

CYTOSTORTM

PRESERVATION OF CELLS & VIRUSES IN SUSPENSION

CS-SNS; CS-LNS



Atelerix Handbook Series Version CS01;1.1.1

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1. COMPONENTS

1.1. KIT CONTENTS

PRODUCT CODE	COMPONENTS	UNITS	UNIT VOLUME	CELL SUSPENSION TO ADD
	CytoStor Vial (Beads)	3 Tubes	0.4ml	-
CS-SNS-03	Gel A (5x)	3 Tubes	0.12ml	0.48ml
	Dissolution Buffer	3 Tubes	1.1ml	-
	CytoStor Vial (Beads)	6 Tubes	0.4ml	-
CS-SNS-06	Gel A (5x)	6 Tubes	0.12ml	0.48ml
	Dissolution Buffer	6 Tubes	1.1ml	-
	CytoStor Vial (Beads)	12 Tubes	0.4ml	-
CS-SNS-12	Gel A (5x)	12 Tubes	0.12ml	0.48ml
	Dissolution Buffer	12 Tubes	1.1ml	-
	CytoStor Vial (Beads)	24 Tubes	0.4ml	-
CS-SNS-24	Gel A (5x)	24 Tubes	0.12ml	0.48ml
	Dissolution Buffer	24 Tubes	1.1ml	-
	CytoStor Vial (Beads)	50 Tubes	0.4ml	-
CS-SNS-50	Gel A (5x)	50 Tubes	0.12ml	0.48ml
	Dissolution Buffer	50 Tubes	1.1ml	-
	CytoStor Vial (Beads)	3 Tubes	2ml	-
CS-LNS-03	Gel A (5x)	3 Tubes	0.6ml	2.4ml
	Dissolution Buffer	3 Tubes	5ml	-
	CytoStor Vial (Beads)	6 Tubes	2ml	-
CS-LNS-06	Gel A (5x)	6 Tubes	0.6ml	2.4ml
	Dissolution Buffer	6 Tubes	5ml	-
	CytoStor Vial (Beads)	12 Tubes	2ml	-
CS-LNS-12	Gel A (5x)	12 Tubes	0.6ml	2.4ml
	Dissolution Buffer	12 Tubes	5ml	-
	CytoStor Vial (Beads)	24 Tubes	2ml	-
CS-LNS-24	Gel A (5x)	24 Tubes	0.6ml	2.4ml
	Dissolution Buffer	24 Tubes	5ml	-
	CytoStor Vial (Beads)	50 Tubes	2ml	-
CS-LNS-50	Gel A (5x)	50 Tubes	0.6ml	2.4ml
	Dissolution Buffer	50 Tubes	5ml	-

Note: Remove components from 2-8°C for at least 20 minutes before use.

1.2. COMPONENTS TO BE SUPPLIED BY THE USER

• 1000µl Pipettes and Tips • Cell Culture Medium • Cells or viruses!!

1.3. BEFORE YOU BEGIN USING CYTOSTOR™

- Ensure CytoStor™ kits have not passed the expiry date stated on the packaging.
 Atelerix does not recommend using kits after this date.
- 2. Read the troubleshooting guide on page 12 to see our list of frequently asked questions. For any further queries, please email us at Sales@atelerix.co.uk.
- 3. Consult the Cell Density and Loading guide (page 4 or 7 for cells, page 10 for viruses) and the Cell and Virus Storage Temperature Guide on page 13.
- 4. CytoStor™ is intended for use solely in accordance with this protocol using the components provided within the kit.

2. PROTOCOL OVERVIEW





3. CYTOSTOR™ (CS-SNS) FOR THE ENCAPSULATION OF CELLS

3.1. GELATION

- 1. Ensure that all components are allowed to equilibrate to room temperature before use and that gels are at the bottom of their tubes. Conduct all steps in a laminar flow hood at room temperature.
- 2. Resuspend cells in the appropriate volume of culture medium (See 3.2 Cell Density and Loading Guide on page 4 for more information), ensuring thorough distribution of cells.
- 3. Add 0.48 mL of the cell culture suspension to the vial containing 0.12 mL of Gel A.
- 4. Gently mix until homogenous, with a pipette, ensuring that no bubbles are introduced (see troubleshooting guide on page 12).
- 5. Add 0.6 mL of the cells / Gel A mix to the CytoStor Vial¹.
- 6. Place the cap back on the tube and gently invert the gel / bead mixture several times until the beads are evenly distributed throughout the gel. Gently flick the tube to settle the contents, ensuring a tight seal (the gel will cure in situ within approximately 30 minutes, sample is ready to ship after 1 hour).
- 7. Store away from light in a polystyrene box at the recommended temperature for the cell type encapsulated. See the table on page 13 or, for the most up to date recommendations on storage temperatures and times, visit our Compatibility section on our website or contact sales@atelerix.co.uk.

