

Scapova™



**PVA Microcarriers
for cell mass culture**

kuraray

Development Concept of SCAPOVA™

SCAPOVA™ series are microcarriers produced by Kuraray. They are made from PVA (Polyvinyl alcohol). SCAPOVA™ can solve following issues in the manufacturing cells for regenerative medicine.

**Microcarriers
for regenerative
medicine**

1

**Obtain enough
number of cells**

2

**Assure medical-
grade safety**

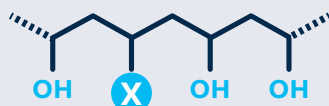
Manufacturing Method of SCAPOVA™

1



Polyvinyl Alcohol (PVA)

It is resin with high elasticity and solubility. It also have high biocompatibility.



Partially modify the hydroxyl groups

2

3



Ink

Solution of new modified PVA.

4

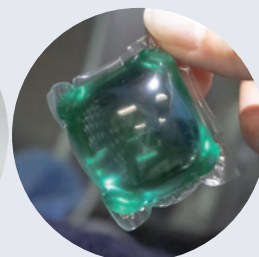


PVA Microcarriers(non-coated)

Micron-sized hydrogel beads are made by cross-linking PVA.

Kuraray's other PVA products

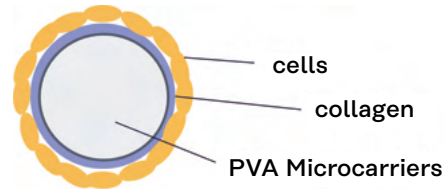
- Polarizing film in LCD
- Water-soluble films for detergent



SCAPOVA™ CL

SCAPOVA™ CL

Clinically usable collagen is coated on the particle surface. Endotoxin and virus in the collagen have been reduced or inactivated.



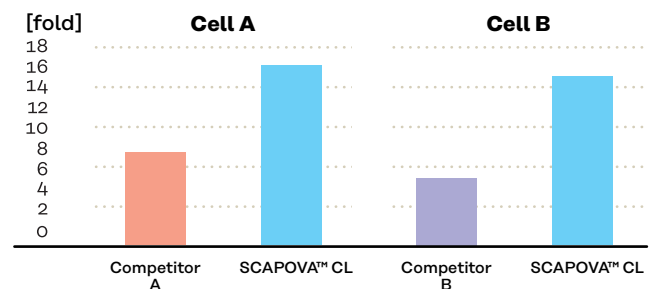
Characteristics of SCAPOVA™ CL

1

High cell proliferation rate

Therapeutic cells, like MSCs, can easily expand on SCAPOVA™ CL.

Human cells cultured on each microcarriers for 7 days. Each bar shows the fold of the initial cell number.



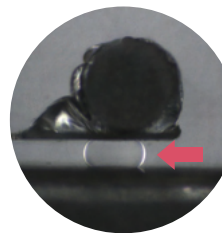
2

Quality management in accordance with medical standards

SCAPOVA™ CL is made from Kuraray's PVA*. We conduct strict quality management. And there is extremely low risk of contamination.

*Polyvinyl Alcohol

Safety test	Result
Cytotoxicity	Negative (Tested in accordance with ISO 10993.)
Genotoxicity	
Systemic acute toxicity	
Implantation test	
Leachable	Tested in accordance with BPOG guideline.
Extractable	

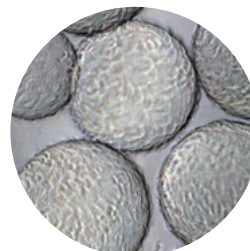


This image shows compression test of SCAPOVA™

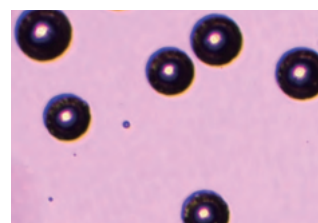
3

Ease of handling

SCAPOVA™ CL is "Ready to use." No need for sterilization and cleaning before using. It swells quickly in PBS or culture medium. Swollen particles are highly transparent. Therefore, cells can be easily observed with a phase contrast microscope.



High transparency makes it easy to observe cells by microscopy



Immediately after SCAPOVA™ CL is added to the culture medium



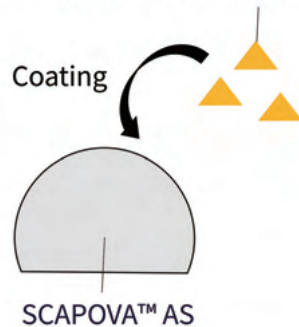
10 seconds after adding culture medium

SCAPOVA™ AS

SCAPOVA™ AS

The particle surface is specially processed and activated.
By activation, any cell adhesion molecules can be coated.

Cell adhesion molecules



Characteristic of SCAPOVA™ AS

1

Suitable for various cell culture

A wide variety of cells can be cultured by coating optimal cell adhesion molecules for target cells.

