



EasySep™ Human CD34 Positive Selection Kit

Positive Selection

Catalog #18056

For processing 5×10^9 cells



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Document #28386 | Version 2_2_0

Description

Isolate highly purified CD34+ cells from fresh or previously frozen mobilized human peripheral blood or bone marrow mononuclear cells (MNCs), or from previously frozen cord blood MNCs by immunomagnetic positive selection.

- Fast and easy-to-use
- Up to 99% purity
- No columns required

This kit targets CD34+ cells for positive selection with an antibody recognizing the CD34 surface marker. Desired cells are labeled with antibodies and magnetic particles, and separated without columns using an EasySep™ magnet. Unwanted cells are simply poured off, while desired cells remain in the tube. Isolated cells are immediately available for downstream applications such as flow cytometry, culture, or DNA/RNA extraction.

If isolating CD34+ cells from fresh cord blood, we recommend using the EasySep™ Human Cord Blood CD34 Selection Kit II (Catalog #17896). If isolating CD34+ cells from fresh whole blood or buffy coat, we recommend using the EasySep™ Whole Blood CD34 Selection Kit (Catalog #18086).

Component Descriptions

COMPONENT NAME	COMPONENT #	QUANTITY	STORAGE	SHELF LIFE	FORMAT
EasySep™ CD34 Positive Selection Cocktail	18056C.1	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™ Magnetic Nanoparticles Positive Selection	18150	1 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 or BONE MARROW

Prepare a MNC suspension from whole mobilized peripheral blood or whole bone marrow by centrifugation over a density gradient medium (e.g. Lymphoprep™, Catalog #07801). For more rapid MNC preparation, use the SepMate™ RUO (Catalog #86450/86415) or SepMate™ IVD* (Catalog #85450/85415) cell isolation tube. Older samples may require a longer centrifugation over the density gradient medium in order to reduce contamination by hypodense granulocytes. Alternatively, the RosetteSep™ Human Granulocyte Depletion Cocktail (Catalog #15624) can be used to deplete total granulocytes before beginning the EasySep™ protocol.

If using previously frozen MNCs, 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 in recommended medium.

* 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.

SAMPLES WITH HIGH CD34+ CONTENT (> 5%)

Please contact us at techsupport@stemcell.com for protocols to further isolate CD34+ cells from samples that have been previously enriched for CD34+ cells.

Recommended Medium

EasySep™ Buffer (Catalog #20144), RoboSep™ Buffer (Catalog #20104), or PBS containing 2% fetal bovine serum and 1 mM EDTA. 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 Table 1 for detailed instructions regarding the EasySep™ procedure for each magnet.

Table 1. EasySep™ Human CD34 Positive Selection Kit 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.	- If starting with $< 2 \times 10^7$ cells, resuspend cells in 0.1 mL - If starting with $2 \times 10^7 - 2 \times 10^8$ cells, resuspend cells at 2×10^8 cells/mL - If starting with $> 2 - 5 \times 10^8$ cells, resuspend cells in 1 mL	- If starting with $< 5 \times 10^7$ cells, resuspend cells in 0.25 mL - If starting with $5 \times 10^7 - 2 \times 10^8$ cells, resuspend cells at 2×10^8 cells/mL - If starting with $2 - 5 \times 10^8$ cells, resuspend cells in 1 mL - If starting with $5 \times 10^8 - 2 \times 10^9$ cells, resuspend cells at 5×10^8 cells/mL
	Add sample to required tube.	5 mL (12 x 75 mm) polystyrene round-bottom tube (e.g. Corning Catalog #352058)	14 mL (17 x 100 mm) polystyrene round-bottom tube (e.g. Corning Catalog #352057)
2	Add Selection Cocktail to sample.	100 μ L/mL of sample	100 μ L/mL of sample
	Mix and incubate.	RT for 15 minutes	RT for 15 minutes
3	Mix Magnetic Particles.	Pipette up and down more than 5 times	Pipette up and down more than 5 times
4	Add Magnetic Particles to sample.**	50 μ L/mL of sample	50 μ L/mL of sample
	Mix and incubate.	RT for 10 minutes	RT for 10 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 < 1 mL - Top up to 10 mL for samples ≥ 1 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 off the supernatant. Remove the tube from the magnet; this tube contains the isolated cells.	Discard supernatant	Discard supernatant
7	Repeat steps as indicated.**	Steps 5 and 6, four more times (total of 5 x 5-minute separations)	Steps 5 and 6, three more times (total of 4 x 5-minute separations)
8	Resuspend cells in desired medium. Be sure to collect cells from the sides of the 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.