For directions on shipping your samples, see Section 3.4.

*Use the CytoStor Vial containing beads provided for encapsulation, storage, and release.





3.2. CELL DENSITY AND LOADING GUIDE

3.2.1. CELL LOADING GUIDE

Desired Cell Load Per Sample (x10 ⁶)	Cell Suspension Concentration (x10 ⁶ cells/mL)	Volume of Cell Suspension per Sample (mL)
1	2.08	0.48
2	4.17	0.48
3	6.25	0.48
4	8.33	0.48
5	10.42	0.48
6	12.50	0.48
7	14.58	0.48
8	16.67	0.48
9	18.75	0.48
10	20.83	0.48

For multiple encapsulations, multiply the volume of cell suspension (C) by the number of samples required.

3.2.2. RECOMMENDED CELL LOAD FOR DIFFERENT CELL TYPES

Cell Diameter	Example Cell Types	Recommended Cell Load per Encapsulation
4 – 10 Microns	Lymphocytes e.g T-Cells, PBMC's	2 x 10 ⁷ Cells
11 – 15 Microns	Fibroblasts CHO Cells HEK Cells	≤ 1 x 10 ⁷ Cells
16 – 30 Microns	Mesenchymal Stem Cells Monocytes Hepatocytes	≤ 1 x 10 ⁷ Cells

CytoStor™ has no lower limit to the number of cells which can be stored, however we recommend using enough cells to be easily recovered after centrifugation.

3.3. RELEASE

- Ensure that all components and samples are allowed to equilibrate to room temperature before use and conduct all steps in a laminar flow hood at room temperature.
- 2. Using a pipette tip or syringe and needle, infuse 1 mL Dissolution Buffer into the bottom of the gel by piercing the gel.
- 3. Place the cap back on the tube and allow the gel to dissolve by occasionally agitating the tube by gentle inversion or rocking for 10 minutes (see troubleshooting guide on page 12).
- 4. Sediment cells by centrifugation at your cells' usual centrifugation settings (e.g. 350RCF for 5 minutes), remove supernatant, and re-suspend cells in medium of choice.

3.4. SHIPPING YOUR CELLS

Use appropriate controlled temperature packaging² when preparing cells for shipping to reduce the effect of ambient temperature change on the encapsulated cells during transit.

²For best cell recovery upon arrival, we recommend using the ICECATCH Solid Ambient or Cool Shipping Boxes.

Find out more & shop at https://www.atelerix.co.uk/pages/variants-collection-page-accessories





4. CYTOSTOR™ LARGE (CS-LNS) FOR THE ENCAPSULATION OF CELLS

4.1. GELATION

- 1. Ensure that all components are allowed to equilibrate to room temperature before use and that gels are at the bottom of their tubes. Conduct all steps in a laminar flow hood at room temperature.
- 2. Resuspend cells in the appropriate volume of culture medium (See 4.2 Cell Density and Loading Guide on page 7 for more information), ensuring thorough distribution of cells.
- 3. Add 2.4 mL of the cell culture suspension to the vial containing 0.6 mL of Gel A.
- 4. Gently mix until homogenous, with a pipette, ensuring that no bubbles are introduced (see troubleshooting guide on page 12).
- 5. Add 2.4 mL of the cells / Gel A mix to the CytoStor Vial¹.
- 6. Place the cap back on the tube and gently invert the gel / bead mixture several times until the beads are evenly distributed throughout the gel. Gently flick the tube to settle the contents, ensuring a tight seal (the gel will cure in situ within approximately 30 minutes, sample is ready to ship after 1 hour).
- 8. Store away from light in a polystyrene box at the recommended temperature for the cell type encapsulated. See the table on page 13 or, for the most up to date recommendations on storage temperatures and times, visit our Compatibility section on our website or contact sales@atelerix.co.uk.

