

EasySep™ Human CD45 Depletion Kit II

Positive Selection

Catalog #17898

For processing 2 x 10⁹ cells



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Description

Deplete CD45+ cells from fresh or previously frozen human peripheral blood mononuclear cells (PBMCs).

- · Fast, easy-to-use and column-free
- Up to 4 log depletion of CD45+ cells
- · Isolated cells are untouched

This kit targets CD45+ cells for removal with an antibody recognizing the CD45 cell surface marker. Unwanted cells are labeled with antibodies and magnetic particles, and separated without columns using an EasySepTM magnet. Desired cells are simply poured off into a new tube. Isolated cells are immediately available for downstream applications such as flow cytometry, culture, or DNA/RNA extraction.

Component Descriptions

COMPONENT NAME	COMPONENT #	QUANTITY	STORAGE	SHELF LIFE	FORMAT
EasySep™ Human CD45 Depletion Cocktail II	17898C	1 x 1 mL	Store at 2 - 8°C. Do not freeze.	Stable until expiry date (EXP) on label.	A combination of monoclonal antibodies in PBS.
EasySep™ Dextran RapidSpheres™ 50101	50101	2 x 1 mL	Store at 2 - 8°C. Do not freeze.	Stable until expiry date (EXP) on label.	A suspension of magnetic particles in water.

PBS - phosphate-buffered saline

Components may be shipped at room temperature (15 - 25°C) but should be stored as indicated above.

Sample Preparation

For available fresh and frozen samples, see www.stemcell.com/primarycells.

PERIPHERAL BLOOD

Prepare a PBMC suspension from whole blood by centrifugation over a density gradient medium (e.g. Lymphoprep[™], Catalog #07801). For more rapid PBMC preparation, use the SepMate[™] RUO (Catalog #86450/86415) or SepMate[™] IVD* (Catalog #85450/85415) cell isolation tube.

If using previously frozen PBMCs, incubate the cells with DNase I Solution (Catalog #07900) at a concentration of 100 μg/mL at room temperature (15 - 25°C) for at least 15 minutes prior to labeling and separation. Filter aggregated suspensions through a 40 μm Cell Strainer (Catalog #27305) for optimal results.

After preparation, resuspend cells at 1 x 10^8 cells/mL in recommended medium.

Recommended Medium

EasySep™ Buffer (Catalog #20144), RoboSep™ Buffer (Catalog #20104), or PBS containing 2% fetal bovine serum (FBS) and 1 mM EDTA. Medium should be free of Ca++ and Mg++.

^{*} SepMate™ IVD is only available in select regions where it is registered as an In Vitro Diagnostic (IVD) device for the isolation of mononuclear cells (MNCs) from whole blood or bone marrow by density gradient centrifugation. In all other regions SepMate™ is available for research use only (RUO).



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Directions for Use - Manual EasySep™ Protocols

See page 1 for Sample Preparation and Recommended Medium. Refer to Table 1 for detailed instructions regarding the EasySep™ procedure for each magnet.

Table 1. EasySep™ Human CD45 Depletion Kit II Protocol

		EASYSEP™ MAGNETS		
STEP	INSTRUCTIONS	EasySep™ (Catalog #18000)	"The Big Easy" (Catalog #18001)	
1	Prepare sample at the indicated cell concentration within the volume range.	1 x 10^8 cells/mL 0.1 - 2 mL NOTE: If starting with fewer than 1 x 10^7 cells, resuspend cells in 0.1 mL	1 x 10^8 cells/mL 0.25 - 8.5 mL NOTE: If starting with fewer than 2.5 x 10^7 cells, resuspend cells in 0.25 mL	
	Add sample to required tube.	5 mL (12 x 75 mm) polystyrene round-bottom tube (e.g. Catalog #38007)	14 mL (17 x 95 mm) polystyrene round-bottom tube (e.g. Catalog #38008)	
2	Add Depletion Cocktail to sample.	50 μL/mL of cells	50 μL/mL of cells	
	Mix and incubate.	RT for 5 minutes	RT for 5 minutes	
3	Vortex RapidSpheres™. NOTE: Particles should appear evenly dispersed.	30 seconds	30 seconds	
4	Add RapidSpheres™ to sample.	75 μL/mL of sample	75 μL/mL of sample	
	Mix and incubate.	RT for 3 minutes	RT for 3 minutes	
5	Add recommended medium to top up the sample to the indicated volume. Mix by gently pipetting up and down 2 - 3 times.	Top up to 2.5 mL	 Top up to 5 mL for samples < 4 mL Top up to 10 mL for samples ≥ 4 mL 	
	Place the tube (without lid) into the magnet and incubate.	RT for 5 minutes	RT for 5 minutes	
6	Pick up the magnet, and in one continuous motion invert the magnet and tube,* pouring the enriched cell suspension into a new tube.	Use new 5 mL tube	Use new 14 mL tube	
7	Remove the tube from the magnet and place the new tube (without lid) into the magnet and incubate for a second separation.	RT for 5 minutes	RT for 5 minutes	
8	Pick up the magnet, and in one continuous motion invert the magnet and tube,* pouring the enriched cell suspension into a new tube.	Isolated cells are ready for use	Isolated cells are ready for use	

RT - room temperature (15 - 25°C)

^{*} Leave the magnet and tube inverted for 2 - 3 seconds, then return upright. Do not shake or blot off any drops that may remain hanging from the mouth of the tube.



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Directions for Use – Fully Automated RoboSep™ Protocol

See page 1 for Sample Preparation and Recommended Medium. Refer to Table 2 for detailed instructions regarding the RoboSep™ procedure.

Table 2. RoboSep™ Human CD45 Depletion Kit II Protocol

STEP	INSTRUCTIONS	RoboSep™ (Catalog #20000 and #21000)				
1	Prepare sample at the indicated cell concentration within the volume range.	1 x 10^8 cells/mL 0.5 - 8.5 mL NOTE: If starting with fewer than 5 x 10^7 cells, resuspend cells in 0.5 mL				
	Add sample to required tube.	14 mL (17 x 95 mm) polystyrene round-bottom tube (e.g. Catalog #38008)				
2	Select protocol.	Human CD45 Depletion II 17898				
3	Vortex RapidSpheres™. NOTE: Particles should appear evenly dispersed.	30 seconds				
4	Load the carousel.	Follow on-screen prompts				
	Start the protocol.	Press the green "Run" button				
5	Unload the carousel when the run is complete. Remove the tube containing the enriched cells.	Isolated cells are ready for use				

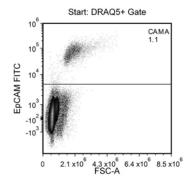
Notes and Tips

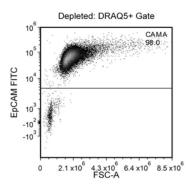
ASSESSING PURITY

For purity assessment of residual CD45+ cells by flow cytometry use one of the following fluorochrome-conjugated antibody clones:

- · Anti-Human CD45 Antibody, Clone HI30 (Catalog #60018; partially blocked), or
- · Anti-Human CD45 Antibody, Clone 2D1 (Catalog #60123; partially blocked)

Data





In the example above, CAMA cells were seeded into PBMCs at a starting frequency of 1.1% (98.9% CD45+; gated on DRAQ5™ for nucleated cells). The CAMA cell (EpCAM+) content of the depleted fraction is 98%, which is a 4.0 log depletion of CD45+ cells.

NOTE: EpCAM is an antibody against an epithelial cell surface antigen expressed on CAMA cells.

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