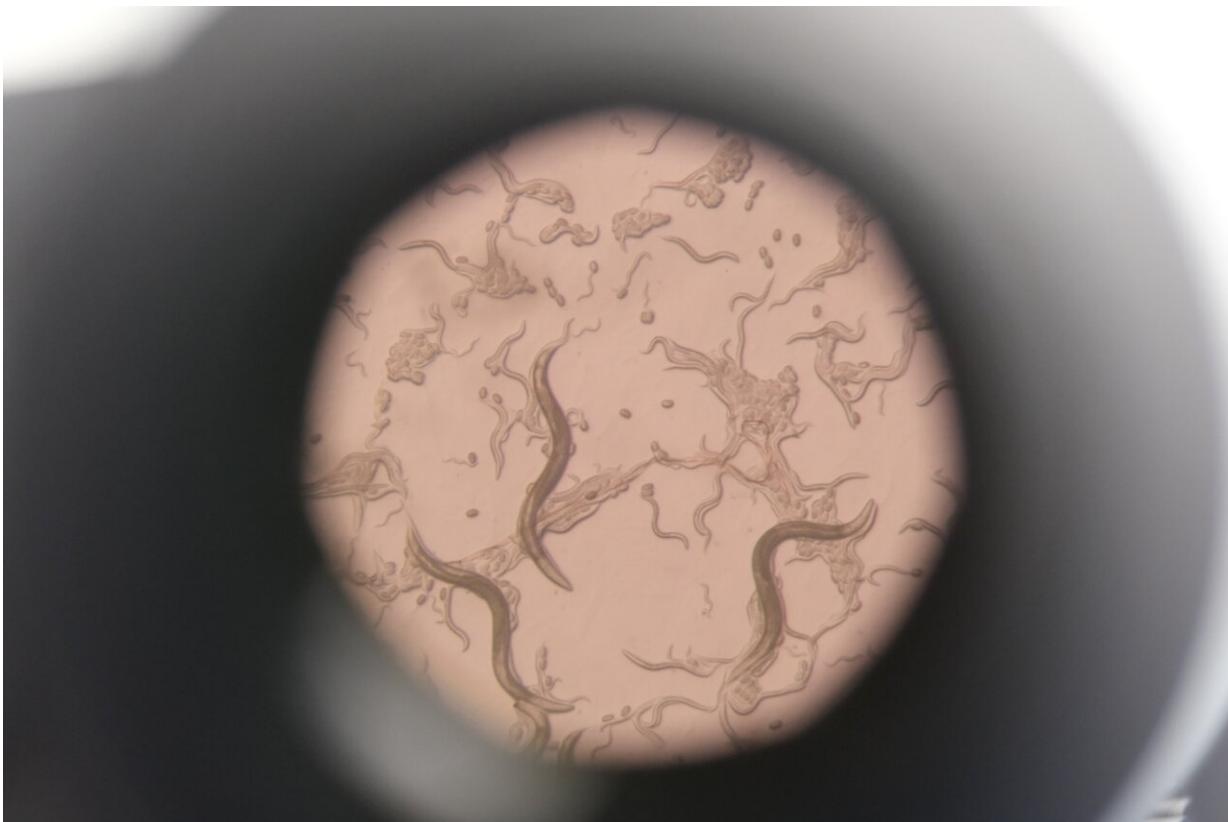


Manipulating stress response in cells could help slow down aging

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The lab experiments were conducted on *C. elegans*, a type of roundworm. While *C. elegans* is significantly anatomically simpler than a human, it relies on many of the same genes that humans do to control the division of cells and to programme faulty cells to die. Credit: NTU Singapore

Scientists at Nanyang Technological University, Singapore (NTU

Singapore) have found that a stress response in cells, when "switched on" at a post-reproductive age, could be the key to slow down aging and promote longevity.

In lab experiments on a type of roundworm that shares similarities with humans, the NTU Singapore team found that switching on this [stress response](#) in aged worms by feeding them a high-glucose diet extended their lifespan as compared to worms fed a normal diet.

This is the first time a link between this stress response and aging has been uncovered, said the NTU team of their findings published in *Nature Communications*.

While further studies are needed to gain a deeper understanding of this link, the scientists said their findings open the door to the development of therapies that could delay the onset or even tackle age-related disorders such as cancer, dementia, and stroke.

Cell biologist and study lead Associate Professor Guillaume Thibault from the NTU School of Biological Sciences says that "aging is a critical risk factor for a variety of human pathologies, from metabolic diseases such as diabetes to cancer and neurodegenerative diseases. From a public health perspective, determining the cellular pathways that underpin the aging process could take us one step closer to developing novel therapeutic strategies to treat age-related disorders."

