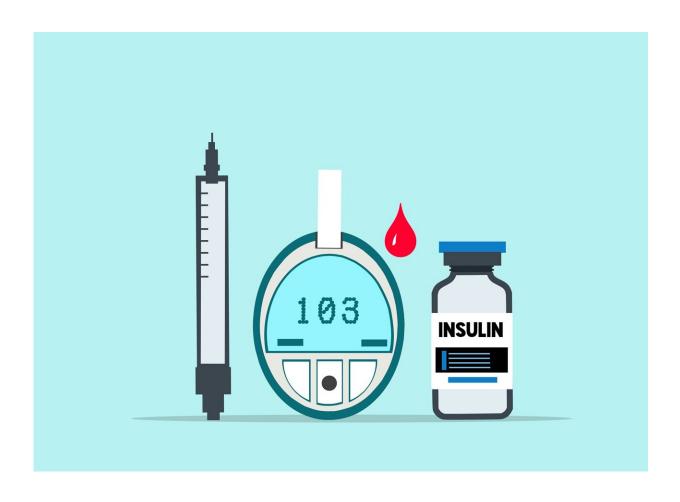


Improving potential stem cell-based treatments for patients with type 1 diabetes

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Following encouraging results from pre-clinical research and a recent clinical trial on stem cell-based treatments for patients with type 1



diabetes, researchers are now working on maximizing the function and potential of stem cell-based treatments for future application in patients.

More than 8 billion people worldwide live with type 1 diabetes (T1D) according to recent estimates. In T1D, the so-called <u>beta cells</u> in the pancreas do not make enough of the hormone insulin due to the destruction of beta-cells by the <u>immune system</u> leading to blood sugar levels rising above <u>normal levels</u>. Over time, T1D can lead to severe organ damage. There is no cure for T1D, and the <u>standard treatment</u> is daily supplementation of insulin, but some <u>patients</u> have difficulties maintaining their blood sugar levels within the normal range and risk developing complications from the disease.

An alternative treatment option being investigated for T1D patients is the transplantation of beta-cells, and news reports of a recently initiated clinical trial (Vertex NCT04786262) has generated hope that stem cellderived beta-cells could become an alternative and renewable treatment for patients with T1D, addressing the constant shortage of cadaveric beta cells.

In recent years, researchers have made great progress in generating large numbers of stem cell-derived beta cells (sBCs) in the lab, but current methods to make mature, functioning sBCs and to remove any remaining residual stem cells which may cause tumors in patients are costly and labor-intense, and therefore difficult to implement in large scale.

A new study published in the journal *Stem Cell Reports* has now come up with a simple method to address those issues. In their work, Holger Russ and colleagues from the University of Florida, U.S., found that dividing and potentially tumorigenic non-BCs can be eliminated from sBC cultures by a brief treatment with a chemotherapeutic drug.

This method not only prevented graft overgrowth and tumor formation



after transplantation into diabetic mice but also enhanced the maturation and functionality of sBCs. Accordingly, this method may reduce the time and cost involved for producing safe and efficient sBC transplants, and eventually contribute to enhancing access to this type of treatment for more patients with T1D.

More information: Holger A. Russ, Enrichment of stem cell derived pancreatic beta-like cells and controlled graft size through pharmacological removal of proliferating cells, *Stem Cell Reports* (2023). DOI: 10.1016/j.stemcr.2023.05.010. www.cell.com/stem-cell-reports 2213-6711(23)00189-3

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