

Negative Selection

Catalog #18170

For processing 100 mL bone marrow



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Description

Deplete red blood cells (RBCs) directly from human bone marrow by immunomagnetic negative selection. This kit can also be used to deplete RBCs from other sample types (see Table 1).

The benefits of this kit include:

- · 99.9% RBC depletion without the need for density gradient centrifugation, sedimentation or lysis
- · Fast, easy-to-use and column-free
- · Isolated cells are untouched

This kit targets RBCs for removal with antibodies recognizing specific cell surface markers. Unwanted cells are labeled with antibodies and magnetic particles, and separated without columns using an EasySep™ magnet. RBC-depleted nucleated cells are simply collected into a new tube and are immediately available for downstream applications such as flow cytometry, culture, or DNA/RNA extraction.

• This is the Product Information Sheet (PIS) for depleting RBCs from bone marrow. If depleting RBCs from other sample types, refer to the applicable PIS Document Number (see Table 1).

Table 1. Applicable Document Number for Other Sample Types

SAMPLE TYPE	DOCUMENT NUMBER
Whole blood	10000003345
Buffy coat	1000005628
Cord blood	10000005630
Leukapheresis	10000005627

Component Descriptions

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

EasySep™ RBC Depletion Reagent (Catalog #18170)

COMPONENT NAME	COMPONENT #	QUANTITY	STORAGE	SHELF LIFE	FORMAT
EasySep™ RBC Depletion Reagent	18170	10 mL	Store at 2 - 8°C. Do not freeze.	Stable until expiry date (EXP) on label.	A suspension of magnetic particles and monoclonal antibodies in PBS.

PBS - phosphate-buffered saline

EasySep™ RBC Depletion Reagent for RoboSep™ (Catalog #18170RF)

COMPONENT NAME	COMPONENT #	QUANTITY	STORAGE	SHELF LIFE	FORMAT		
EasySep™ RBC Depletion Reagent	18170C	4 x 2.5 mL	Store at 2 - 8°C.	Stable until expiry date (EXP) on label.	A suspension of magnetic particles and monoclonal antibodies in PBS.		

PBS - phosphate-buffered saline

Sample Preparation

BONE MARROW

- 1. Dilute the sample 5- to 10- fold in recommended medium and mix gently by pipetting up and down.
- 2. Centrifuge cells at 300 x g for 10 minutes with the brake on. Remove and discard the supernatant.
- 3. Resuspend the cell pellet with recommended medium. Filter the sample through a pre-wetted 70 µm strainer to remove bone fragments, cell aggregates, and debris. Rinse the strainer with recommended medium.
- 4. Centrifuge cells at 300 x g for 10 minutes with the brake off.
- 5. Carefully remove and discard the plasma, without disturbing the cell pellet.
- 6. Resuspend the cell pellet to the original sample volume with recommended medium.

To avoid loss of monocytes, EDTA must be added to the bone marrow sample to a final concentration of 6 mM prior to labeling and separation (see step 2, Tables 2 - 4).

NOTE: An EDTA stock solution greater than 0.05 M is recommended to avoid over diluting the start sample.

Recommended Medium

PBS containing 2% fetal bovine serum (FBS). Medium should be free of Ca++ and Mg++.





Directions for Use - Manual EasySep™ Protocols

See page 1 for Sample Preparation and Recommended Medium. Refer to Tables 2 and 3 for detailed instructions regarding the EasySep™ procedure for each magnet.

Table 2. EasySep™ RBC Depletion Reagent Protocol for BONE MARROW

=.		EASYSEP™ MAGNETS				
STEP	INSTRUCTIONS	EasySep™ (Catalog #18000)	"The Big Easy" (Catalog #18001)			
	Prepare sample within the volume range.	0.5 - 1 mL	1 - 5 mL			
1	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 EDTA to sample.	At a final concentration of 6 mM EDTA	At a final concentration of 6 mM EDTA			
3	Dilute sample with recommended medium.	Equal volume to sample	Equal volume to sample			
4	Vortex Depletion Reagent. NOTE: Reagent should appear evenly dispersed.	30 seconds	30 seconds			
5	Add Depletion Reagent to sample and mix.	50 μL/mL of original sample volume NOTE: No incubation, IMMEDIATELY proceed to next step	50 μL/mL of original sample volume NOTE: No incubation, IMMEDIATELY proceed to next step			
6	Place the tube (without lid) into the magnet and incubate.	RT for 5 minutes	RT for 5 minutes			
7	Pick up the magnet, and in one continuous motion invert the magnet and tube,* pouring the cell suspension into a new tube.	Use a new 5 mL tube	Use a new 14 mL tube			
8	Add Depletion Reagent to the new tube containing the depleted cells and mix.	Use same volume as in step 5 NOTE: No incubation, IMMEDIATELY proceed to next step	Use same volume as in step 5 NOTE: No incubation, IMMEDIATELY proceed to next step			
9	Remove the tube from the magnet and place the tube from step 8 (without lid) into the magnet and incubate for a second separation.	RT for 5 minutes	RT for 5 minutes			
10	Pick up the magnet, and in one continuous motion invert the magnet and tube,** pouring the cell suspension into a new tube.	Isolated cells are ready for use	Isolated cells are ready for use			

RT - room temperature (15 - 25°C)

^{*} Following the first magnetic separation, the collected cells may contain a significant amount of RBCs and may look similar to the original human bone marrow sample.

