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Datasheet for ABIN1305243 Anti-Human CD19 Magnetic Particles

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Overview

Quantity:	5 mL
Target:	CD19
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	Magnetic Particles
Application:	Separation (Sep)
Product Details	
Brand:	BD IMag™
Target Details	
Target:	CD19
Alternative Name:	CD19 (CD19 Products)
Background:	BD IMag [™] anti-human CD19 Particles - DM are magnetic nanoparticles that have monoclonal antibody conjugated to their surfaces. These particles are optimized for the positive selection or depletion of CD19-bearing leukocytes using the BD IMagnet [™] . CD19 is expressed during all stages of B-cell differentiation and maturation, except plasma cells. CD19 is also present on follicular dendritic cells. It is not found on T cells or on normal granulocytes. Peripheral Blood Mononuclear Cells (PBMC) are labeled with BD IMag [™] anti-human CD19 Particles - DM according to the Magnetic Labeling Protocol. This labeled cell suspension is then placed within the magnetic field of the BD IMagnet [™] (Cat. No. 552311). Labeled cells migrate toward the

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 1/4 | Product datasheet for ABIN1305243 | 07/26/2024 | Copyright antibodies-online. All rights reserved. magnet (positive fraction), leaving the unlabeled cells in suspension so they can be drawn off (negative fraction). The tube is then removed from the magnetic field for resuspension of the positive fraction. The separation is repeated twice to increase the purity of the positive fraction. The magnetic separation steps are diagrammed in the Separation Flow Chart. After the positive fraction is washed, the small size of the magnetic particles allows the positive fraction to be further evaluated in downstream applications such as flow cytometry.

Application Details

Protocol:	1. Prepare PBMC from anti-coagulated human blood, preferably by density gradient
	centrifugation using Ficoll-Paque™. Remove clumps of cells and/or debris by passing the
	suspension through a 70-µm nylon cell strainer.
	2. Dilute BD IMag™ Buffer (10X) (Cat. No. 552362) 1:10 with sterile distilled water or prepare 1X
	BD IMag™ buffer by supplementing Phosphate Buffered Saline with 0.5% BSA, 2 mM EDTA, and
	0.09% sodium azide). Store at 4°C.
	3. Wash cells with an excess volume of 1X BD IMag™ buffer, and carefully aspirate all the
	supernatant.
	4. Vortex the BD IMag™ anti-human CD19 Particles - DM thoroughly, and add 50 μl of particles
	for every 10^7 total cells.
	5. MIX THOROUGHLY. Incubate at room temperature for 30 minutes.
	6. Bring the BD IMag™-particle labeling volume up to 1 -8 x 10^7 cells/ml with 1X BD IMag™
	buffer, and immediately place the tube on the BD IMagnet™. Incubate for 8 - 10 minutes.
	7. With the tube on the BD IMagnet™, carefully aspirate off the supernatant. This supernatant
	contains the negative fraction.
	8. Remove the tube from the BD IMagnet™, and add 1X BD IMag™ buffer to the same volume as
	in step 6. Gently resuspend cells by pipetting up and down, and return the tube to the BD
	IMagnet™ for another 2 - 4 minutes.
	9. With the tube on the BD IMagnet™, carefully aspirate off the supernatant and discard.
	10. Repeat Steps 8 and 9.
	11. After the final wash step, resuspend the positive fraction in an appropriate buffer or media,
	and proceed with desired downstream application(s). The concentration of BD IMag™ anti-
	human CD19 Particles - DM suggested in this protocol has been optimized for the purification
	of CD19 positive B lymphocytes from human peripheral blood. When labeling target cell
	populations present at lower frequencies, fewer BD IMag™ particles can be used. Conversely,
	when labeling target cell populations that are present at higher frequencies, more particles
	should be used. To determine the optimal concentration of the BD IMag $^{ m m}$ anti-human CD19

Application Details	
	Particles - DM for a particular application, a titration in two-fold increments is recommended. NOTE: Avoid nonspecific labeling by working quickly and keeping incubation times to a minimum.
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Aqueous buffered solution containing BSA and ≤ 0.09 % sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C
Storage Comment:	Store undiluted at 4° C. Page 1 of 3551520 Rev. 3
Publications	
Product cited in:	Bradbury, Goldmacher, Tedder: "The CD19 signal transduction complex of B lymphocytes. Deletion of the CD19 cytoplasmic domain alters signal transduction but not complex formation with TAPA-1 and Leu 13." in: Journal of immunology (Baltimore, Md. : 1950) , Vol. 151, Issue 6, pp. 2915-27, (1993) (PubMed).
	Uckun, Muraguchi, Ledbetter, Kishimoto, OBrien, Roloff, Gajl-Peczalska, Provisor, Koller: " Biphenotypic leukemic lymphocyte precursors in CD2+CD19+ acute lymphoblastic leukemia and their putative normal counterparts in human fetal hematopoietic tissues." in: Blood , Vol. 73, Issue 4, pp. 1000-15, (1989) (PubMed).
	Nadler, Anderson, Marti, Bates, Park, Daley, Schlossman: "B4, a human B lymphocyte- associated antigen expressed on normal, mitogen-activated, and malignant B lymphocytes." in: Journal of immunology (Baltimore, Md. : 1950), Vol. 131, Issue 1, pp. 244-50, (1983) (PubMed).

Flow Cytometry

Image 1.



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