

# Fasting-mimicking diet may reverse diabetes

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Credit: CC0 Public Domain

A diet designed to imitate the effects of fasting appears to reverse diabetes by reprogramming cells, a new USC-led study shows.

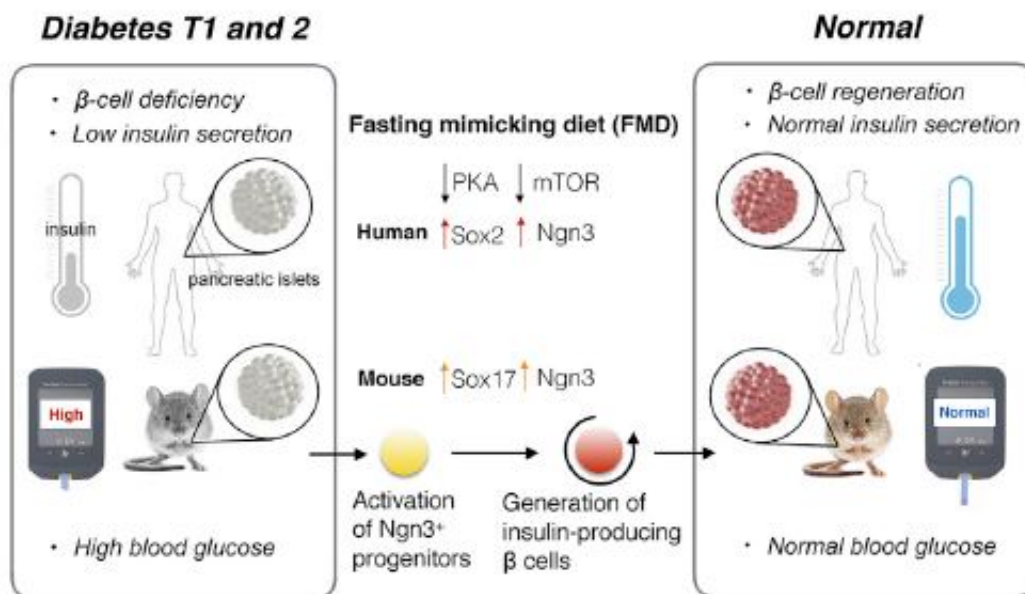
The fasting-like diet promotes the growth of new insulin-producing [pancreatic cells](#) that reduce symptoms of type 1 and type 2 diabetes in

mice, according to the study on mice and [human cells](#) led by Valter Longo, director of the Longevity Institute at the USC Leonard Davis School of Gerontology.

"Cycles of a fasting-mimicking diet and a normal diet essentially reprogrammed non-insulin-producing [cells](#) into [insulin-producing cells](#)," said Longo, who is also a professor of biological sciences at the USC Dornsife College of Letters, Arts and Sciences. "By activating the regeneration of pancreatic cells, we were able to rescue mice from late-stage type 1 and type 2 diabetes. We also reactivated [insulin production](#) in human pancreatic cells from type 1 diabetes patients."

The reprogrammed adult cells and organs prompted a regeneration in which damaged cells were replaced with new functional ones, he said.

The study published on Feb. 23 in the journal *Cell*, is the latest in a series of studies to demonstrate promising health benefits of a brief, periodic diet that mimics the effects of a water-only fast.



This visual abstract depicts the findings of Cheng et al., who show a short-term diet that mimics periodic fasting modulates b-cell number and promotes insulin secretion and glucose homeostasis with implications for both type 1 and type 2 diabetes. Credit: Cheng et al./Cell 2017

## Reversing insulin resistance and depletion

In type 1 and late-stage type 2 diabetes, the pancreas loses insulin-producing beta cells, increasing instability in blood sugar levels. The study showed a remarkable reversal of diabetes in mice placed on the fasting-mimicking diet for four days each week. They regained healthy insulin production, reduced [insulin resistance](#) and demonstrated more stable levels of blood glucose. This was the case even for mice in the later stages of the disease.

The diet cycles switched on genes in the adult mice that are normally active only in the developing pancreases of fetal mice. The genes set off production of a protein, neurogenin-3 (Ngn3); thus, generating new, healthy [insulin-producing beta cells](#).

## Next steps: clinical study

Longo and his team also examined pancreatic cell cultures from human donors and found that, in cells from type 1 diabetes patients, fasting also increased expression of the Ngn3 protein and accelerated insulin production. The results suggest that a fasting-mimicking diet could alleviate diabetes in humans.

Longo and his research team have amassed evidence indicating several health benefits of the fasting-mimicking diet. Their study published [last week](#) in *Science Translational Medicine* demonstrated that the fasting-

mimicking diet reduced risks for cancer, diabetes, heart disease and other age-related diseases in human study participants who followed the special diet for five days each month in a three-month span.

Prior studies on the diet have shown potential for alleviating symptoms of the neurodegenerative disease multiple sclerosis, increasing the efficacy of chemotherapy for cancer treatments, and decreasing visceral fat.

"These findings warrant a larger FDA trial on the use of the fasting-mimicking diet to treat human [diabetes patients](#) to help them produce normal levels of insulin while improving insulin function," Longo said. "Hopefully, people with diabetes could one day be treated with an FDA-approved fasting-mimicking [diet](#) for a few days each month and gain control over their insulin production and blood sugar."

**More information:** Cheng et al: "Fasting-mimicking diet promotes Ngn3-driven  $\beta$ -cell regeneration to reverse diabetes" *Cell*, [DOI: 10.1016/j.cell.2017.01.040](#)

Min Wei et al. Fasting-mimicking diet and markers/risk factors for aging, diabetes, cancer, and cardiovascular disease, *Science Translational Medicine* (2017). [DOI: 10.1126/scitranslmed.aai8700](#)

Provided by University of Southern California

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