Coating experience

Vitronectin / Fibronectin / Laminin / iMatrix-511 / Synthemax® II / Poly-L-lysine / Collagen

2

Animal Origin Free

SCAPOVA™ AS does not contain any animal derived materials.

Using recombinant proteins or synthetic peptides, animal free cell culture can be achieved.

Protocol of coating

SCAPOVA™ AS needs coating of cell adhesion molecules before using.
There are two coating methods.

Method

A

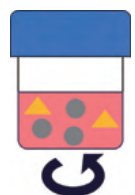
1. Prepare the coating material solutions, such as cell adhesive proteins.
2. Add coating material solutions directly to SCAPOVA™ AS in dry condition.
3. Gently mix SCAPOVA™ AS dispersion and then wash with PBS.



Method

B

1. Add culture medium and coating material solution into bioreactor.
2. Add SCAPOVA™ AS into the medium with coating material and mix them.



After coating, the culture procedure is the same as usual. Please contact us for detail protocol.

Product information of SCAPOVA™

SCAPOVA™ CL

SCAPOVA™ AS

Outline date	Particle size(D50)	200 - 250 µm	120 - 200 µm
	Surface	Collagen coated	Activated treatment
	Swelling factor (in PBS)	10	5
	The surface area [/ <i>g</i> dry weight]	2,600 cm ²	1,900 cm ²
	Recommended amount [/ <i>ℓ</i> for MSC]	1.54 g	2.11 g
	Sterilization	Gamma irradiation	Gamma irradiation

※SCAPOVA™ can be available only for research. Please contact us for clinical use.

Quantity & Price	Product name	Quantity	Price	Product code
	SCAPOVA™ CL	1 g	Contact us	M11018SAC1-01GB
		5 g		M11018SAC1-05GB
		10 g		M11018SAC1-10GB
	SCAPOVA™ AS	1 g		M11018PF00-01GB
		5 g		M11018PF00-05GB
		10 g		M11018PF00-10GB

Culturing experience

SCAPOVA™ CL

- hMSC
- human fibroblasts
- VERO cells
- Bovine myoblasts

SCAPOVA™ AS

- hMSC
- hiPS cells



Please contact us for the protocol and application notes.

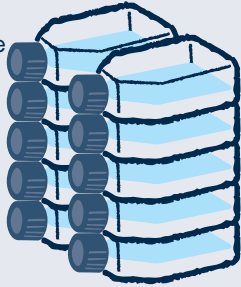
What is a microcarrier?

**Microcarrier is micron-sized bead for cell culture scaffold.
Cells attached to the bead surface and proliferate on the bead.**

Cell culture using flasks

- Many flasks take a lot of space.
- It takes a lot of labor
- It doesn't suit for mass culture

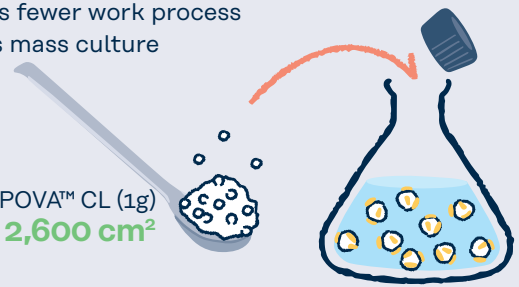
e.g. T175 flasks (10 flasks)
about 1,750 cm²



Cell culture using microcarriers

- It saves space
- It takes fewer work processes
- It suits mass culture

e.g. SCAPOVA™ CL (1g)
about 2,600 cm²



1

Input

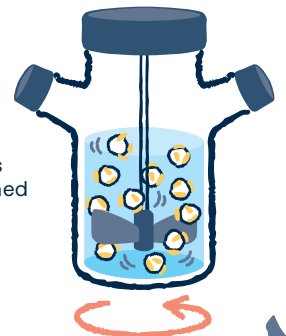
Weigh SCAPOVA™ and put them into culture medium. Any containers, such as flasks and bioreactors, can be used.



2

Cell seeding

Seed cells and leave it for 24 hours (agitate at determined speed). After that, agitate at suitable speed.



3

Medium exchange

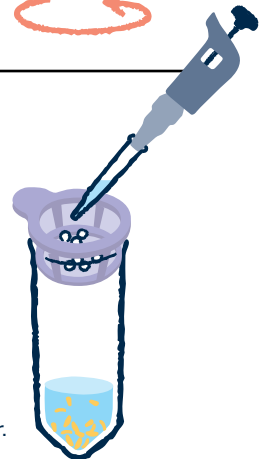
After stopping agitation, SCAPOVA™ settle down within few minutes. Change half of the medium every 3 or 4 days.



4

Cell harvesting

Detach cells from SCAPOVA™ with Trypsin/EDTA solution. Separate cells and SCAPOVA™ by cell strainer.



How to use SCAPOVA™

We have application notes with details on how to use the microcarriers.

Contact us

KURARAY CO., LTD. Life Innovation Promotion Business Department, Research and Development Division

✉ Contact.LIPG@kuraray.com

🌐 <https://www.kuraray.co.jp/microcarriers/>



kuraray