



SOLIDEX[®]-ISOEx Human CD8 Isolation Nanobeads (Column-Based)

Product Instruction Manual

Cat. No.	GM-Tg-hg-TA040-iso-nanoIMB
Product Name (Column-Based)	SOLIDEX [®] -ISOEx Human CD8 Isolation Nanobeads
Storage Temperature	2-8 °C (Do not freeze)

Technical Parameters

Classification	Description
Cell type	CD8+ T Cell
Isolation method	Positive selection
Labeling type	Direct
Magnetic nanobeads type	Non-releasable
Format	Column-based

Product Introduction

CD8+ T cells (cytotoxic T cells) are key effector cells of adaptive immunity, playing a central role in antiviral and anti-tumor immunity by directly killing virus-infected or cancerous cells. Their primary surface marker is CD8, which is co-expressed with CD3 (CD3+CD4-CD8+). Depending on the application scenario, we support two isolation strategies: positive selection and depletion.

By utilizing SOLIDEX[®]-ISOEx Human CD8 Isolation Nanobeads, column-based isolation achieves superior cell purity. Through the specific capture facilitated by the magnetic matrix within the column, high-purity target cells can be effectively obtained. This method is highly compatible with automated workflows. High-purity CD8+ T cells obtained post-isolation facilitate various downstream applications, such as viral infection research—specifically analyzing the role of CD8+ T cells in antiviral immune responses—and supporting applications like CAR-T cell therapy.

Product Components and Specifications

Component	Catalog No.	10 Test (1×10 ⁸ cells)	50 Tests (5×10 ⁸ cells)	100 Tests (1×10 ⁹ cells)
SOLIDEX [®] -ISOEx Human CD8 Isolation Nanobeads (Column-Based)	GM-Tg-hg-TA040-iso-nanoIMB	50 µL	250 µL	0.5 mL

Note: This product is for research and development use only.

Shelf Life: Store at 2-8°C, protected from light, and do not freeze. Under these conditions, the product is valid for 6 months.

Reagents and Equipment Required

A. Cell isolation Column

For positive selection or depletion, recommended for use with GeneMedi SOLIDEX[®]-ISOEx cell isolation columns: M Column for standard throughput, Cat. No.: **GMP-ISOEx-Column-M**; L Column for high throughput, Cat. No.: **GMP-ISOEx-Column-L**. Comparable columns from other mainstream brands are also compatible.

B. Magnetic Separator

C. Cell isolation buffer

Phosphate-buffered saline (PBS), pH 7.2, containing 0.5% bovine serum albumin (BSA) and 2 mM EDTA. (user-supplied).

Note:

- (1) BSA can be replaced by other proteins such as human serum albumin (HSA), human serum, or fetal bovine serum (FBS).
- (2) Degas the Cell isolation buffer before use, as air bubbles may block the Column.
- (3) Keep the Cell isolation buffer cold (2-8°C).

Protocol

A. Cell Labeling

- a. Count the peripheral blood mononuclear cells (PBMCs). Add 200 μ L of cell isolation buffer per 1×10^7 cells and resuspend the cells.
- b. Add 5 μ L of SOLIDEX[®]-ISOEx Human CD8 Isolation Nanobeads per 1×10^7 cells. Mix gently but thoroughly, and incubate at 2-8°C for 10 minutes.
- c. Add 1 mL of cell isolation buffer per 1×10^7 cells. Centrifuge and discard the supernatant. Resuspend the cell pellet in 3 mL of cell isolation buffer. The cells are now ready for the subsequent cell isolation step.

Note:

- a. The nanobeads must be thoroughly mixed prior to use by pipetting up and down.
- b. The cell isolation buffer must be pre-cooled to 2-8°C or on ice.

B. Cell Isolation

- a. Place the LS isolation column on the magnetic separator and rinse with 3 mL of cell isolation buffer.
- b. Once the cell isolation buffer has completely drained, apply the 3 mL cell suspension to the column. Place a clean 15 mL centrifuge tube underneath to collect the flow-through.
- c. After the liquid has drained from the column, wash the column once by adding 5 mL of cell isolation buffer. Collect the flow-through in the same centrifuge tube; this fraction contains the non-CD8+ T cells.
- d. Remove the isolation column from the magnetic separator and place it into a clean 15 mL centrifuge tube. Add 5 mL of cell isolation buffer and use the plunger to directly flush the liquid out of the column; this collected fraction contains the CD8+ T cells.

Notes

- A. Avoid freezing during use and storage of the Nanobeads.
- B. It is recommended to use low-binding pipette tips and centrifuge tubes to prevent loss of Nanobeads due to adsorption.
- C. Before aspirating the Nanobeads, mix them gently. Avoid bubble formation during mixing.
- D. This product is used for research use only.

Validation Data from GeneMedi

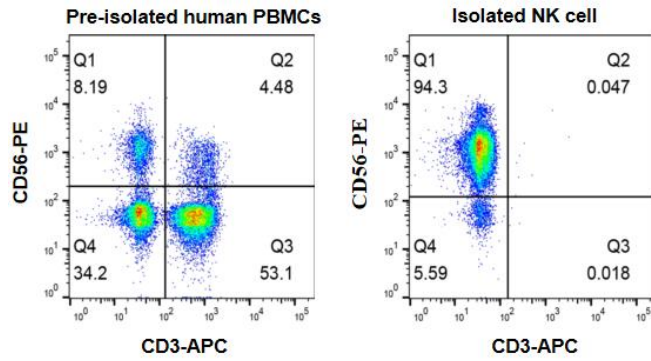


Figure 1. Isolation of high-purity T cells achieved by the SOLIDEX®-ISOEx Human CD8 Isolation Nanobeads (Column-based). Figure 1. Isolation of high-purity T cells achieved by SOLIDEX®-ISOEx Human CD8 Isolation Nanobeads (Column-based): To evaluate the purity of the isolated T cells, CD8⁺ T cells were isolated from human peripheral blood mononuclear cells (PBMCs). Cells pre- and post-isolation were labeled with CD3-APC and CD8-PE antibodies for flow cytometric analysis. The purity of CD8⁺ T cells pre-isolation and post-isolation was 16.6% and 94.6%, respectively.