

StemSep® Abbreviated Procedure:

Refer to StemSep® Cell Separation Technical Manual for additional information.
www.stemcell.com/technical/28416_ssmanual.pdf

1. Prepare cells at a concentration of 5×10^7 cells/mL or within the acceptable range of $2 - 8 \times 10^7$ cells/mL in separation medium (see Notes and Tips).
2. Add normal rat serum to cell suspension at 50 μ L/mL of cells (e.g. for 1 mL of cells, add 50 μ L of serum). Incubate at 4°C (refrigerator) for 15 minutes.
3. Add StemSep® Enrichment Cocktail at 35 μ L/mL of cells (e.g. for 1 mL of cells add 35 μ L of cocktail). Mix well.
4. Incubate at 4°C (refrigerator) for 15 minutes.
5. Wash and resuspend cells ($2 - 8 \times 10^7$ cells/mL) in separation medium.
6. Add Anti-Biotin TAC at a concentration of 100 μ L/mL of cells (e.g. for 1 mL of cells add 100 μ L of anti-biotin TAC). Mix well.
7. Incubate at 4°C (refrigerator) for 15 minutes.
8. Add Magnetic Colloid at a concentration of 60 μ L/mL of cells (e.g. for 1 mL of cells add 60 μ L of colloid). Mix well.
9. Incubate at 4°C (refrigerator) for 15 minutes.
10. During incubation prepare column (refer to diagrams opposite) as follows:
 - a) Using Table 3 (see Notes and Tips) determine the appropriate column size based on cell number.
Note: Do not insert column from the front of the magnet. Lower column slowly from above down into the gap of magnet.
 - b) Gravity Feed - Place column in magnet and assemble. Prime with priming medium (see Notes and Tips) from the bottom up by depressing plunger of side syringe *slowly*.** Check for air bubbles. Proceed with 10d.
****Note:** 0.1" column is primed quickly.
 - c) Pump Feed - Place column in magnet and assemble. Prime with priming medium (see Notes and Tips) from the bottom up at appropriate speed (see Table 1). Check for air bubbles. Proceed with 10d.
 - d) Wash from the top down with appropriate volume of separation medium (see Table 2).
Note: Do not let column run dry at anytime during priming, loading or washing of the column.
11. Load sample.
12. Wash from the top down with separation medium, collecting the sample volume plus the appropriate column wash volume as flowthrough (see Table 2). The enriched cells are now ready for use.

Table 1. Flow Rates and Pump Settings



Column Size	Priming		Loading Sample and Washing	
	mL/min	pump setting*	mL/min	pump setting*
0.6"	0.6	3.0	2.0	10.0
0.5"	0.3	1.5	1.0	5.0
0.3"	0.2	1.0	0.6	3.0

*Pump setting for 4-channel pump supplied by StemCell Technologies only.