^{**} To minimize RBC contamination in the isolated cells, pour off the sample along a clean area of the tube (i.e. the opposite side to where the sample was poured in).





Table 3. EasySep™ RBC Depletion Reagent Protocol for BONE MARROW

		EASYSEP™ MAGNETS						
STEP	STED INSTRUCTIONS		EasyEights™ (Catalog #18103)				Easy 50	Nation
SIEP IN	INSTRUCTIONS	THE PERSON NAMED IN COLUMN	5 mL tube	14 mL	tube	111111111111111111111111111111111111111	(Catalog #18002)	
	Collect sample within the volume range.	0.5 - 1 mL		1 - 5 mL		5 - 20 mL		
1	Add sample to required tube.	poly	5 mL (12 x 75 mm) styrene round-bottom tube (e.g. Catalog #38007)	14 mL (17 x 95 mm) polystyrene round-bottom tube (e.g. Catalog #38008)		50 mL (30 x 115 mm) conical t (e.g. Catalog #38010)	tube	
2	Add EDTA to sample.	At a fina	I concentration of 6 mM EDTA	At a final concentration of 6 mM EDTA			At a final concentration of 6 mM	EDTA
3	Dilute sample with recommended medium.	E	Equal volume to sample	Equal volume	e to sample)	Equal volume to sample	
4	Vortex Depletion Reagent. NOTE: Reagent should appear evenly dispersed.		30 seconds	30 seconds			30 seconds	
5	Add Depletion Reagent to sample and mix.		mL of original sample volume incubation, IMMEDIATELY proceed to next step	50 μL/mL of original sample volume NOTE: No incubation, IMMEDIATELY proceed to next step			50 μL/mL of original sample vol NOTE: No incubation, IMMEDIATELY to next step	
6	Place the tube (without lid) into the magnet and incubate.	RT for 5 minutes		RT for 5 minutes		RT for 10 minutes		
7	Carefully pipette*** (do not pour) the cell suspension into a new tube. NOTE: Collect the entire clear fraction from top to bottom. For optimal recovery, also collect a small volume of RBCs (up to 10% of the starting sample volume).	Use a new 5 mL tube		Use a new ⁻	14 mL tube	e Use a new 50 mL tube		
8	Add Depletion Reagent to the new tube containing the depleted cells and mix.	Use same volume as in step 5 NOTE: No incubation, IMMEDIATELY proceed to next step		Use same volur NOTE: No incubation, IN to next	MEDIATEL'		Use same volume as in step 5 NOTE: No incubation, IMMEDIATELY proce to next step	
9	Remove the tube from the magnet and place the tube from step 8 (without lid) into the magnet and incubate for a second separation.	RT for 5 minutes		RT for 5 minutes		RT for 10 minutes		
10	Carefully pipette*** (do not pour) the cell suspension into a new tube.	Use a new 5 mL tube		Use a new 14 mL tube		Use a new 50 mL tube		
11	Remove the tube from the magnet and place the tube from step 10 (without lid) into the magnet and incubate for a third separation.		RT for 5 minutes	RT for 5 minutes			RT for 10 minutes	
12	Carefully pipette*** (do not pour) the cell suspension into a new tube.	Isola	ated cells are ready for use	Isolated cells are ready for use			Isolated cells are ready for us	se

RT - room temperature (15 - 25°C)

^{***} Collect the entire supernatant, all at once, into a single pipette (e.g. for EasyEights[™] 5 mL tube use a 2 mL serological pipette [Catalog #38002]; for EasyEights[™] 14 mL tube use a 10 mL serological pipette [Catalog #38004]).





Directions for Use – Fully Automated RoboSep™ Protocol

See page 1 for Sample Preparation and Recommended Medium. Refer to Table 4 for detailed instructions regarding the RoboSep™ procedure. NOTE: If using RoboSep™-S, ensure the software is at least v.1.2.0.2 and a carousel compatible with this product is installed. Contact us at techsupport@stemcell.com for more information.

Table 4. RoboSep™ RBC Depletion Reagent Protocol for BONE MARROW

STEP	INSTRUCTIONS	RoboSep [™] (Catalog #21000)			
	Prepare sample within the volume range.	1 - 5 mL			
1	Add sample to required tube.	14 mL (17 x 95 mm) polystyrene round-bottom tube (e.g. Catalog #38008)			
2	Add EDTA to sample.	At a final concentration of 6 mM EDTA			
3	Select protocol.	Human RBC Depletion 18170 - WB CB BM			
4	Vortex Depletion Reagent. NOTE: Reagent should appear evenly dispersed.	30 seconds			
5	Load the carousel.	Follow on-screen prompts			
5	Start the protocol.	Press the green "Run" button			
6	Unload the carousel when the run is complete.	Isolated cells are ready for use			

Notes and Tips

Not suitable for use with downstream magnetic positive selection products.

ASSESSING PURITY

For purity assessment of residual RBCs by flow cytometry, use the following fluorochrome-conjugated antibody clone:

· Anti-Human CD235ab (Glycophorin A/B), Clone HIR2 (Catalog #60111)

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