



HLA

BUFFY COAT OR WHOLE BLOOD LYMPHOID SELECTION KIT

CATALOG #18684HLA

THIS PRODUCT INFORMATION SHEET IS PROVIDED FOR USE WITH ROBOSEP™ (SECTION A) OR "THE BIG EASY" SILVER EASYSEP™ MAGNET (SECTION B).

A) FULLY AUTOMATED PROTOCOL USING ROBOSEP™ (CATALOG #20000).

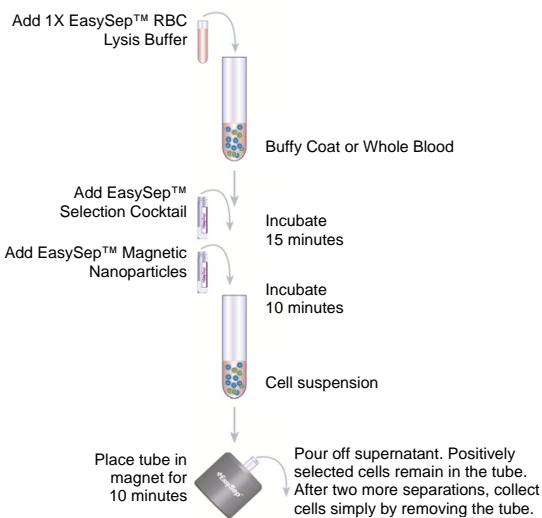
This procedure is used for processing up to 4.5 mL of buffy coat or whole blood per separation.

1. Collect whole blood in a heparinized blood collection tube. Lymphoid cells can be positively selected directly from unprocessed whole blood, or from buffy coat if preferred. If a buffy coat is required, process the collected blood as directed (see Notes and Tips, reverse side). Transfer a maximum of 4.5 mL whole blood or buffy coat to a 14 mL (17 x 100 mm) polystyrene tube. Cells must be placed in a 14 mL polystyrene tube to properly fit into the RoboSep™ carousel.

Falcon™ 14 mL Polystyrene Round-Bottom Tubes (BD Biosciences, Catalog #352057) are recommended.

2. Add 1X EasySep™ RBC Lysis Buffer (see Notes and Tips, reverse side) at a ratio of 1 part lysis buffer to 1 part sample. Mix well.
3. Select the appropriate RoboSep™ protocol:
 - Human Lymphoid WB Positive Selection 18684HLA
- If a modified RoboSep™ protocol is required, please contact STEMCELL Technologies' Technical Support at techsupport@stemcell.com.
4. Mix the EasySep™ Magnetic Nanoparticles to ensure that they are in a uniform suspension by vigorously pipetting up and down more than 5 times. Vortexing is not recommended.
5. Load the RoboSep™ carousel as directed by the on-screen prompts. When all desired quadrants are loaded, press the green "Run" button. All cell labeling and separation steps will be performed by RoboSep™.
6. When cell separation is complete, remove the tube containing the isolated cells from the magnet and resuspend cells in an appropriate amount of desired medium. Be sure to collect any cells that may be stuck to the sides of the tube. The positively selected cells are now ready for use.

MANUAL EASYSEP™ PROTOCOL DIAGRAM



FOR RESEARCH USE ONLY. NOT INTENDED FOR HUMAN OR ANIMAL DIAGNOSTIC OR THERAPEUTIC USES.

Components:

- EasySep™ HLA WB Lymphoid Positive Selection Cocktail 3 x 1.0 mL
- EasySep™ Magnetic Nanoparticles Positive Selection 3 x 1.0 mL
- EasySep™ 10X RBC Lysis Buffer 10 mL

REQUIRED EQUIPMENT:

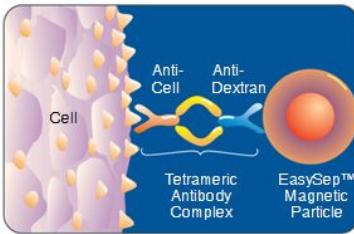
"The Big Easy" EasySep™ Magnet (Catalog #18001), or RoboSep™ (Catalog #20000).

PRODUCT DESCRIPTION AND APPLICATIONS:

EasySep™ HLA WB Lymphoid Positive Selection Cocktail and EasySep™ Magnetic Nanoparticles label CD3+ and CD19+ cells for magnetic separation. These reagents are designed to positively select CD3+ and CD19+ cells from fresh buffy coat or whole blood.

EASYSEP™ LABELING OF HUMAN CELLS:

Target cells are specifically labeled with dextran-coated magnetic particles using bispecific Tetrameric Antibody Complexes (TACs). These complexes recognize both dextran and the target cell surface antigen (Figure 1). Magnetically labeled cells are then separated from unlabeled cells using the EasySep™ procedure (reverse side).

**Figure 1.**

Schematic Drawing of EasySep™ TAC Magnetic Labeling of Human Cells.

