

**Product Name**

**Name:** mSRCgel™ Extracellular Matrix, High Protein Concentration, Reduced Growth Factor extracted from EHS mouse sarcoma  
[+] Phenol Red

**Cat. No.:** C2032-0005, C2032-0010

**Size:** 5 mL, 10 mL

**Product Description**

The extracellular matrix (ECM) can provide structural support for cells and tissues in a dynamic three-dimensional network of macromolecules. The ECM is a molecular network which holds bioactive molecules and growth factors together. It is of vital importance that it controls the basal behaviors and characteristics of cells such as adhesion, migration, polarity, differentiation, proliferation, and apoptosis.

**mSRCgel** is a natural basement membrane (BM) extracted from **mouse sarcoma** cells, a type of connective tissue tumor. mSRCgel is high in ECM proteins, including laminin, collagen IV, heparan sulfate proteoglycan (perlecan), entactin, and many essential growth factors.

**Application**

In vivo angiogenesis studies and 3D tumor models.  
Protein concentration: 16 - 26 mg/mL

**Storage and Stability**

The product should be kept at **-20°C**. Avoid multiple freeze-thaws.  
The product is **light-sensitive** and therefore should not be left in the light.  
Shelf life: Stable until expiry date on the label

**Procedure****Example for Procedures:****Subcutaneous tumor formation in mice****1. Equipment, reagents and consumables**

- Equipmen: Biosafety cabinet, cell incubator, low temperature horizontal centrifuge, and inverted microscope.
- Reagents: mSRCgel, basal medium, a medium containing 10% fetal bovine serum, 1 × PBS, trypsin solution.
- Consumables: sterile pipette tips, 10 cm cell culture dish, sterile microcentrifuge tubes, the amounts of disposable syringes and other consumables may be adjusted according to the experimental design.

**2. Example protocol**

- a) Put mSRCgel in an ice box and put it (in the refrigerator) at 4°C so that the gel can slowly melt overnight;
- b) Select target cells grown in good condition and logarithmically, discard the supernatant, add 2 mL of 1×PBS solution to wash gently, and discard the liquid;
- c) Add 1 mL of 0.25% trypsin solution into the petri dish, let it stand for 10 seconds, discard the

trypsin, and continue the digestion at room temperature for 1~3 minutes with the residual trypsin.

- d) When the cells become round (keep in mind that they cannot be digested until the cell edge is clear), add 1 mL of the medium containing 10% fetal bovine serum to terminate the digestion. Carefully pipette the cells and collect the cell suspension into a 15 mL plastic centrifuge tube.
- e) Centrifuge at 1,000 rpm for 3 minutes and discard the supernatant. After washing the cells with PBS once, re-suspend the cells by with serum-free basal medium, and take 10  $\mu$ L of the cell suspension for cell count.
- f) HepG2 cells: Each mouse should be inoculated with 5 million cells. The required volume of the cell suspension is centrifuged in a sterile 1.5 mL plastic centrifuge tube at 1,000 rpm for 3 min. The supernatant is discarded and serum-free medium or PBS is added until the total volume is 300  $\mu$ L.

HCT - 116 cells: Each mouse should be inoculated with 1 million cells. The required volume of the cell suspension is centrifuged in a sterile 1.5 mL plastic centrifuge tube at 1,000 rpm for 3 min. The supernatant is discarded and serum-free medium or PBS is added until the total volume is 300  $\mu$ L.

MIA - PaC - 2 cells: Each mouse should be inoculated with 10 million cells. The required volume of the cell suspension is centrifuged in a sterile 1.5 mL plastic centrifuge tube at 1,000 rpm for 3 min. The supernatant is discarded and serum-free medium or PBS is added until the total volume is 500  $\mu$ L.

- g) Mixing with mSRCgel: The cell suspension and mSRCgel are mixed in a 1:1 ratio at 4°C.
- h) Subcutaneous injection: The left hand of a nude mouse is fixed and the cell suspension is injected subcutaneously into the right back of the nude mice. During the inoculation, the needle is inserted a little deeper, about 1 cm deep, to reduce the overflow of cell suspension from the site of injection, and the inoculation volume is 100 $\mu$ L. For MIA-PaC-2 cells, the volume injected is 200  $\mu$ L/piece.
- i) Recording data: The nude mice are put back into the cage for further feeding, and the tumor volume is measured regularly according to the experimental design. The data are recorded to draw a curve and the nude mice are euthanized after the tumor volumes grow to just less than 2,000 mm<sup>3</sup>. The tumors is then removed and photographed.

## Quality Control

mSRCgel Extracellular Matrix is tested for the presence of bacteria, fungi, and mycoplasma. In addition, osmolality, protein concentration, endotoxin, gel stability, and biological activity are tested.

## Precaution and Disclaimer

For research use only, not for clinical diagnosis, and treatment.