"While our study found that a high-glucose diet could be useful to slow down aging and promote longevity in aged worms, we are not recommending that the aged population should now turn to a [high-sugar diet](#). What this study does show is that triggering certain stress responses in cells may translate to longevity, and that activating this stress response with a drug might be critical to decelerate cellular aging."

Aside from showing that the effect of manipulating this stress response in aged worms, the NTU scientists also showed that the same response, when 'switched off' in young worms fed a high-glucose diet, helped them to live longer than worms on a normal diet.

Commenting as an independent expert, Professor Rong Li, Director of the Mechanobiology Institute at the National University of Singapore says that "metabolic diseases have serious consequences in the elderly if left untreated. This work is impactful because the scientists identified a cellular pathway, called the unfolded [protein](#) response, which affects lifespan in animals fed a high glucose diet."

"They found that inhibiting this pathway dramatically extended the lifespan of these animals. They therefore propose that targeting this pathway may extend lifespan in humans with metabolic disorder."

This study is aligned with the research pillar of the University's NTU2025 five-year [strategic plan](#), which focuses on health and society as one area with potential for significant intellectual and societal impact.

How the cell's stress response is activated

Cells produce a stress response when stressors (such as an excess of glucose) cause a build-up of problematic "unfolded" proteins in the cell. The stress response, called the unfolded protein response, works to clear up these problematic proteins to restore balance in the cell.

Aging could also lead to an accumulation of unfolded proteins due a natural decline in the ability of the cell's machinery to produce healthy proteins, triggering the same stress response.

The molecular machinery in the cell tackles this build-up through its "stress sensors," which initiate a series of molecular mechanisms to

rescue the cell from this stress. If the overload of unfolded proteins is not resolved, the prolonged unfolded protein response induces cell death instead.

Unfolded protein response in aged worms led to healthier aging

To investigate how the unfolded protein response affects longevity in animals, the scientists induced this response in adult roundworms (*Caenorhabditis elegans*) using glucose. While *C. elegans* is significantly anatomically simpler than a human, it relies on many of the same genes that humans do to control the division of cells and to program faulty cells to die.

The scientists fed some of the worms a high-glucose diet at two different life stages: young i.e. at the start of their adulthood (Day 1), and at a post-reproductive age (Day 5), when the worms are aged and no longer fertile. A control group of worms were fed a normal diet throughout.

The scientists found that the aged worms given a high-glucose diet lived for 24 days—almost twice the lifespan of the young worms given the same diet (13 days). Worms on a normal diet lived for 20 days.

Aside from living longer, the aged worms on a high-glucose diet were more agile and had more energy storage cells as compared to worms given a normal diet, suggesting healthier aging.

Prolonged stress response in young worms led to cell death

A day after feeding the worms a high-glucose diet, the NTU scientists monitored the activity of the three stress sensors that are each responsible for a cellular pathway in the unfolded protein response.

They found that one of the stress sensors, IRE1, was significantly more active in young worms compared to aged worms.

When the scientists removed the gene coding for IRE1 in worms to "switch off" the cellular pathway the stress sensor initiates, they found that young worms fed a high-glucose diet from Day 1 lived for 25 days—twice as long as when the IRE1 gene was intact.

This suggests that the increased activity of stress sensor IRE1 seen in young worms fed a high-glucose diet from Day 1—what the scientists say is a prolonged unfolded protein response—was responsible for shortening their lifespan.

Assoc Prof Thibault says that they "believe that the high-glucose diet fed to the aged worms stimulated their otherwise sluggish [unfolded protein response](#) and switched on certain [cellular pathways](#), tackling not just the stress caused by excess glucose but also other aging-related stress, restoring cellular stability."

"In contrast, young [worms](#) subjected to a high-glucose diet provoked unresolved stress in the cells due to an overactivated IRE1. This prolonged activation led the cells to initiate cell death instead."

The findings suggest that a drug that reduces the activity of IRE1 while increasing the activity of the other two [stress](#) sensors could potentially be developed to decelerate cellular aging and consequently extend lifespans, he added.

More studies and findings will need to be conducted in the roundworms to further dissect the complex mechanism behind the lifespan extension induced by a high-glucose diet, as well as how this mechanism interacts with other processes in [cells](#).

More information: Caroline Beaudoin-Chabot et al, The unfolded protein response reverses the effects of glucose on lifespan in chemically-sterilized *C. elegans*, *Nature Communications* (2022). [DOI: 10.1038/s41467-022-33630-0](#)

Provided by Nanyang Technological University

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