Table 2. Column Wash Volume

Column Size	Column Wash Volume
0.6"	25 mL
0.5"	15 mL
0.3"	8 mL
0.1"	1.5 mL

PROCEDURE

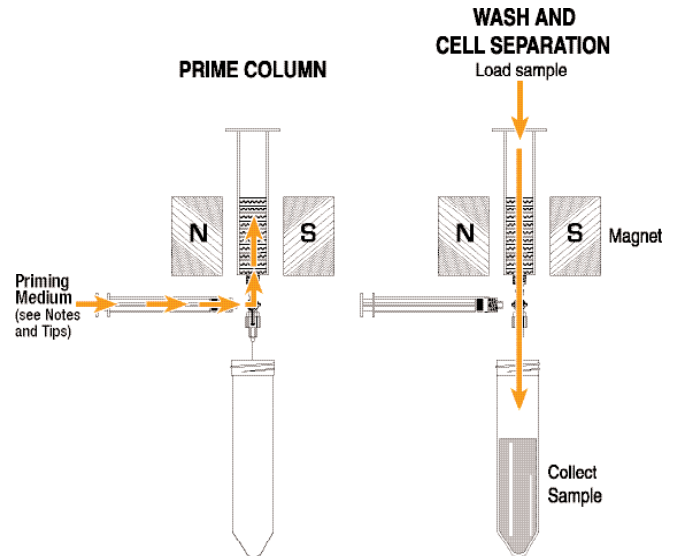
Version 3.0.1

Negative Selection

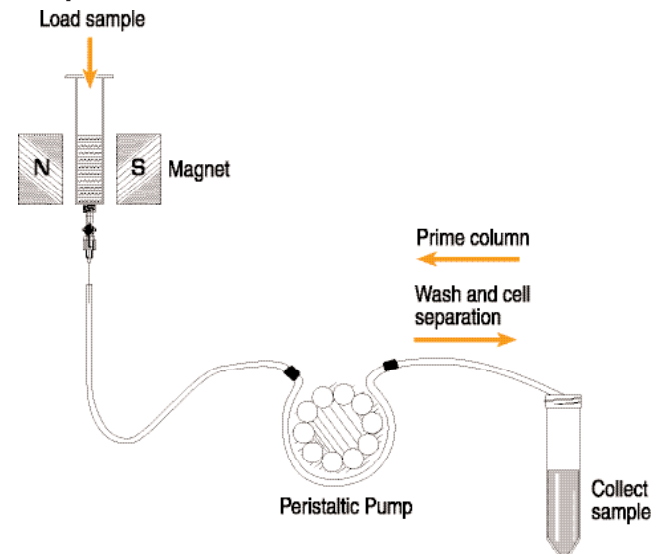
Mouse CD4⁺ T Cell Enrichment Kit

CATALOG #13052 (13062)

Gravity feed:



Pump feed:



StemCell Technologies

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FOR RESEARCH USE ONLY

#28840

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Catalog #13052 (13062)For labeling 10^9 (5×10^9) total cells**Components:**

• StemSep® Mouse CD4 ⁺ T Cell Enrichment Cocktail	700 μ L (5 x 700 μ L)
• Anti-Biotin TAC	2.0 mL (5 x 2.0 mL)
• Magnetic Colloid	1.5 mL (5 x 1.5 mL)
• Rat Serum	2.0 mL (5 x 2.0 mL)

**Product Information Sheet****REQUIRED EQUIPMENT:**

StemSep® Magnet (Catalog #11030, 11050, 11060, or 11070) or a magnet with the strength of at least 0.5 Tesla, and StemSep® Negative Selection Columns (see Table 3, Notes and Tips).

PRODUCT DESCRIPTION AND APPLICATIONS:

The StemSep® Mouse CD4⁺ T Cell Enrichment Kit is designed to highly enrich CD4⁺ T cells from suspensions of mouse spleen cells. The recovered CD4⁺ T cells have not been labeled with antibody.

STEMSEP® LABELING OF MOUSE CELLS:

Antigens on the surface of unwanted cells are first labeled with biotinylated monoclonal antibodies (Figure 1). Cells are then linked to magnetic dextran iron particles using bispecific Tetrameric Antibody Complexes (TAC). These complexes recognize both biotin and dextran on the magnetic colloid/nanoparticle (Figure 1). Magnetically labeled cells are then separated from unlabeled cells by passing them through a magnetic separation column placed in a magnet.

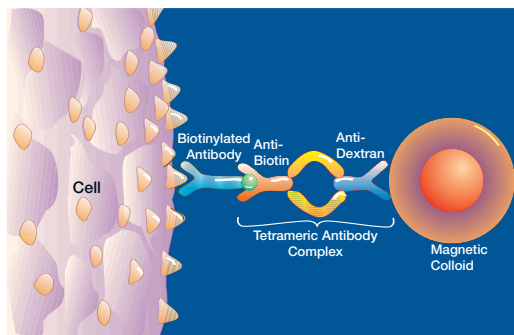


Figure 1.
Schematic Drawing of StemSep® TAC Magnetic Labeling of Mouse Cells.

NOTES AND TIPS:**Column Preparation:**

- Use the appropriate column size (see Table 3).
- Check all the connections during priming and washing to ensure they do not leak.
- Prime the column from the bottom up.
- Use priming medium (see below) to prime the column.
- Ensure that there are no air bubbles in the column.
- Use separation medium (see below) to wash the column. The protein present in the separation medium prevents cells from binding non-specifically to the column.
- Ensure that the column does not run dry at any time.

