



Version 2

Working

Th9 Polarization of Mouse CD4+ Cells [↗](#)

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EXTERNAL LINK

<https://www.biolegend.com/protocols/th9-polarization-of-mouse-cd4--cells-protocol/4243/>

PROTOCOL STATUS

Working

MATERIALS TEXT


- Sterile PBS
- Cell culture medium (IMDM supplemented with 10% FBS)
- Sterile plastic petri dishes
- RBC Lysis Buffer (Cat. No. [420301](#))
- Anti-mouse CD3ε, clone 145-2C11 (LEAF™ format, Cat. No. [100314](#))
- Anti-mouse CD28, clone 37.51, (LEAF™ format, Cat. No. [102112](#))
- Mouse MojoSort™ CD4 T-cell Isolation Kit (Cat. No. [480005](#))
- Anti-mouse IFN-γ, clone XMG1.2, (LEAF™ format, Cat. No. [505812](#))
- Recombinant mouse IL-2 (carrier-free) (Cat. No. [575402](#))
- Recombinant mouse IL-4 (carrier-free) (Cat. No. [574302](#))
- Recombinant human TGF-β1 (carrier-free) (Cat. No. [580702](#))
- Monensin Solution (Cat. No. [420701](#))
- PMA (Phorbol 12-myristate 13-acetate) (Cat. No. P8139 from Sigma)
- Ionomycin (Cat. No. I0634 from Sigma)

Isolation of CD4+ Cells From Lymph Nodes

- 1 Harvest lymph nodes (superficial cervical, mandibular, axillary, inguinal, and mesenteric) from mice.
- 2 Tease lymph nodes through a sterile 70-μm nylon cell strainer to obtain single-cell suspensions in complete IMDM containing 10% FCS (complete medium).
- 3 Resuspend cells in complete medium and use your favorite method to isolate CD4⁺ cells. Consider using our [MojoSort™ Mouse CD4 T Cell Isolation Kit](#).

Th9 Polarization of CD4+ Cells:

- 4 On day 0, coat 60 x 15mm of plastic petri dishes with anti-mouse CD3ε, clone 145-2C11 (5μg/ml). Incubate at 37°C for 2 hours or 4°C overnight. Aseptically decant antibody solution from the plate. Wash plate 3 times with sterile PBS. Discard liquid. ⌚ 02:00:00
- 5 Plate CD4+ cells at 10 x 10⁶/5 ml/dish. Culture cells for 3 days in the presence of anti-mouse CD28, clone 37.51 (5μg/mL), recombinant human TGF-β1 (10ng/mL), recombinant mouse IL-4 (10ng/mL), recombinant mouse IL-2 (20ng/mL), and anti-mouse IFN-γ, clone XMG1.2 (10μg/mL).

6 On day 3, wash cells once and then restimulate in complete medium with 500ng/ml PMA and 500ng/mL ionomycin, in the presence of monensin for 6 hours.  06:00:00

7 After harvesting, the cells are ready for staining.

Note: Recombinant human TGF- β is effective for stimulating mouse cells.



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