For directions on shipping your samples, see Section 4.4.

*Use the CytoStor Vial containing beads provided for encapsulation, storage, and release.

4.2. CELL DENSITY AND LOADING GUIDE

4.2.1. CELL LOADING GUIDE

Desired Cell Load Per Sample (x10°)	Cell Suspension Concentration (x10° cells/mL)	Volume of Cell Suspension per Sample (mL)
5	2.08	2.40
10	4.17	2.40
15	6.25	2.40
20	8.33	2.40
25	10.42	2.40
30	12.50	2.40
35	14.58	2.40
40	16.67	2.40
45	18.75	2.40
50	20.83	2.40

For multiple encapsulations, multiply the volume of cell suspension (C) by the number of samples required.

4.2.2. RECOMMENDED CELL LOAD FOR DIFFERENT CELL TYPES

Cell Diameter	Example Cell Types	Recommended Cell Load per Encapsulation
4 – 10 Microns	Lymphocytes e.g T-Cells, PBMC's	1 x 10 ⁸ Cells
11 – 15 Microns	Fibroblasts CHO Cells HEK Cells	≤ 5 x 10 ⁷ Cells
16 – 30 Microns	Mesenchymal Stem Cells Monocytes Hepatocytes	≤ 5 x 10 ⁷ Cells

CytoStor™ has no lower limit to the number of cells which can be stored, however we recommend using enough cells to be easily recovered after centrifugation.







4.3. RELEASE

- 1. Ensure that all components and samples are allowed to equilibrate to room temperature before use and conduct all steps in a laminar flow hood at room temperature.
- 2. Using a pipette or syringe and needle, infuse 5 mL Dissolution Buffer into the bottom of the gel by piercing the gel.
- 3. Place the cap back on the tube and allow the gel to dissolve by occasionally agitating the tube by gentle inversion or rocking for 10 minutes (see troubleshooting guide on page 12).
- 5. Sediment cells by centrifugation at your cells' usual centrifugation settings (e.g. 350RCF for 5 minutes), remove supernatant, and re-suspend cells in medium of choice.

4.4. SHIPPING YOUR CELLS

Use appropriate controlled temperature packaging² when preparing cells for shipping to reduce the effect of ambient temperature change on the encapsulated cells during transit.

²For best cell recovery upon arrival, we recommend using the ICECATCH Solid Ambient or Cool Shipping Boxes.

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5. CYTOSTOR™ (CS-SNS) FOR ENCAPSULATION OF VIRUSES

5.1. GELATION

- Ensure that all components are allowed to equilibrate to room temperature before
 use and that gels are at the bottom of their tubes. Conduct all steps in a laminar
 flow hood at room temperature.
- 2. Resuspend virus in the appropriate volume of medium (See 5.2 Viral Titre and Loading Guide on page 10 for more information), ensuring thorough distribution of virus.
- 3. Add 0.48 mL of the virus suspension to the vial containing 0.12 mL of Gel A.
- 4. Gently mix until homogenous, with a pipette, ensuring that no bubbles are introduced (see troubleshooting guide on page 12).
- 5. Add 0.6 mL of the virus / Gel A mix to the CytoStor Vial¹.
- 6. Place the cap back on the tube and gently invert the **gel / bead** mixture several times until the beads are evenly distributed throughout the gel. Gently flick the tube to settle the contents, ensuring a tight seal (the gel will cure in situ within approximately 30 minutes, sample is ready to ship after 1 hour).
- 7. Store away from light in a polystyrene box at room temperature (15°C 25°C).

For directions on shipping your samples, see Section 5.4.

*Use the CytoStor Vial containing beads provided for encapsulation, storage, and release.





5.2. VIRAL TITRE AND LOADING GUIDE

5.2.1. LOADING GUIDE

Desired Viral Load per Sample	Virus Titre	Volume of Virus Suspension per Sample
(x 10 ⁸)	(x 10 ⁸ IFU/mL)	(mL)
1	2.08	0.48
2	4.17	0.48
3	6.25	0.48
4	8.33	0.48
5	10.42	0.48
6	12.50	0.48
7	14.58	0.48
8	16.67	0.48
9	18.75	0.48
10	20.83	0.48

For multiple encapsulations, multiply the volume of viral suspension (C) by the number of samples required.