Preparing a Single Cell Suspension. Spleen: Disrupt spleen into 5 mL of separation medium. Further disperse clumps by gently pipetting up and down several times. Remove remaining clumps of cells and debris by passing cell suspension through a 70 μ m mesh nylon strainer. Centrifuge, discard supernatant and resuspend cells at 5×10^7 /mL (a range of $2 - 8 \times 10^7$ /mL is acceptable) in separation medium.

Recommended Media: Using degassed media reduces the chance of developing air bubbles in the column. Air bubbles cause channeling in the column reducing the capacity of the column and potentially compromising purity. Media should be Ca^{++} and Mg^{++} free.

- **Priming Medium:** Use PBS (Catalog #37350), either at room temperature or degassed, without serum or other protein.
- **Separation Medium:** Use PBS + 2% FBS (Catalog #07905) or Hank's (Catalog #37250) + 2% FBS.

Assessing Purity. Purity of CD4⁺ T cells can be measured using flow cytometry by staining with a fluorochrome-conjugated anti-CD4 antibody (e.g. anti-CD4 FITC antibody, Catalog # 10702).

Table 3. Column Capacity: Recommended Number of Mouse Nucleated Cells in the Start Suspension for Various Column Sizes

Column Size	Catalog # Gravity	Catalog # Pump	Column Capacity Based on Cell Number	Will Fit Magnet Size
0.6"	12061	12062	$10^9 - 1.5 \times 10^9$	green, blue, black
0.5"	12051	12052	$5 \times 10^7 - 3 \times 10^8$	green, blue, black
0.3"	12031	12032	$2 \times 10^7 - 10^8$	all sizes
0.1"	12021	-	$10^5 - 2 \times 10^7$	red, green

TYPICAL STEMSEP® MOUSE CD4⁺ T CELL ENRICHMENT PROFILE:Start: 21% CD4⁺ CellsEnriched: 91% CD4⁺ Cells

The CD4⁺ T cell content of the enriched fraction typically ranges from 80 to 94%.

COMPONENT DESCRIPTIONS:**StemSep® Mouse CD4⁺ T Cell Enrichment Cocktail** code #13052C

This cocktail contains a combination of biotinylated monoclonal antibodies purified from rat ascites fluid or hybridoma culture supernatant by affinity chromatography using Protein A or Protein G Sepharose. These antibodies are directed against cell surface antigens on mouse hematopoietic cells (CD8, CD11b (Mac-1), CD45R/B220, Ly-6G/C (Gr-1), TER119). It should be noted that this product is a biological reagent, and as such cannot be completely characterized or quantified. Some variability is unavoidable. Supplied in phosphate buffered saline with 0.1% BSA.

Anti-Biotin TAC code #13050C

A combination of mouse and rat monoclonal antibodies purified using Protein G Sepharose. The monoclonal antibody subclass is IgG₁. The antibodies form tetrameric antibody complexes (TAC) which are directed against both biotin and dextran. Supplied in phosphate buffered saline.

Magnetic Colloid code #10051

A colloidal suspension of magnetic dextran iron particles in USP saline, pH 7.0 – 7.5.

Rat Serum code #13551

Serum to prevent non-specific binding of rat antibodies to mouse cells. Certified mycoplasma-free, sterile if unopened.

STABILITY AND STORAGE:**StemSep® Mouse CD4⁺ T Cell Enrichment Cocktail and Anti-Biotin TAC**

Stable at 4°C for one year and two years, respectively. Do not freeze these products. Contents sterile in unopened tube.

Magnetic Colloid

Stable at -20°C for one year. Stable at 4°C for six weeks. Contents sterile in unopened tube. This product may be shipped frozen or at room temperature and must be frozen or refrigerated upon receipt. **Thaw and vortex before storing at -20°C if not completely frozen upon receipt.** Repeated freezing and thawing is possible but not recommended. Vortex before re-freezing.

Rat Serum

Stable for at least two years when stored at -20°C. Stable for two months when stored at 4°C. Contents sterile in unopened tube.

See Material Safety Data Sheet for more information.

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