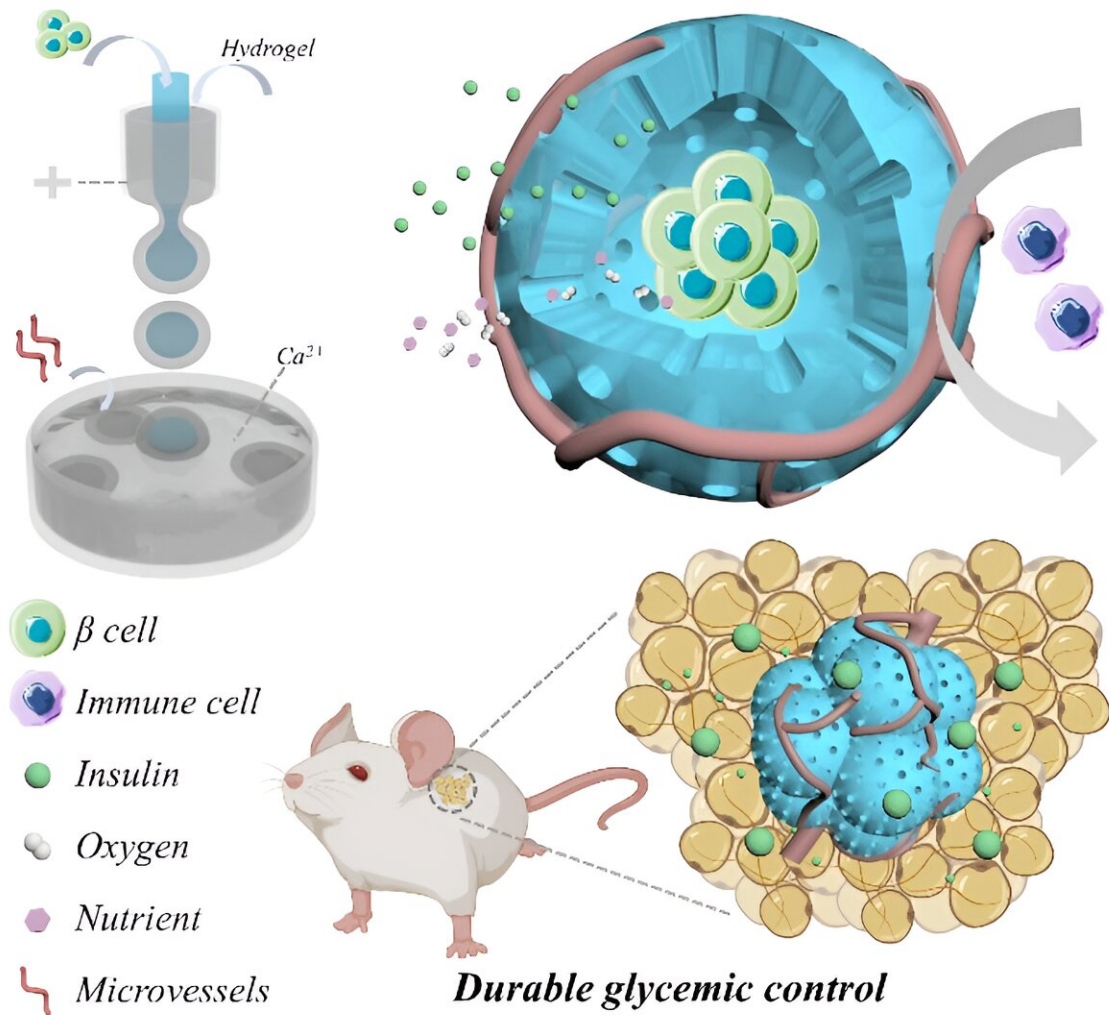


# Biomimetic artificial islet model: A new way to control high blood sugar

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Biomimetic artificial islet model for glycaemic control. Credit: Prof. Ling Li from Southeast University and Prof. Luoran Shang from Fudan University

Diabetes mellitus (DM) is a metabolic disease characterized by pancreatic malfunction and poor production of insulin. Imagine your pancreas is a factory for insulin, a key hormone that keeps your blood sugar level. Inside this factory are tiny assembly lines called pancreatic islets.

When these islets get damaged, it's like a mini-disaster in the factory. The assembly lines malfunction, and they can't produce enough insulin. This can lead to problems with regulating blood sugar, which is a big deal for your health.

For the most common condition resulting from islet damage, type 1 diabetes, there is no cure. Treatment focuses on managing blood sugar levels through [insulin injections](#). However, although exogenous insulin can temporarily stabilize blood sugar, frequent injections bring a heavy burden and may cause complications.

Islet transplantation, as an alternative strategy, faces the challenges of donor shortage and long-term immunosuppression. To address these issues, researchers have developed biomaterial encapsulating islets/islet cells to reconstruct islet function. Despite some progress, long-term survival and sustained insulin secretion are still constrained by immune attack and insufficient nutrient supply.

Microfluidic technology has been used to produce hydrogels with customizable structures due to its precise fluid manipulation capabilities. These microcapsules facilitate oxygen and nutrient exchange and possess adaptive structural features to encapsulate other drugs to regulate cell fate. Nonetheless, accurately simulating the complete structure of native islets remains challenging.

Current methods for recapitulating islets focus on the vascularization of the microcapsules.

Inspired by natural islet structures, a team of Chinese scientists developed a biomimetic artificial islet model based on microcapsules, which integrates a microvascular network and uses microfluidic high-voltage electrospray technology to achieve fine regulation of blood sugar levels.

These experimental results provide strong support for the prospects of vascularized microcapsules as a biomimetic artificial islet model and highlight their important value in the field of diabetes treatment. This innovative technology not only provides new treatment options for diabetic treatment, but also opens up new directions for future medical research.

Prof. Ling Li mentioned, "We had developed a vascularized microcapsule as a biomimetic artificial islet model (v-MCs) through a microfluidic strategy and used it for blood glucose control. The design of the microcapsules is inspired by the vascular network structure of natural islets. It has a unique core-shell structure that allows nutrients and oxygen to enter smoothly while effectively protecting the encapsulated islet cells from attack by host immune cells. In addition, v-MCs also can respond to glucose and exchange molecules.

"Through transplantation experiments in mice, we found that v-MCs can significantly improve the abnormal blood sugar levels, [food intake](#) and body weight of diabetic mice, and effectively improve glucose tolerance, showing its effective potential as a treatment for diabetes.

"Further histological analysis verified the safety of v-MCs implanted in vivo. These results indicate that the biomimetic artificial islet model is of great value in the treatment of diabetes, and its future application prospects are particularly broad in the fields of regenerative medicine and [tissue engineering](#), worthy of further research and expectations."

The research has been recently [published](#) in the journal *Materials Futures*.

**More information:** Jingbo Li et al, Biomimetic artificial islet model with vascularized microcapsule structures for durable glyceimic control, *Materials Futures* (2024). [DOI: 10.1088/2752-5724/ad47ce](https://doi.org/10.1088/2752-5724/ad47ce)

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