

Product datasheet

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ARG54230 anti-CD135 / FLT3 antibody [BV10A4] (PE)

Package: 50 tests Store at: 4°C

Summary

Isotype

Product Description PE-conjugated Mouse Monoclonal antibody [BV10A4] recognizes CD135 / FLT3

Tested Reactivity Hu
Species Does Not React With Ms
Tested Application FACS

Specificity The mouse monoclonal antibody BV10A4 (BV10) reacts with CD135 (FLT3, FLK2, STK1), a 130160 kDa

type III receptor tyrosine kinase that is involved in early steps of hematopoiesis.

Host Mouse

Clonality Monoclonal

Clone BV10A4

Target Name CD135 / FLT3

Immunogen BV-173 leukemic cell line

Conjugation PE

Alternate Names CD135; FLK2; Receptor-type tyrosine-protein kinase FLT3; FLK-2; STK-1; STK1; FL cytokine receptor;

FLT-3; Stem cell tyrosine kinase 1; Fetal liver kinase-2; Fms-like tyrosine kinase 3; CD antigen CD135; EC

2.7.10.1

lgG1

Application Instructions

Application table	Application	Dilution
	FACS	20 μl / 10^6 cells
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form Liquid

Purification Note The purified antibody is conjugated with R-Phycoerythrin (PE) under optimum conditions. The

conjugate is purified by size-exclusion chromatography and adjusted for direct use. No reconstitution is

necessary.

Buffer PBS, 15 mM Sodium azide and 0.2% (w/v) high-grade protease free BSA

Preservative 15 mM Sodium azide

Stabilizer 0.2% (w/v) high-grade protease free BSA

Storage instruction Aliquot and store in the dark at 2-8°C. Keep protected from prolonged exposure to light. Avoid

repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be

gently mixed before use.

Bioinformation

Database links GeneID: 2322 Human

Swiss-port # P36888 Human

Gene Symbol FLT3

Gene Full Name fms-related tyrosine kinase 3

Background CD135 / FLT3, also known as FLK2 or STK-1 is a receptor tyrosine kinase that plays important roles in

hematopoiesis. After binding of Flt3 ligand (FL), CD135 homodimerizes and stimulates proliferation, differentiation and protects the cell from apoptosis. The loss of CD90 and gain of CD135 expression marks the loss of self-renewal in hematopoietic stem cell population. Detectable CD135 expression appears first at low levels on the surface of primitive multilineage progenitor cells and disappears during defined stages of B-cell development, but is upregulated and maintained during maturation of monocytes. CD135 is also expressed on thymocytes, dendritic cell progenitors and on mature dendritic

cells, as well as on various malignant hematopoietic cells.

Function Tyrosine-protein kinase that acts as cell-surface receptor for the cytokine FLT3LG and regulates

differentiation, proliferation and survival of hematopoietic progenitor cells and of dendritic cells. Promotes phosphorylation of SHC1 and AKT1, and activation of the downstream effector MTOR. Promotes activation of RAS signaling and phosphorylation of downstream kinases, including

MAPK1/ERK2 and/or MAPK3/ERK1. Promotes phosphorylation of FES, FER, PTPN6/SHP, PTPN11/SHP-2, PLCG1, and STAT5A and/or STAT5B. Activation of wild-type FLT3 causes only marginal activation of STAT5A or STAT5B. Mutations that cause constitutive kinase activity promote cell proliferation and

resistance to apoptosis via the activation of multiple signaling pathways. [UniProt]

Research Area Immune System antibody; Signaling Transduction antibody

Calculated Mw 113 kDa

PTM N-glycosylated, contains complex N-glycans with sialic acid.

Autophosphorylated on several tyrosine residues in response to FLT3LG binding. FLT3LG binding also increases phosphorylation of mutant kinases that are constitutively activated. Dephosphorylated by PTPRJ/DEP-1, PTPN1, PTPN6/SHP-1, and to a lesser degree by PTPN12. Dephosphorylation is important

for export from the endoplasmic reticulum and location at the cell membrane.

Rapidly ubiquitinated by UBE2L6 and the E3 ubiquitin-protein ligase SIAH1 after autophosphorylation,

leading to its proteasomal degradation.