

# Root molecular cause of declining health in the old found by scientists

February 10 2011, By Bill Schaller

---



Harvard scientists found that the basic cause of age-related health decline is malfunctioning telomeres — the end caps on cells’ chromosomes that protect them against DNA damage. Credit: Kris Snibbe/Harvard Staff Photographer

Scientists at the Harvard-affiliated Dana-Farber Cancer Institute say they have identified the root molecular cause of a variety of ills brought on by advanced age, including waning energy, failure of the heart and other organs, and metabolic disorders such as diabetes.

“What we have found is the core pathway of aging connecting several

age-related biological processes previously viewed as independent of each other,” said Ronald A. DePinho, senior author of a report posted online by the [journal Nature](#). The first author, Ergun Sahin, is a member of the DePinho Lab and an instructor in medicine at Harvard Medical School (HMS).

DePinho, who is the director of Dana-Farber’s Belfer Institute for Applied Cancer Science and also a professor of medicine at HMS, said that although the studies were conducted in mice, “The findings bear strong relevance to human aging, as this core pathway can be directly linked to virtually all known genes involved in aging, as well as current targeted therapies designed to mitigate the toll of aging on health.”

The scientists found that the basic cause of age-related health decline is malfunctioning telomeres — the end caps on cells’ chromosomes that protect them against DNA damage. As cells reach their predetermined limit of times that they can divide, the telomeres become shortened and frayed, making the chromosomal ends vulnerable to increased rates of unrepaired DNA damage.

Faced with this increasing reservoir of injured DNA, cells activate a gene, p53, that sounds an alarm and shuts down the cells’ normal growth and division cycle, ordering them to rest until the damage can be repaired or, if not, to self-destruct.

Scientists previously had blamed this emergency shutdown and cell death for age-related deterioration of organs whose cells divide rapidly and are rejuvenated by reserves of adult stem cells. Such tissues include skin, intestinal lining, and blood cells, among others, which generate trillions of new cells each day of life.

However, left unanswered is how cells with less cell division, such as the heart or the liver, sustain equivalent levels of aging. The scientists felt if

they could solve this mystery, they might gain new insights into how DNA damage could lead to age-related decline across all organs.

The new findings demonstrate that the telomere dysfunction and activation of p53 also trigger a wave of cellular and tissue degeneration that links telomeres to well-known mechanisms of aging that are not simply related to rapid growth and division. In other words, telomere dysfunction is not just one culprit in the declining health of advanced age. It's the kingpin, according to DePinho and his colleagues.

DePinho published a study in *Nature* in January 2011 that demonstrated it was possible to reverse the symptoms of extreme aging in mice by increasing their levels of telomerase, the enzyme that maintains the health of the telomeres.

In this new, larger role, the telomere dysfunction also sets off an array of reactions leading to diminished health and longevity. For example, muscles suffer a loss of mitochondria, a cell's chemical power plant, causing waning vitality and failure of the heart and other organs. Risks of metabolic disorders such as [diabetes](#) are increased.

In addition, the process weakens the body's antioxidant defenses against the damaging molecules known as reactive oxygen species, or "free radicals," that accumulate with age and exposure to stress. Until now, some researchers had labeled the decline in mitochondria or the buildup of free radicals as the primary causes of age-related ills. The new work integrates these seemingly disparate mechanisms into one unified theory of aging.

Telomere dysfunction causes this wave of metabolic and organ failure, the scientists found, because when the p53 gene is activated, it represses the functions of two master regulators of metabolism, PGC1-alpha and PGC1-beta. This dialing down of the regulators diminishes metabolic

processes needed to provide energy and resist stress. In the mouse experiments, the scientists showed that “knocking out” p53 in mice released the brakes on PGC1-alpha and PGC1-beta.

“This is the first study that directly links telomere dysfunction to regulators of the mitochondria and antioxidant defense via p53,” DePinho said. “The discovery of this new pathway of aging integrates a lot of different ideas people have had and gives us a better understanding of the aging process.”

By unifying several major pathways of aging under the umbrella of telomere dysfunction, he said, the findings may yield new targets for therapies. The discoveries also may underlie the relatively sudden and rapid failure of the body leading to the end of life.

“Because telomere dysfunction weakens defenses against damage by free radicals, or reactive oxygen species,” DePinho said, “we think this exposes telomeres to an accelerated rate of damage which cannot be repaired and thereby results in even more organ deterioration. In effect, it sets in motion a death spiral.”

Provided by Harvard University

Citation: Root molecular cause of declining health in the old found by scientists (2011, February 10) retrieved 19 April 2024 from <https://medicalxpress.com/news/2011-02-root-molecular-declining-health-scientists.html>

<p>This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.</p>
--