



**Mouse Splenocytes  
(MS)**  
Catalog #M5540-57

**Cell Specification**

The spleen stores and purifies erythrocytes, metabolizes hemoglobin, and recycles iron. The spleen also provides a critical function for the immune system by mounting a primary immune response to antigens in the blood and synthesizing antibodies [1]. Splenocytes are mononuclear cells derived from the spleen and include T-lymphocytes, B-lymphocytes, NK-cells, and NK T-cells. Primary mouse splenocytes (MS) can be used to isolate CD4+ T-cells, CD8+ T-cells and CD45R+ B cells.

MS from ScienCell Research Laboratories are isolated from normal adult C57BL/6 mouse spleen. MS are depleted of splenic macrophages, cryopreserved after isolation, and delivered frozen. Each vial contains at least 10 million cells in 1 ml volume. MS are quality control tested for viability. MS are negative for mycoplasma, bacteria, yeast, and fungi. MS can be maintained for a limited period of time in culture using the conditions provided by ScienCell Research Laboratories and *are not intended for long-term culture*.

**Recommended Medium**

It is recommended to use HematoGro Medium (HeGM, Cat. #5501) for short-term maintenance of MS *in vitro*.

**Product Use**

MS are for research use only. They are not approved for human or animal use, or for application in *in vitro* diagnostic procedures.

**Storage**

Upon receiving, directly and immediately transfer the cells from dry ice to liquid nitrogen and keep the cells in liquid nitrogen until they are needed for experiments.

**Shipping**

Dry ice.

**References**

[1] Mebius R, Kraal G. (2005) "Structure and function of the spleen." *Nat Rev Immunol* 5(8): 606-616.

## Instructions for culturing cells

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**Caution:** Cryopreserved cells are very delicate. Thaw the vial in a 37°C water bath and return the cells to culture as quickly as possible with minimal handling!

*Note: Experiments should be well organized before thawing MS. It is recommended that MS are purified or used for experiments as quickly as possible after thawing the cells. Cells are not intended for long term culture.*

### Initiating the culture:

1. Prepare complete medium (HeGM, Cat. #5501). Thaw HeGS and P/S solution at 37°C. Gently tilt the tubes several times to ensure the contents are completely mixed before adding to the medium. Spray the medium bottle and tubes with 70% ethanol, and wipe to remove excess liquid. In a sterile field, remove the caps without touching the interior threads with fingers. Add HeGS and P/S solution to the medium and mix well.
2. Add 15 ml of complete medium to a T-75 flask. Leave the vessel in the sterile field and proceed to thaw the cryopreserved cells.
3. Place the frozen vial in a 37°C water bath. Hold and rotate the vial gently until the contents completely thaw. Remove the vial from the water bath promptly, wipe it down with 70% ethanol and transfer it to the sterile field.
4. Remove the cap carefully without touching the interior threads. Gently resuspend and dispense the contents of the vial into the equilibrated, culture vessel.

*Note: Dilution and centrifugation of cells after thawing are not recommended since these actions are more harmful to the cells than the effect of residual DMSO in the culture.*

5. Replace the cap or lid, and gently rock the vessel to distribute the cells evenly. Loosen cap if necessary to allow gas exchange.
6. Return the culture vessel to the incubator.
7. Cells should be promptly purified and used to isolate CD4+ T-cells, CD8+ T-cells and CD45R+ B cells.

**Caution:** Handling animal derived products is potentially biohazardous. Always wear gloves and safety glasses when working with these materials. Never mouth pipette. We recommend following the universal procedures for handling products of human origin as the minimum precaution against contamination [1].

[1] Grizzle WE, Polt S. (1988) "Guidelines to avoid personal contamination by infective agents in research laboratories that use human tissues." *J Tissue Culture Methods*. 11: 191-9.