

Anti-Human CD38 Antibody, Clone AT-1, FITC

Mouse monoclonal IgG1 antibody against human, rhesus, cynomolgus CD38, FITC-conjugated

Catalog #100-1578

100 Tests

20 µL/test

Product Description

This monoclonal antibody reacts with CD38, an ~45 kDa type II integral membrane glycoprotein that is part of the ADP-ribosyl cyclase family. CD38 is widely expressed at variable levels on hematopoietic cells including early B and T cell lineages, activated B and T cells, NK cells, monocytes, and dendritic cells. It is strongly expressed in plasma cells. CD38 is also expressed in some non-hematopoietic tissues such as brain, kidney, muscle, and pancreas. CD38 is an ectoenzyme that acts as both an ADP-ribosyl cyclase and ADP-ribose (ADPR) hydrolase for the synthesis and hydrolysis of cyclic ADPR (cADPR) and NAADP. cADPR acts as a second messenger for intracellular calcium mobilization to facilitate glucose-induced insulin secretion. CD38 functions as a signaling receptor and thus associates with various molecules, one of which is CD31, to mediate unique cell-type specific signaling. On hematopoietic cells, CD38 mediates adhesion and the activation, proliferation, and differentiation of mature B and T cells, and the apoptosis of myeloid and lymphoid progenitor cells. CD38 monoclonal antibodies have been used to study B and T cell differentiation and activation, to inhibit B lymphopoiesis, and to protect B cells from apoptosis. It has been shown that the AT-1 antibody mutually competes for binding with OKT10 and SUN-4B7, and all three clones only bind the native (not reduced) form of CD38.

Target Antigen:	CD38
Alternative Names:	ADP-ribosyl cyclase, cADPR, Cyclic ADP-ribose hydrolase, T10
Gene ID:	952
Species Reactivity:	Human, Rhesus, Cynomolgus, Pigtailed macaque
Host Species:	Mouse
Clonality:	Monoclonal
Clone:	AT-1
Isotype:	IgG1, kappa
Immunogen:	Human T cell line CCRF-CEM

Applications

Verified Applications: FC

Reported Applications: FC

Abbreviations: CellSep: Cell separation; ChIP: Chromatin immunoprecipitation; FA: Functional assay; FACS: Fluorescence-activated cell sorting; FC: Flow cytometry; FCXM: Flow cytometric crossmatch assay; FISH: Fluorescence in situ hybridization; ICC: Immunocytochemistry; IF: Immunofluorescence microscopy; IHC: Immunohistochemistry; IHC-F: Immunohistochemistry (frozen-tissue); IHC-P: Immunohistochemistry (paraffin-embedded); IP: Immunoprecipitation; NMR: Nuclear magnetic resonance spectroscopy; RIA: Radioimmunoassay; WB: Western blotting

Properties

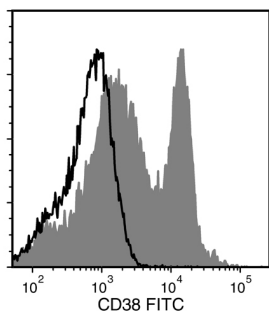
Product Formulation: Phosphate-buffered saline containing 0.1% bovine serum albumin and less than 0.1% sodium azide

Purification: The antibody was purified by affinity chromatography.

Stability and Storage: Product stable at 2 - 8°C when stored undiluted. Do not freeze. Protect product from prolonged exposure to light. For product expiry date, contact techsupport@stemcell.com.

Directions for Use: For flow cytometry, the suggested use of this antibody is 20 μ L per 1×10^6 cells in 100 μ L. It is recommended that the antibody be titrated for optimal performance for each application.

Data



Flow cytometry analysis of human peripheral blood mononuclear cells (PBMCs) labeled with Anti-Human CD38 Antibody, Clone AT-1, FITC (filled histogram), or Mouse IgG1, kappa Isotype Control Antibody, Clone MOPC-21, FITC (Catalog #60070FI; solid line histogram).

Related Products

For a complete list of antibodies, including other conjugates, sizes, and clones, as well as related products available from STEMCELL Technologies, visit www.stemcell.com/antibodies, or contact us at techsupport@stemcell.com.

References

- Ausiello CM et al. (2000) Functional topography of discrete domains of human CD38. *Tissue Antigens* 56(6): 539–47.
- Cosker F et al. (2010) The ecto-enzyme CD38 is a nicotinic acid adenine dinucleotide phosphate (NAADP) synthase that couples receptor activation to Ca²⁺ mobilization from lysosomes in pancreatic acinar cells. *J Biol Chem* 285(49): 38251–9.
- Deaglio S et al. (1998) Human CD38 (ADP-ribosyl cyclase) is a counter-receptor of CD31, an Ig superfamily member. *J Immunol* 160(1): 395–402.
- Deaglio S et al. (2007) CD38/CD19: a lipid raft-dependent signaling complex in human B cells. *Blood* 109(12): 5390–8.
- Graeff R et al. (2006) Acidic residues at the active sites of CD38 and ADP-ribosyl cyclase determine nicotinic acid adenine dinucleotide phosphate (NAADP) synthesis and hydrolysis activities. *J Biol Chem* 281(39): 28951–7.
- Ferrero E et al. (2004) Characterization and phylogenetic epitope mapping of CD38 ADPR cyclase in the cynomolgus macaque. *BMC Immunol* 5: 21.
- Hoshino S et al. (1997) Mapping of the catalytic and epitopic sites of human CD38/NAD⁺ glycohydrolase to a functional domain in the carboxyl terminus. *J Immunol* 158(2): 741–7.
- Kader M et al. (2013) Blocking TLR7- and TLR9-mediated IFN- α production by plasmacytoid dendritic cells does not diminish immune activation in early SIV infection. *PLoS Pathog* 9(7): e1003530.
- Terstappen LW et al. (1991) Sequential generations of hematopoietic colonies derived from single nonlineage-committed CD34+CD38-progenitor cells. *Blood* 77(6): 1218–27.
- Vaisitti T et al. (2010) CD38 increases CXCL12-mediated signals and homing of chronic lymphocytic leukemia cells. *Leukemia* 24(5): 958–69.
- Weinfurter JT et al. (2011) Cross-reactive T cells are involved in rapid clearance of 2009 pandemic H1N1 influenza virus in nonhuman primates. *PLoS Pathog* 7(11): e1002381.

PRODUCTS ARE FOR RESEARCH USE ONLY AND NOT INTENDED FOR HUMAN OR ANIMAL DIAGNOSTIC OR THERAPEUTIC USES UNLESS OTHERWISE STATED.

Copyright © 2024 by STEMCELL Technologies Inc. All rights reserved including graphics and images. STEMCELL Technologies & Design, STEMCELL Shield Design, and Scientists Helping Scientists are trademarks of STEMCELL Technologies Canada Inc. All other trademarks are the property of their respective holders. While STEMCELL has made all reasonable efforts to ensure that the information provided by STEMCELL and its suppliers is correct, it makes no warranties or representations as to the accuracy or completeness of such information.