



# WT SV40 MEF

CRL-2907™

## Description

WT SV40 MEF is a fibroblast cell line that was isolated from a mouse embryo. It can be paired with MEF cell lines, generated from BCL-2 family member knockout mice (eg, ATCC CRL-2908, ATCC CRL-2909, ATCC CRL-2910, ATCC CRL-2911, ATCC CRL-2912, ATCC CRL-2913), as a control to study the molecular mechanisms of cell apoptosis and the BCL-2 family signaling pathway. This cell line was deposited by S Korsmeyer.

**Organism:** *Mus musculus*, mouse

**Cell Type:** fibroblast

**Tissue:** Embryo

**Morphology:** fibroblast-like

**Growth properties:** Adherent

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## Storage Conditions

**Product format:** Frozen

**Storage conditions:** Vapor phase of liquid nitrogen

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## Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic use, any human or animal consumption, or any diagnostic use.

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## BSL 2

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories*

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ATCC highly recommends that appropriate personal protective equipment is always used when handling vials. For cultures that require storage in liquid nitrogen, it is important to note that some vials may leak when submersed in liquid nitrogen and will slowly fill with liquid nitrogen. Upon thawing, the conversion of the liquid nitrogen back to its gas phase may result in the vial exploding or blowing off its cap with dangerous force creating flying debris. Unless necessary, ATCC recommends that these cultures be stored in the vapor phase of liquid nitrogen rather than submersed in liquid nitrogen.

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## Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at [www.atcc.org](http://www.atcc.org).

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## Growth Conditions

**Temperature:** 37°C

**Atmosphere:** 95% Air, 5% CO<sub>2</sub>

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## Handling Procedures

### Unpacking and storage instructions:

1. Check all containers for leakage or breakage.

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2. Remove the frozen cells from the dry ice packaging and immediately place the cells at a temperature below  $-130^{\circ}\text{C}$ , preferably in liquid nitrogen vapor, until ready for use.

**Complete medium:** The base medium for this cell line is ATCC-formulated IMDM Catalog No. 30-2005. To make the complete growth medium, add the following components to the base medium: 10 % Fetal Bovine Serum and 1x Non-essential amino acids.

**Handling Procedure:**

To insure the highest level of viability, thaw the vial and initiate the culture as soon as possible upon receipt. If upon arrival, continued storage of the frozen culture is necessary, it should be stored in liquid nitrogen vapor phase and not at  $-70^{\circ}\text{C}$ . Storage at  $-70^{\circ}\text{C}$  will result in loss of viability.

1. Thaw the vial by gentle agitation in a  **$37^{\circ}\text{C}$**  water bath. To reduce the possibility of contamination, keep the O-ring and cap out of the water. Thawing should be rapid (approximately 2 minutes).
2. Remove the vial from the water bath as soon as the contents are thawed, and decontaminate by dipping in or spraying with 70% ethanol. All of the operations from this point on should be carried out under strict aseptic conditions.
3. Transfer the vial contents to a centrifuge tube containing 9.0 mL complete culture medium. and spin at approximately  $125 \times g$  for 5 to 7 minutes.
4. Resuspend cell pellet with the recommended complete medium (see the specific batch information for the culture recommended dilution ratio). and dispense into a new culture flask. It is important to avoid excessive alkalinity of the medium during recovery of the cells. It is suggested that, prior to the addition of the vial contents, the culture vessel containing the complete growth medium be placed into the incubator for at least 15 minutes to allow the medium to reach its normal pH (7.0 to 7.6). pH (7.0 to 7.6).
5. Incubate the culture at  **$37^{\circ}\text{C}$**  in a suitable incubator. A 5%  $\text{CO}_2$  in air atmosphere is recommended if using the medium described on this product sheet.

**Subculturing procedure:** Volumes used in this protocol are for  $75 \text{ cm}^2$  flasks; proportionally reduce or increase amount of dissociation medium for culture vessels of other sizes.

1. Remove and discard culture medium.
2. Briefly rinse the cell layer with  $\text{Ca}^{++}/\text{Mg}^{++}$  free Dulbecco's phosphate-buffered

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saline (D-PBS) or 0.25% (w/v) Trypsin - 0.53 mM EDTA solution to remove all traces of serum which contains trypsin inhibitor.

3. Add 1.0 to 2.0 mL of Trypsin-EDTA solution to flask and observe cells under an inverted microscope until cell layer is dispersed (usually within 5 to 15 minutes).

Note: To avoid clumping do not agitate the cells by hitting or shaking the flask while waiting for the cells to detach. Cells that are difficult to detach may be placed at 37.0°C to facilitate dispersal.

4. Add 6.0 to 8.0 mL of complete growth medium and aspirate cells by gently pipetting.
5. Add appropriate aliquots of the cell suspension to new culture vessels. An inoculum of  $5 \times 10^3$  to  $1 \times 10^4$  viable cells/cm<sup>2</sup> is recommended.
6. Incubate cultures at 37°C. At subculture, cell concentration is between  $1 \times 10^5$  to  $2 \times 10^5$  cells/cm<sup>2</sup>.

**Subcultivation ratio:** A subcultivation ratio of 1:8 to 1:20 is recommended.

**Medium renewal:** Every 2 to 3 days

**Reagents for cryopreservation:** Fetal Bovine Serum supplemented with 10% (v/v) DMSO (ATCC 4-X)

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### Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: WT SV40 MEF (ATCC CRL-2907)

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### References

References and other information relating to this material are available at [www.atcc.org](http://www.atcc.org).

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