

Anti-Human CD235a (Glycophorin A) Antibody, Clone 2B7, FITC

Mouse monoclonal IgG1 antibody against human CD235a (glycophorin A), FITC-conjugated

Catalog #100-1672

100 Tests

20 µL/test

Product Description

This monoclonal antibody reacts with CD235a (Glycophorin A), a 10 kDa type I sialoglycoprotein present in the cell membrane of erythrocytes and erythroid precursors as a homodimer. Glycophorin A bears the antigenic determinants for the MN and Ss blood groups and has been proposed to provide a large mucin-like surface to erythrocytes that acts to minimize aggregation in circulation. Glycophorin A is first detectable on morphologically recognizable erythroid precursors just after the colony-forming unit erythroid (CFU-E) stage, and reaches its maximal expression at the late normoblast stage. Anti-glycophorin is useful in combination with anti-transferrin receptor (CD71) to identify distinct stages of erythroid differentiation since CD71 expression precedes Glycophorin A expression, but is lost during maturation of normoblasts into mature red blood cells (RBCs). Peptide-binding ELISA data indicate that the epitope of the 2B7 antibody is located in the extracellular domain of CD235a within the sequence Ala54 - Ser66 (AATPRAHEVSEIS).

Target Antigen:	CD235 (Glycophorin)
Alternative Names:	Glycophorin A, GYPA, MN sialoglycoprotein, MNS blood group, PAS-2, Sialoglycoprotein alpha
Gene ID:	2993
Species Reactivity:	Human
Host Species:	Mouse
Clonality:	Monoclonal
Clone:	2B7
Isotype:	IgG1, kappa
Immunogen:	Cell lysate containing partially lysed human red blood cells
Conjugate:	FITC (Fluorescein isothiocyanate)

Applications

Verified Applications: FC

Reported Applications: ELISA, 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

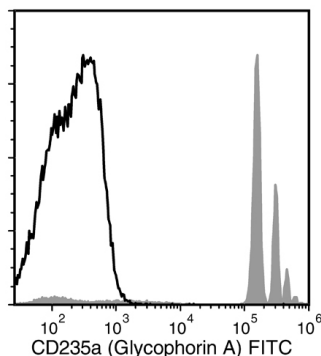
Product Formulation: Phosphate-buffered saline containing 0.1% bovine serum albumin and less than 0.1% sodium azide

Purification: The antibody was purified by affinity chromatography.

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

Directions for Use: For flow cytometry, the suggested use of this antibody is 20 µL per 1×10^6 cells in 100 µL. This volume is usually appropriate for labeling samples containing mature RBCs, but may be too high for labeling immature RBCs in samples that have been depleted of mature RBCs. It is recommended that the antibody be titrated for optimal performance for each application.

Data



Flow cytometry analysis of human whole blood labeled with Anti-Human CD235a (Glycophorin A) Antibody, Clone 2B7, 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

Loken MR et al. (1987) Flow cytometric analysis of human bone marrow: I. Normal erythroid development. *Blood* 69(1): 255–63.

Paes BCMF et al. (2020) Generation of hematopoietic stem/progenitor cells with sickle cell mutation from induced pluripotent stem cell in serum-free system. *Hematol Transfus Cell Ther.* 2021 Apr-Jun;43(2):156-64.

Robinson J et al. (1981) Expression of cell-surface HLA-DR, HLA-ABC and glycophorin during erythroid differentiation. *Nature* 289(5793): 68–71.

Ruiz JP (2019) Robust generation of erythroid and multilineage hematopoietic progenitors from human iPSCs using a scalable monolayer culture system. *Stem Cell Res* 41: 101600.

Wu Y et al. (2019) Highly efficient therapeutic gene editing of human hematopoietic stem cells. *Nat Med* 25(5): 776–83.

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