Stem Cell Preservation: Editorial

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Cell preservation is an essential technology for widespread distribution and applications of class cells. Ancient cryopreservation via slow-freezing or vitrification provides long-run storage, however, needs cytotoxic cryoprotectants (CPA) and tedious account loading/unloading, cooling, and convalescent procedures. Physiological condition storage around 0–4 °C is an alternate technique however solely works for a brief amount thanks to its high storage temperatures.

Here, we report on the deep-supercooling (DSC) preservation of human adipose-derived stem cells at deep subzero temperatures without freeze for extended storage. Enabled by surface protection with an immiscible oil part, cell suspension will be preserved in an exceedingly liquid state at −13 °C and −16 °C for seven days with high cell viability, retention of stemness, attachment, and multilineage differentiation capacities. These results demonstrate that DSC is an associate improved short-run preservation approach to supply off-the-peg cell sources for booming cell-based drugs and applied science.

Cell-seeded implants are an integral part of regenerative drugs strategy that aims to exchange blistered tissue and restore tissue function. Pluripotent stem cells are unit promising cell candidates for the event of regenerative drugs therapies as they need the power to self-renew and commit towards various cell sorts. In vivo, stem cells reside in an exceedingly dynamic niche, a stem cell-specific microenvironment that possesses chemical, biological and mechanical cues, that drive the vegetative cell fate and renewal. The affiliation between stem cells and their niche may be a two-way relationship consisting of each cell–cell interaction and cell–extracellular matrix (ECM) interactions.

An alternate regenerative drugs approach is that the manipulation of the vegetative cell microenvironment. Hence, novel methods are developed as well as 3D biomaterials and bioreactor technologies providing topographic, chemical and mechanical cues to recreate the vegetative cell niche. Understanding the mechanisms dominant vegetative cell fate and therefore the dynamic nature of the vegetative cell niche can alter researchers to duplicate this stem cell-specific microenvironment, and thus, harness and management the dear attributes that stem cells possess. This chapter elucidates the importance of pluripotent stem cells and their dynamic niche in regenerative drugs. It any presents novel methods to duplicate chemical, topographic and mechanical stimuli that area unit essential for the regulation of vegetative cell fate and therefore tissue regeneration.

Regenerative drugs may be a promising field that aims to develop therapies for presently unmanageable diseases. These approaches embody cell–cell and cell–scaffold implants. Self-renewal and therefore the ability of pluripotent stem cells to commit towards explicit cell lineages in response to mechanical, chemical and physical stimuli makes them the perfect building block for such therapies. Embryonic stem cells (ESCs) are unit isolated throughout embryological development. Their usage in regenerative drugs is controversial, and thus, adult stem cells are additional completely investigated for his or her potential use for tissue regeneration. Currently, evoked pluripotent stem cells (iPSCs) are unit being investigated as an alternate supply for pluripotent stem cells thanks to their origin and clinical potential.

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