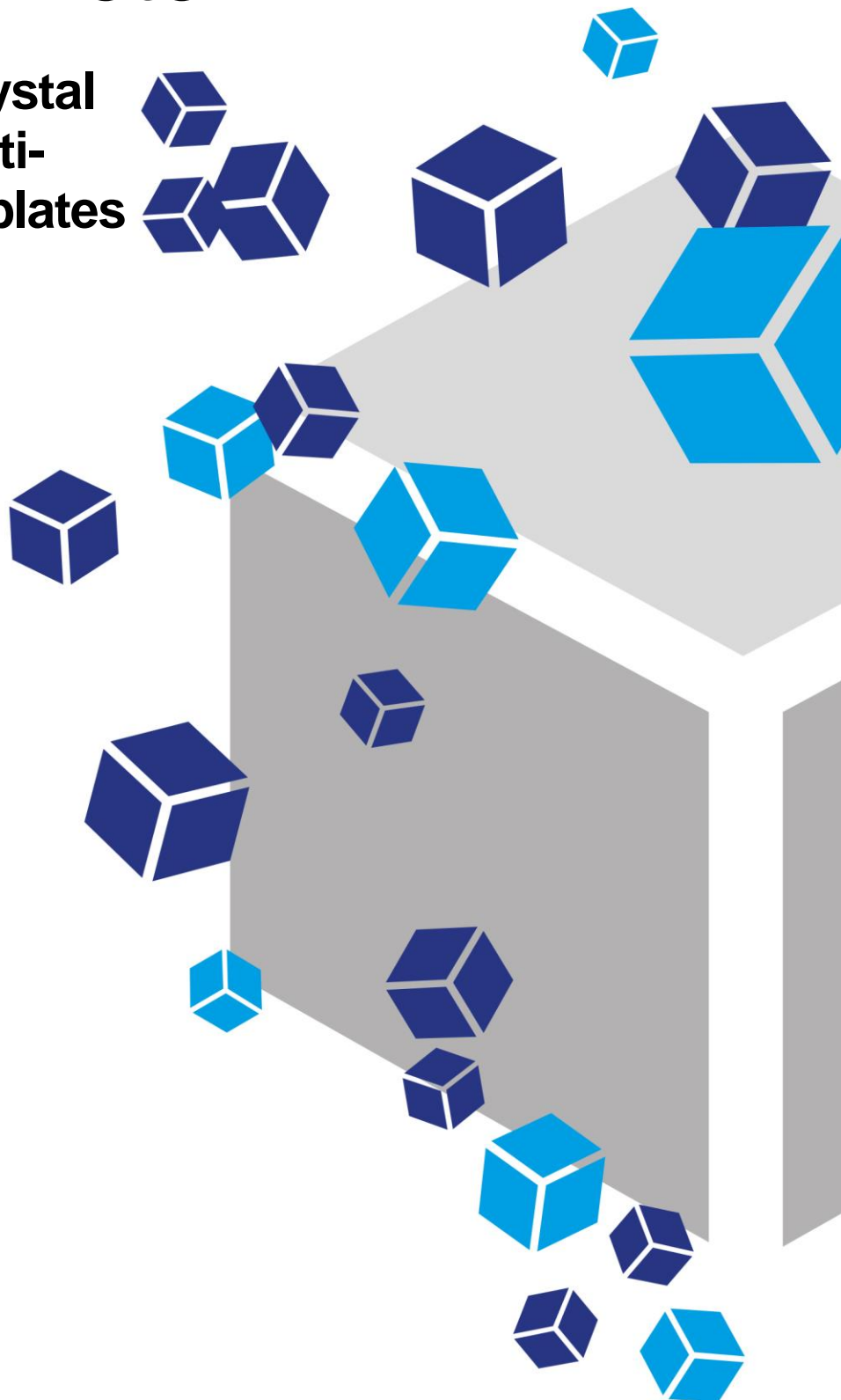


# Application Note

**Creating PODS<sup>®</sup> crystal monolayers on multi-well tissue culture plates**



# Creating PODS<sup>®</sup> crystal monolayers on multi-well tissue culture plates

## Introduction to PODS<sup>®</sup>

### The challenge with soluble growth factors

Many proteins, especially growth factors and cytokines, when used as a reagent, degrade quickly, rapidly losing their bioactivity. This fragility hampers research and significantly limits the therapeutic potential of proteins.

