

## Human fibroblasts, dermis (HFIB-D), cryo

Cat.-No.: 121 0411 (500,000 cells / cryovial)

### Maintenance of HFIB-D

Check the cryovial for signs of damage during dispatch. Since the cryopreserved HFIB-D arrived on dry ice, they have to be transferred immediately to a deep freezer (-80°C) for a storage time of up to one week or to be thawed according to the procedure stated below. For

In the case of longer storage, transfer the vial directly to liquid nitrogen (-196°C).

### Thawing of cryopreserved HFIB-D:

1. Transfer the cryovial straight to a water bath with a temperature of 37°C. To achieve a rapid thawing, gently stir the cryovial repeatedly. Few seconds before thawing is completed transfer the vial into a laminar flow hood. Decontaminate the outer surface of the cryovial, and proceed immediately with the following steps under sterile conditions.
2. Within in a laminar flow hood, transfer aseptically the thawed cell suspension into a 15 ml centrifuge tube. Add 10 ml of fibroblast growth medium, seal the tube with its screw cap, and centrifuge it at 250xg for 5 min.
3. Remove aseptically the supernatant, and re-suspend the cell pellet in 10 ml of provitro's fibroblast growth medium using a serological pipette. **Check cell number and viability.**
4. Transfer the entire volume of HFIB-D suspension into two 25 cm<sup>2</sup> culture flasks (5 ml each). The advantage of using two 25 cm<sup>2</sup> flasks instead of one larger flask is that it helps to reduce the risk of losing all cells. So, if you experience difficulties in passaging the first 25 cm<sup>2</sup> flask, there is still another flask left for further use. In order to prevent contamination make sure that there are no traces of medium left on the inner/outer part of the flask neck.
5. Place the cell culture flasks in an incubator at 37°C, steam saturated and 5 % (V/V) CO<sub>2</sub>. Close the screw lids on each culture flask by half a turn only to allow gas exchanges to take place.
6. Change the fibroblast growth medium after the first 24 hrs to remove unattached cells, then every 48-72hrs. **Feed the cells only with culture medium that has been warmed up.**
7. The cells are ready for sub-culturing having reached a confluence of 75 %. If the cell layer is allowed to grow too dense they will suffer irreversible contact inhibition leading to ceased proliferation. **For subculturing, use the reagents recommended in the accompanying analysis certificate, only.**
8. Recommended seeding density of HFIB-D: > 4,000 cells per cm<sup>2</sup>

### Description:

Following provitro's standard operating procedures, the HFIB-D cultures are isolated from human tissue, then transferred into a primary cell culture, and finally aliquoted and frozen.

### Proliferative capacity of HFIB-D:

Provitro's HFIB-D cultures are derived from original tissue (in vivo state) using careful methods,. They are not transformed or mutated and have a limited *in-vitro* lifespan. All HFIB-D batches are tested by provitro for their proliferative capacity using provitro's culture reagents as outlined in the accompanying certificate of analysis.

### Quality control:

All HFIB-D cultures from provitro are subject to comprehensive quality tests, summarized in the accompanying certificate of analysis.

### Warning note:

Concerning use of biological material:

Provitro's primary cell cultures are of human or animal origin, and no known test procedures can ensure the total absence of infectious agents. All products should therefore be handled following safety precautions as if they were infectious.

### In vitro laboratory use only.

Not intended for any human or animal diagnostic or therapeutic use.