibody

Catalog Number:	114401, 114402
Size:	25 μg, 100 μg
Target Name:	CD49d, VLA-4 α , Integrin α 4, ITGA4
Regulatory Status:	RUO

PRODUCT DETAILS

Clone:	9F10
Application:	Flow Cytometry
Reactivity:	Human
Format:	Purified
Isotype:	Mouse IgG1
Antibody Type:	Monoclonal
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide
Protein Concentration:	0.5 mg/mL
Storage&Handling:	The antibody solution should be stored between 2°C and 8°C
Isotype Controls:	301401
Antibody Family:	Human Antibodies

BACKGROUND INFORMATION

CD49d, also known as integrin α 4, is a cell surface adhesion molecule expressed on leukocytes including lymphocytes, monocytes, and eosinophils. It plays a central role in immune cell trafficking by mediating adhesion to vascular endothelium and facilitating transmigration into tissues during immune surveillance and inflammation. CD49d pairs with β 1 (CD29) or β 7 integrins to form the heterodimers α 4 β 1 (VLA-4) and α 4 β 7, each with distinct tissue-homing functions.

Structurally, CD49d is a type I transmembrane glycoprotein with a large extracellular domain responsible for ligand binding, a single-pass transmembrane region, and a short cytoplasmic tail that links to intracellular signaling and cytoskeletal machinery. Its principal ligands include VCAM-1 (vascular cell adhesion molecule-1) and fibronectin for α 4 β 1, and MAdCAM-1 (mucosal addressin cell adhesion molecule-1) for α 4 β 7, enabling tissue-specific adhesion and migration.

In disease, CD49d contributes to chronic inflammation and autoimmune disorders such as multiple sclerosis and inflammatory bowel disease by promoting leukocyte infiltration into target tissues. It is also implicated in cancer, particularly in leukemias, where high CD49d expression correlates with enhanced survival and poor prognosis.

Therapeutically, CD49d is a validated target. Monoclonal antibodies such as natalizumab block α 4 integrins to reduce immune cell migration, providing clinical benefit in multiple sclerosis and Crohn's disease. Ongoing strategies aim to refine targeting while minimizing immunosuppression-related risks.