### Protein Micro-depots

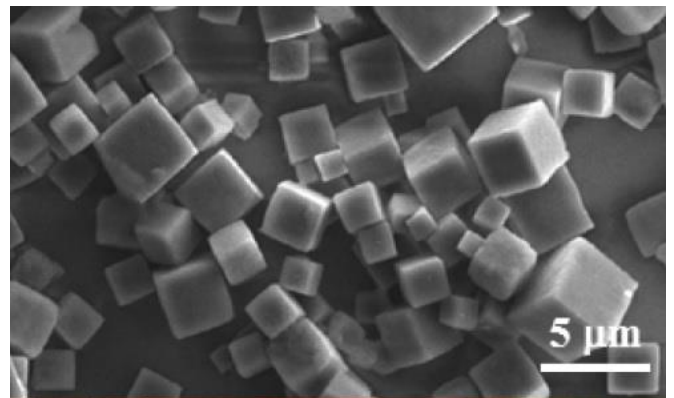
Development of a technology that can continuously replenish active protein from a local, microscopic store, has been a significant challenge, but one that could transform the fields of cell culture and medicine by allowing greater control over the growth of cells.

### Introducing PODS<sup>®</sup>

PODS<sup>®</sup> technology has made the goal of a micro-depot for proteins a reality. PODS<sup>®</sup> is a sustained release system which continuously replenishes proteins from millions of local microscopic stores which can be placed next to (or at a distance from) cells, either randomly or in precise locations. Just like cells, these micro-depots release a steady stream of bioactive protein. This protein can be limited to local surroundings or dispersed more widely, or made to form a gradient.

### How does it work?

At the heart of PODS<sup>®</sup> is an extraordinary polyhedrin protein. This specific polyhedrin protein has the unique ability to encase cargo proteins within perfect, transparent, cubic, micro-sized crystals, much smaller than the cells. These protein crystals form admixtures of the polyhedrin and cargo proteins which slowly degrade releasing the biologically active cargo protein.



### How can PODS<sup>®</sup> help my research?

PODS<sup>®</sup> are tough and will withstand physical and chemical stress, so you can handle them with ease. PODS<sup>®</sup> can be made to release intact cargo protein over days, weeks or even months. Using PODS<sup>®</sup> you can readily create a steady-state protein environment in microscopic detail wherever you want, tailored exactly to your requirements. This is the power of PODS<sup>®</sup>. PODS<sup>®</sup> proteins are now available for many growth factors and cytokines and are already being used in many leading world-class research labs. PODS<sup>®</sup> protein applications include:

- Micropatterning
- Physiological, stable gradient formation
- Bioinks for 3D printing
- Microcarriers
- Functionalizing scaffolds
- Microfluidics (lab on a chip)
- Improved and simplified stem cell culture
- Therapeutic protein delivery

## Methods

Adhering PODS<sup>®</sup> crystals to tissue culture (TC) multi-well plates or inserts

1. Dilute PODS<sup>®</sup> crystals into PBS and pipette the suspension into wells of a TC multi-well plate or TC inserts (see table below for recommended PODS<sup>®</sup> crystal amounts and PBS volumes).
2. Spin TC multi-well plate or inserts in wells in a centrifuge with plate rotor for 20 minutes at 3000 x g.
3. Remove supernatant and allow PODS<sup>®</sup> crystals to dry onto wells/inserts, either by leaving the plate at room temperature for 1 h with the lid removed or overnight in the fridge with the lid closed.
4. Add cell suspension or media to wells by gently pipetting against the side of the well.