NOTES AND TIPS:

EASYSEP™ 10X RBC LYSIS BUFFER Lysis buffer is supplied as a 10X concentrate. Prepare 1X lysis buffer at least 1 hour before use by adding 1 part 10X lysis buffer to 9 parts distilled or Type 1 water. Mix gently and completely before use.

RECOMMENDED MEDIUM The recommended medium is RoboSep™ Buffer (Catalog #20104) or phosphate-buffered saline (PBS) + 2% fetal bovine serum (FBS) (Catalog #07905) with 1 mM EDTA. Medium should be Ca++ and Mg++ free.

PREPARING A BUFFY COAT Positive selection of lymphoid cells from buffy coat uses less reagent per mL of blood and reduces donor variability (see below). Add 1 part recommended medium to 1 part whole blood. Centrifuge at 200 x g for 10 minutes at room temperature (15 - 25°C) with the break off. Remove the concentrated leukocyte band (this is the buffy coat), plus a small portion of the plasma and concentrated red blood cells, and transfer to a 14 mL polystyrene tube. The purpose of this step is to concentrate leukocytes approximately 5-fold while maintaining the same hematocrit.

DONOR VARIABILITY Certain donors express one or more soluble serum factors that can cause cross-linking with magnetic nanoparticles. This may result in visible aggregates in the enriched cell fraction following positive selection. These aggregates may appear as a distinct, high side-scatter population on FSC vs. SSC plots during flow cytometry analysis of the enriched fraction. This population consists solely of particles, with no cells or platelets present, as determined by labeling with fluorescently-conjugated antibodies against dextran, CD41 and CD45.

Potential aggregation can be avoided by preparing a buffy coat before cell separation (see above), or by washing away the donor plasma. Dilute the sample 2-fold in the recommended medium, and centrifuge at 300 x g for 10 minutes. Remove as much plasma as possible without disturbing the white and red blood cells, then resuspend sample to original volume with recommended medium before beginning the separation procedure.

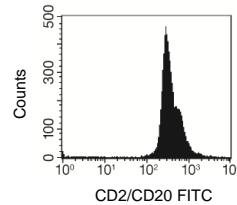
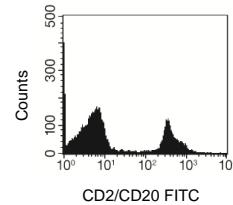
If the samples have not been washed, any aggregates can be gated out during flow cytometry analysis of the enriched fraction based on their FSC vs. SSC characteristics, or by their lack of CD45 expression.

ASSESSING PURITY The EasySep™ HLA WB Lymphoid Positive Selection Cocktail uses the anti-CD3 antibody clone UCHT-1 and the anti-CD19 antibody clone AE1. Both may block some anti-CD3 and anti-CD19 clones used to assess purity by flow cytometry. We recommend using a combination of fluorochrome-conjugated anti-CD2 (e.g. Catalog #10501) and anti-CD20 antibodies (e.g. Catalog #10510). Alternatively, a secondary fluorochrome-conjugated antibody, such as FITC-labeled sheep anti-mouse IgG can also be used to assess purity.

TYPICAL EASYSEP™ HLA LYMPHOID CELL SELECTION PROFILE:

Start: 33.1% Lymphoid Cells

Selected: 99% Lymphoid Cells



Starting with fresh whole blood, the lymphoid content of the isolated fraction (as assessed by anti-CD2 and anti-CD20 staining) typically ranges from 97.9 - 99.9%.

COMPONENT DESCRIPTIONS:**EASYSEP™ HLA WB LYMPHOID POSITIVE SELECTION COCKTAIL CODE #18684HC**

This cocktail contains a combination of monoclonal antibodies purified from hybridoma culture supernatant by affinity chromatography using Protein A or Protein G Sepharose. These antibodies are bound in bispecific TACs which are directed against CD3 or CD19, and dextran. The mouse monoclonal antibody subclass is IgG. This cocktail is supplied in PBS and contains an antibody against human Fc receptor. It should be noted that this product is a biological reagent, and as such cannot be completely characterized or quantified. Some variability is unavoidable.

EASYSEP™ MAGNETIC NANOPARTICLES POSITIVE SELECTION CODE #18150H

A suspension of magnetic dextran iron particles in water.

EASYSEP™ 10X RBC LYSIS BUFFER CODE #20110

Concentrated buffer used to lyse red blood cells prior to cell labeling and separation.

STABILITY AND STORAGE:**EASYSEP™ HLA WB LYMPHOID POSITIVE SELECTION COCKTAIL****EASYSEP™ MAGNETIC NANOPARTICLES POSITIVE SELECTION**

Product stable at 2 - 8°C until expiry date as indicated on label. Contents have been sterility tested. Do not freeze this product. This product may be shipped at room temperature (15 - 25°C), and should be refrigerated upon receipt.

EASYSEP™ 10X RBC LYSIS BUFFER

Product stable at room temperature (15 - 25°C) until expiry date as indicated on label. Once diluted 1X Lysis Buffer is stable at 2 - 8°C for 3 months. Do not freeze.

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