5.2.2. RECOMMENDED VIRAL LOAD

Sample	Example Types	Recommended Viral Load per Encapsulation
Virus	Lentivirus,	
	Coronavirus, Viral	x 10 ⁹ IFU
	Vectors	

CytoStor TM has no lower limit to the number of virus which can be stored, however we recommend using enough cells to be easily recovered after centrifugation.

5.3. RELEASE

- 1. Ensure that all components and samples are allowed to equilibrate to room temperature before use and conduct all steps in a laminar flow hood at room temperature.
- 2. Using a pipette tip or syringe and needle, infuse 1 mL Dissolution Buffer into the bottom of the gel by piercing the gel.
- 3. Place the cap back on the tube and allow the gel to dissolve by occasionally agitating the tube by gentle inversion or rocking for 10 minutes (see troubleshooting guide on page 12).
- 4. Virus can then either be:
 - a. Sedimented by ultracentrifugation. Following ultracentrifugation, remove supernatant and re-suspended virus in medium of choice.
 - b. Diluted in a medium of choice (minimum 1 in 4 dilution recommended).

5.4. SHIPPING YOUR VIRUS

Use appropriate controlled room temperature packaging² when preparing virus for shipping to reduce the effect of ambient temperature change on the encapsulated virus during transit.

²For best cell recovery upon arrival, we recommend using the ICECATCH Solid Ambient or Cool Shipping Boxes.

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6. TROUBLESHOOTING GUIDE

Problem / Question	Guidance	
I have air bubbles in the gel	Air trapped within the gel will affect preservation, so	
after mixing with my media, is	bubbles should be eliminated before mixing with the	
this a problem?	beads. Allow time for the mixture to settle and the	
	bubbles to travel to the surface before addition.	
Can I use the kit to	We would recommend TissueReady™ for the	
encapsulate organoids and	encapsulation and storage of organoids and	
/ or spheroids?	spheroids.	
Can I ship the Dissolution	Yes, the Dissolution Buffer is stable at a wide range of	
Buffer in the same package	temperatures and can be shipped together with the	
as the samples?	encapsulated samples.	
What are the recommended	A guide to the recommended storage times and	
storage times and	temperatures can be found on our <u>Compatibility</u>	
temperatures for my cell	section on our website. If you cannot find any	
type?	recommendations for your cell type, please contact	
	Sales@atelerix.co.uk.	
Can I reuse the contents of	No, there should only be sufficient volume for a set	
the kit if I don't use it all?	number of encapsulations per kit. Any spare reagents	
	will not be sufficient to perform any additional	
	encapsulations properly.	
Can I use PBS instead of	No, PBS should not be used at any point as it inhibits	
media when encapsulating	and slowly reverses gelation.	
samples?		
Can I split the kit into	No, we do not recommend removing the GelBase	
smaller tubes to get more	Beads from the tubes supplied or deviating from the	
encapsulations?	volumes stated.	
What if the beads have not	This sometimes occurs when the medium used to	
fully dissolved after 10	store the beads fortifies the gel. Extend the incubation	
minutes? Can the cells be	until the beads dissolve - this should not take much	
allowed to sit in the buffer	longer than 15 minutes. Please contact	
for longer?	Sales@atelerix.co.uk.for information on how long	
	cells may be allowed to incubate in the Dissolution	
	Buffer.	

7. STATEMENTS

7.1. KIT STORAGE AND STABILITY

This kit is stable at 2-8°C for up to 12 months. Atelerix does not recommend using the kit after the expiry date stated on the packaging.

7.2. CELLULAR MATERIAL

Please ensure that cell cultures are free of fungal and bacterial contamination before proceeding.

7.3. TRADEMARKS

Cyto $Stor^{TM}$ is a trademark of Atelerix Ltd.

8. APPENDIX: CELL AND VIRUS STORAGE TEMPERATURE GUIDE

Cell Type	Recommended Storage Conditions	Tested Storage Time
PBMCs	2-8°C	3-Days
Jurkat Cells	15-25°C	14-Days
MSCs	15-20°C	21-Days
Monocytes	2-8°C	5-Days
Fibroblasts	15-20°C	14-Days
Viruses	15-25°C	14-Days

If you cannot find any recommendations for your cell type, please visit our <u>Compatibility</u> section on our website or contact <u>Sales@atelerix.co.uk</u>.



NOTES

