

Do I need to change the type of media I use if I am using FGF2 Discs?

No, FGF2 DISCs can be used with the media you normally use for your cell cultures. The only change is that with DISCs you will feed less often and use less medium. Using DISCs and performing fewer feeds improves stem cell pluripotency.

Will FGF2 DISCs interfere with my downstream application of the cells?

FGF2 DISCs have been shown to have a positive impact on downstream applications of cells because they increase pluripotency and reduce unwanted cell differentiation.

Are FGF2 DISCs difficult to add or remove from my cultures?

No, FGF2 DISCs can be easily added using sterile forceps and removed by using a low vacuum or pipette.

What if I want a different concentration, can the protein release be adjusted?

Yes, the protein concentration can be adjusted based on the number of DISCs and amount of medium. See the table on the right for examples.

Number of FGF2-DISCs	Volume of Medium	FGF2 Release Level	Example Culture Dish
 1	0.5 ml	40 ng/ml	24 well plate
 1	1 ml	20 ng/ml	12 well plate
 1	2 ml	10 ng/ml	6 well Plate
 1	4 ml	5 ng/ml	T25 Flask
 2	2 ml	20 ng/ml	6 well plate
 3	2 ml	30 ng/ml	6 well plate
 4	2 ml	40 ng/ml	6 well plate

What if I observe a lower proliferation of pluripotent stem cells (PSCs) after a passage?

We suggest increasing your seeding density (i.e. decrease the split ratio).

What should I do if I see pH changes in my cultures?

Cultures should be monitored, and additional medium changes performed as needed. In high density cultures, this is especially important: we recommend increasing the frequency of medium changes or adding buffers to your medium, such as sodium bicarbonate or HEPES, to help maintain pH levels.

What should I do if I see increased cell debris in my cultures?

When medium is changed less often, it should be expected that cell debris will accumulate and may appear at a higher level than when medium is changed daily. Users have not reported this as being detrimental.

How often do I need to change the DISC?

We recommend changing the DISC every 6-8 days.

Do I have to change a DISC every time I do a medium change?

No, leave the same DISC in the well for about a week and perform medium changes without changing the DISC. For tips and an instructional video on using and removing FGF2 DISCs, click this link:

<https://stemcultures.com/resources/how-to-use-discs/>

Can DISCs work with other cell types?

Yes, FGF2 signaling is important for several different cell types.

Can DISCs release other growth factors besides FGF2?

Yes, DISCs are a platform technology and can be custom made. Please contact us at support@stemcultures.com for custom orders.

What do I do if DISCs become hydrated and difficult to handle?

DISCs are hydrogels and readily absorb moisture, which can cause them to stick and become more difficult to handle. We recommend storing DISCs at 4 °C. If DISCs do absorb moisture, they can easily be re-dried in a cell culture hood by leaving the lid ajar for 15-30 minutes.