



MojoSort™ Mouse CD45 Nanobeads Protocol - Depletion 👄

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#### ABSTRACT

**Product description and procedure summary:** The cells targeted by the Nanobeads are depleted by incubating your sample with the directly conjugated magnetic particles. The magnetically labeled fraction is retained by the use of a magnetic separator. After collection of the targeted cells, downstream applications include functional assays, gene expression, phenotypic characterization, etc.

**Note:** This procedure is optimized for the isolation of  $10^7$  to  $2 \times 10^8$  cells per tube. If working with fewer than  $10^7$  cells, keep volumes as indicated for  $10^7$  cells. For best results, optimize the conditions to your specific cell number and tissue. Prepare fresh MojoSort<sup>™</sup> Buffer solution by diluting the 5X concentrate with sterile distilled water. *Scale up volumes if using 14mL tubes and Magnet, and place the tube in the magnet for 10 minutes.* 

**EXTERNAL LINK** 

https://www.biolegend.com/protocols/mojosort-mouse-cd45-nanobeads-protocol-depletion/4751/

### **GUIDELINES**

MojoSort™ magnetic particles can be used with other commercially available magnetic separators, both free standing magnets and column-based systems. Because MojoSort™ protocols are optimized for the MojoSort™ separator, the protocols may need to be adjusted for other systems. Please contact BioLegend Technical Service (tech@biolegend.com) for more information and guidance. We do not recommend using MojoSort™ particles for BD's IMag™ or Life Technologies' DynaMag™.

**Application notes:** To use this product in magnetic separation columns, a titration of the Nanobeads should be performed. Optimal concentration for magnetic separation columns is lot-specific. Please contact BioLegend Technical Service (tech@biolegend.com) for further assistance on how to use MojoSort™ Nanobeads in magnetic separation columns.

# MATERIALS

NAME ~	CATALOG #	VENDOR V
MojoSort™ Magnet	480019	BioLegend
MojoSort™ Buffer	480017	BioLegend
MojoSort™ Mouse CD45 Nanobeads	480027, 480028	BioLegend

## MATERIALS TEXT

- Adjustable pipettes
- 70µm filters (one per sample)
- 5mL (12 x 75mm) or 14mL (17 x 100mm) polypropylene tubes
- Reagents for sample preparation
- Reagents and instruments (Flow cytometer) to determine yield and purity

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- In the final wash of your sample preparation, resuspend the cells in MojoSort™ Buffer by adding up to 4 mL in a 5 mL (12 x 75 mm) polypropylene tube.

**Note:** Keep MojoSort™ Buffer on ice throughout the procedure.

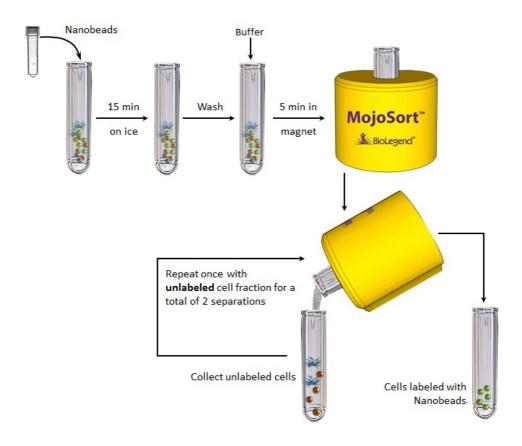
Prepare cells from your tissue of interest or blood without lysing erythrocytes.

- 3 Filter the cells with a 70µm cell strainer, centrifuge at 300xg for 5 minutes, and resuspend in an appropriate volume of MojoSort™ Buffer. Count and adjust the cell concentration to 1 x 10<sup>8</sup> cells/mL.
- 4 Aliquot 100  $\mu$ L of cell suspension (10<sup>7</sup> cells) into a new tube.
- 5 Resuspend the beads by vortexing, maximum speed, 5 touches. Add **10μL of Antibody Nanobeads**. Mix well and **incubate on ice for 15 minutes**. Scale up the volume accordingly if separating more cells. For example, add 100 μL of Nanobeads for separating 1 x 10<sup>8</sup> cells in 1 ml of MojoSort™ Buffer. When working with less than 10<sup>7</sup> cells, use indicated volumes for 10<sup>7</sup> cells.
- 6 Wash the cells by adding MojoSort™ Buffer up to 4mL. Centrifuge the cells at 300xg for 5 minutes.
- 7 Discard the supernatant.
- 8 Add 2.5mL of MojoSort™ Buffer.
  Note: If you observe aggregates, filter the suspension. To maximize yield, you can disrupt the aggregates by pipetting the solution up and down.
- 9 Place the tube in the magnet for 5 minutes.

  Optional: Take a small aliquot before placing the tube in the magnet to monitor purity and yield. Keep unused cells to be used as control or other applications if needed.
- 10 Pour out the unlabeled fraction (THESE ARE YOUR CELLS OF INTEREST). Set aside the labeled fraction.

11 Place the unlabeled fraction tube in the magnet for 5 minutes. This will be a total of 2 separations (once on the labeled cell mixture and once on the unlabeled fraction). Pool the unlabeled fractions. The labeled fractions may be useful as staining controls, to monitor purity/yield, or other purposes.

Optional: Take a small aliquot to monitor purity and yield.



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