** To improve recovery, the Magnetic Particle addition can be increased to 100 μ L/mL and/or the magnetic separation can be reduced by 1 separation to either 4 x 5-minute or 3 x 5-minute separations, depending on the magnet used. To improve purity, an additional round of magnetic separation may be done. The improvement in purity is typically up to an additional 4%. Note that this additional separation will reduce the recovery.

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 CD34 Positive Selection Protocol

STEP	INSTRUCTIONS	RoboSep™ (Catalog #20000 and #21000)	
1	Prepare sample at the indicated cell concentration within the volume range.	- If starting with $< 5 \times 10^7$ cells, resuspend cells in 0.25 mL - If starting with $5 \times 10^7 - 2 \times 10^8$ cells, resuspend cells in 0.25 mL - 1 mL - If starting with $2 - 5 \times 10^8$ cells, resuspend cells in 1 mL - If starting with $5 \times 10^8 - 2 \times 10^9$ cells, resuspend cells in 1 - 4 mL	
	Add sample to required tube.	14 mL (17 x 100 mm) polystyrene round-bottom tube (e.g. Corning Catalog #352057)	
2	Select protocol.	- Human CD34 Positive Selection 18056-high purity - Human CD34 Positive Selection 18056-high recovery	
3	Mix Magnetic Particles.	Pipette up and down more than 5 times	
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 isolated cells and resuspend in desired medium. Be sure to collect cells from the sides of the tube.	Isolated cells are ready for use	

Notes and Tips

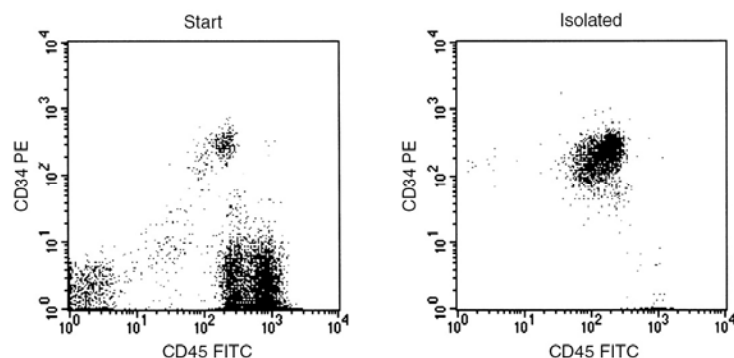
ASSESSING PURITY

The EasySep™ Human CD34 Positive Selection Cocktail uses a class II anti-CD34 antibody clone that may block some class I and II anti-CD34 antibody clones used to assess purity by flow cytometry. Alternate clones for CD34 positive selection are available as custom kits. Contact us at techsupport@stemcell.com for more information. For purity assessment of CD34+ cells by flow cytometry use one of the following class III fluorochrome-conjugated antibody clones:

- Anti-Human CD34 Antibody, Clone 581 (Catalog #60013), 8G12 (Catalog #60121), AC136, or Birma K3, and
- Anti-Human CD45 Antibody, Clone HI30 (Catalog #60018)

NOTE: Flow cytometry analysis of the positively selected cells may show slightly increased side scatter relative to the start sample.

Data



Starting with previously frozen cord blood or fresh mobilized PBMCs, the CD34+ cell content of the isolated fraction typically ranges from 84 - 99%. In the above example, the purities of the start and final isolated fractions are 1.0% and 96.0%, respectively.

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