## Results

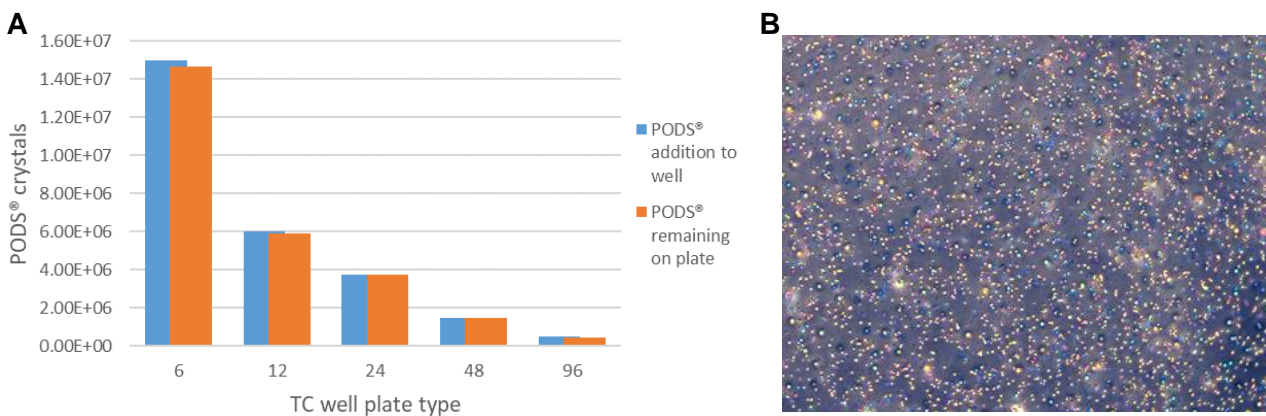
**Recommended maximum PODS<sup>®</sup> crystal amounts and PBS volumes to create PODS<sup>®</sup> monolayers by centrifugation**

Plate type	Area (cm <sup>2</sup> )	Recommended PBS [ $\mu$ l]	Maximum recommended PODS <sup>®</sup> *	Loss of PODS <sup>®</sup> after supernatant removal (%)	Loss of PODS <sup>®</sup> after a full media change (%)
6	9.50	1000	$1.5 \times 10^7$	2.0	4.0
12	3.80	600	$6.0 \times 10^6$	1.5	3.0
24	1.90	300	$3.0 \times 10^6$	0.5	1.5
48	0.95	150	$1.5 \times 10^6$	1.0	9.0**
96	0.32	50	$5.0 \times 10^5$	3.0	5.0

\* Higher densities of PODS<sup>®</sup> crystals can be achieved but will result in higher subsequent losses.

\*\* Higher loss of PODS<sup>®</sup> crystals is due to the geometry of a 48-well plate that prevents pipetting to the side of the well.

### PODS<sup>®</sup> crystals adhered to TC plastic surfaces



Maximum recommended amounts of PODS<sup>®</sup> crystals were spun onto various TC plates.

**(A)** Retention of PODS<sup>®</sup> crystals on TC plastic surfaces were measured after aspiration of supernatant.

**(B)** Brightfield micrograph of a monolayer of dispersed PODS<sup>®</sup> crystals at the maximum recommended density.

## Conclusions

- PODS<sup>®</sup> crystals adhered to TC surfaces as a monolayer will release growth factor evenly across the well.
- Uniformly distributed monolayers of PODS<sup>®</sup> crystals on TC plastic can be easily achieved using a centrifuge with plate rotor.
- Crystal densities of up to  $1.4 \times 10^6$  PODS<sup>®</sup>/cm<sup>2</sup> will readily adhere to TC surfaces as a well-spaced out single layer.
- Medium changes can be performed by simply aspirating conditioned medium as usual without significant loss of PODS<sup>®</sup> crystals.

For more information and a full list of our current PODS<sup>®</sup> growth factors, please visit our website [www.cellgs.com](http://www.cellgs.com).



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