



Rat Sertoli Cells (RSerC)

Catalog #R4520

Cell Specification

Sertoli cells (SerC) are sustentacular cells that line the basal region of the seminiferous tubules. These epithelial supporting cells adopt a pyramid-shaped morphology. SerC directly interacts with the germ cells and provide morphological and nutritional support for spermatogenesis. During spermiogenesis, the final step in spermatogenesis, SerC play a pivotal role in the degradation of residual body that were removed from the mature spermatid. Embryologically, following the determination of the testes, SerC prevent the development of female sex organs through the secretion of Mullerian Inhibiting Factor (1). Tight junctions formed between adjacent SerC make up the blood-testis barrier (BTB) or SerC barrier that is unique to the testis. This physical barrier plays an immunoprotective role in the testis by sequestering the spermatocytes and spermatids (2). The BTB also serves as a crucial barrier to prevent the introduction of external factors into the seminiferous tubules. Understanding the physiology and functions of Rat Sertoli Cells (RSerC) can provide functional insights into the regulation of spermatogenesis and male fertility.

RSerC from ScienCell Research Laboratories are isolated from CD® IGS rat testes. RSerC are cryopreserved at Passage 1 and delivered frozen. Each vial contains $>5 \times 10^5$ cells in 1 ml volume. RSerC are characterized by immunofluorescence with antibodies specific to GATA4 and SOX9. RSerC are negative for mycoplasma, bacteria, yeast, and fungi. RSerC are guaranteed to further culture under the conditions provided by ScienCell Research Laboratories; *however, RSerC are not recommended for long-term cultures due to limited expansion capacity and senescence after subculturing.*

Recommended Medium

It is recommended to use Sertoli Cell Medium (SerCM, Cat. #4521) for culturing RSerC *in vitro*.

Product Use

RSerC 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] Carlsen E, Olsson C, Petersen JH, Andersson AM, Skakkebaek NE. (1999) "Diurnal rhythm in serum levels of inhibin B in normal men: relation to testicular steroids and gonadotropins." *J Clin Endocrinol Metab.* 84(5): 1664-9.
- [2] Washburn RL, Hibler T, Kaur G, Dufour JM. (2022) "Sertoli Cell Immune Regulation: A Double-Edged Sword." *Front Immunol.* 13: 913502.

Instructions for culturing cells

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: RSerC are very sensitive cells and they are not expected to proliferate many times in culture. Experiments should be well organized before thawing the cells. It is recommended that RSerC are used for experiments at earliest passage after initial plating with minimal expansion. If subculture is inevitable, follow the instructions below with special care and it is recommended that the cells only be subcultured once.

Initiating the culture:

Note: ScienCell primary cells must be cultured in a 37°C, 5% CO₂ incubator. Cells are only warranted if ScienCell media and reagents are used and the recommended protocols are followed.

1. Prepare a poly-L-lysine-coated culture vessel (2 µg/cm², T-75 flask is recommended). To obtain a 2 µg/cm² poly-L-lysine-coated culture vessel, add 10 ml of sterile water to a T-75 flask and then add 15 µl of poly-L-lysine stock solution (10 mg/ml, Cat. #0413). Leave the vessel in a 37°C incubator overnight (or for a minimum of one hour).
2. Prepare complete medium. Decontaminate the external surfaces of medium bottle and medium supplement tubes with 70% ethanol and transfer them to a sterile field. Aseptically transfer supplement to the basal medium with a pipette. Rinse the supplement tube with medium to recover the entire volume.
3. Rinse the poly-L-lysine-coated vessel twice with sterile water and then add 20 ml of complete medium. Leave the vessel in the sterile field and proceed to thaw the cryopreserved cells.
4. Place the frozen vial in a 37°C water bath. Hold and rotate the vial gently until the contents completely thaw. Promptly remove the vial from the water bath, wipe it down with 70% ethanol, and transfer it to the sterile field.
5. Carefully remove the cap without touching the interior threads. Gently resuspend and dispense the contents of the vial into the equilibrated, poly-L-lysine-coated 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. It is also important that cells are plated in poly-L-lysine-coated culture vessels to promote cell attachment.
6. Replace the cap or lid of the culture vessel and gently rock the vessel to distribute the cells evenly. Loosen cap, if necessary, to allow gas exchange.
7. Return the culture vessel to the incubator.
8. Do not disturb the culture for at least 16 hours after the culture has been initiated. Refresh culture medium the next day to remove residual DMSO and unattached cells.

Maintaining the culture:

1. Refresh supplemented culture medium the next morning after establishing a culture from cryopreserved cells.
2. Change the medium every three days thereafter, until the culture is approximately 70% confluent.
3. Once the culture reaches 70% confluency, change medium every two days until the culture is approximately 90% confluent.

Subculturing:

1. Subculture when the culture reaches 90-95% confluency.
2. Prepare poly-L-lysine-coated culture vessels ($2\ \mu\text{g}/\text{cm}^2$) one day before subculture.
3. Warm complete medium, trypsin/EDTA solution, 0.05% (T/E, Cat. #0183), T/E neutralization solution (TNS, Cat. #0113), and DPBS (Ca^{++} - and Mg^{++} -free, Cat. #0303) to **room temperature**. We do not recommend warming reagents and medium in a 37°C water bath prior to use.
4. Rinse the cells with DPBS.
5. Add 5 ml DPBS and 5 ml 0.05% T/E solution (Cat. #0183) into flask (in the case of a T-75 flask). Gently rock the flask to ensure complete coverage of cells by T/E solution. Use a microscope to monitor the change in cell morphology.

Note: We recommend using ScienCell 0.05% T/E solution which is optimized to minimize cell damage due to over trypsinization. If 0.25% T/E solution (Cat. #0103) is used, then 9 ml of DPBS and 1 ml of 0.25% T/E solution should be used.

Caution: Do NOT use undiluted trypsin when subculturing primary cells.

6. During incubation, prepare a 50 ml conical centrifuge tube with 5 ml of fetal bovine serum (FBS, Cat. #0500).
7. Once cells completely round up, transfer T/E solution from the flask to the 50 ml centrifuge tube (a small percent of cells may detach) and continue to incubate the flask at 37°C for another minute (no solution in the flask at this time).
8. At the end of incubation, gently tap the side of the flask to dislodge cells from the surface. Check under a microscope to make sure that all cells detach.
9. Add 5 ml of TNS solution to the flask and transfer detached cells to the 50 ml centrifuge tube. Rinse the flask with another 5 ml of TNS to collect the residual cells.
10. Examine the flask under a microscope for a successful cell harvest by looking at the number of cells being left behind; there should be less than 5%.
11. Centrifuge the 50 ml centrifuge tube at 1000 rpm for 5 minutes. Gently resuspend cells in culture medium.
12. Count and plate cells in a new poly-L-lysine-coated culture vessel with the recommended cell density. A seeding density of $10,000\ \text{cells}/\text{cm}^2$ is recommended.

Note: We do not recommend cryopreservation of primary cells by the end user. Refreezing cells may damage them and affect cell performance. ScienCell does not guarantee primary cells cryopreserved by the end user.

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 Cult Methods*. 11: 191-9.