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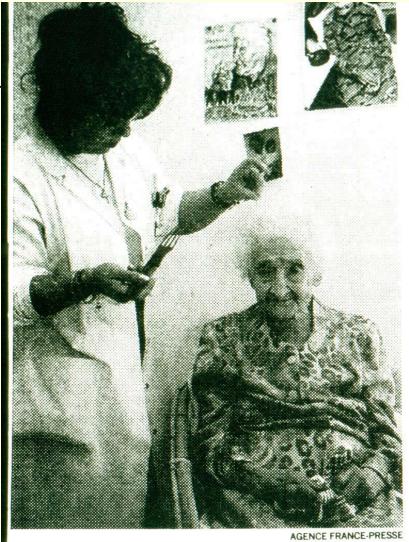


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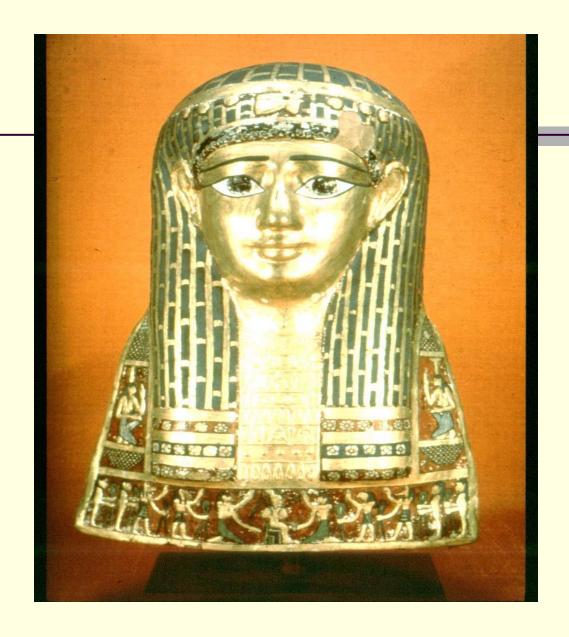
Is there a role for Stem cells in Healthy Ageing

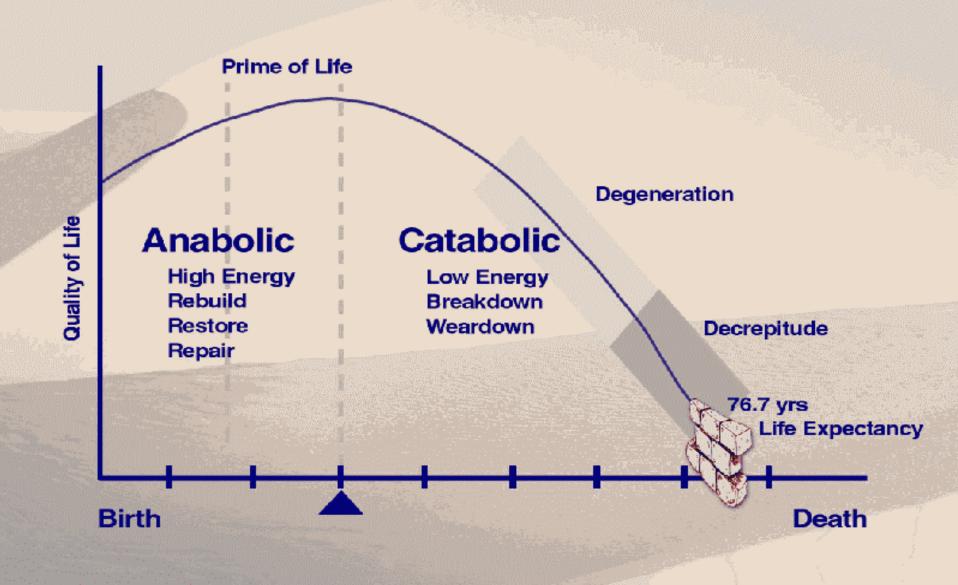
Dr. Rajbans Singh MBBS, MRCP, Dip. Geriatric Medicine Consultant Physician & Gerontologist Healthy Ageing Specialist Pantai Medical Centre Beacon International Specialist Centre

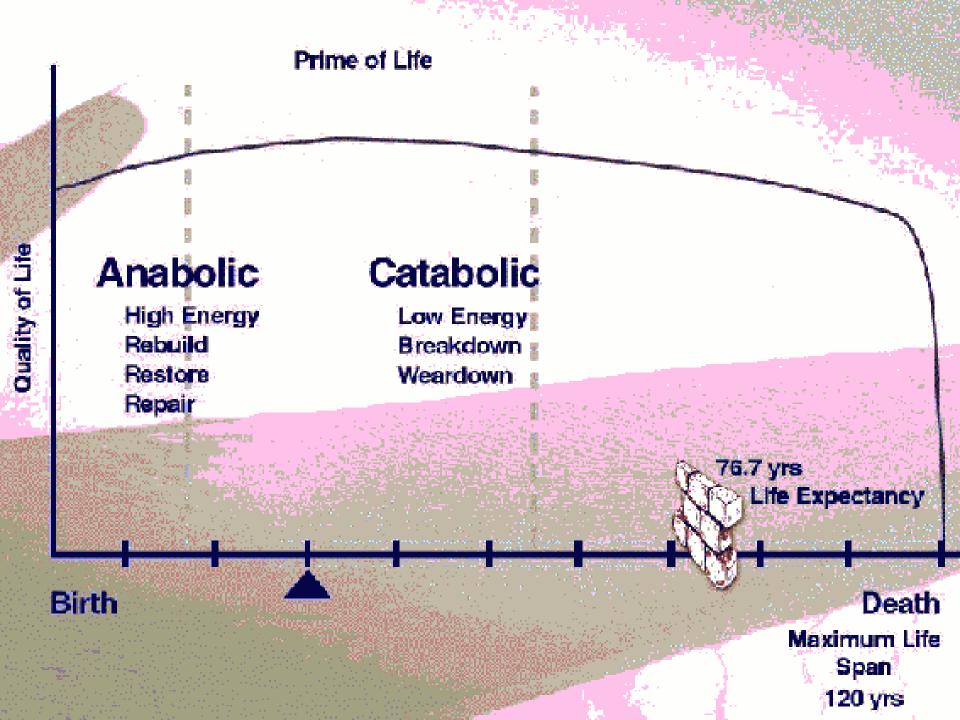


Jeanne Calment gets ready for her 120th birthday, which Guinness says is the current world record.

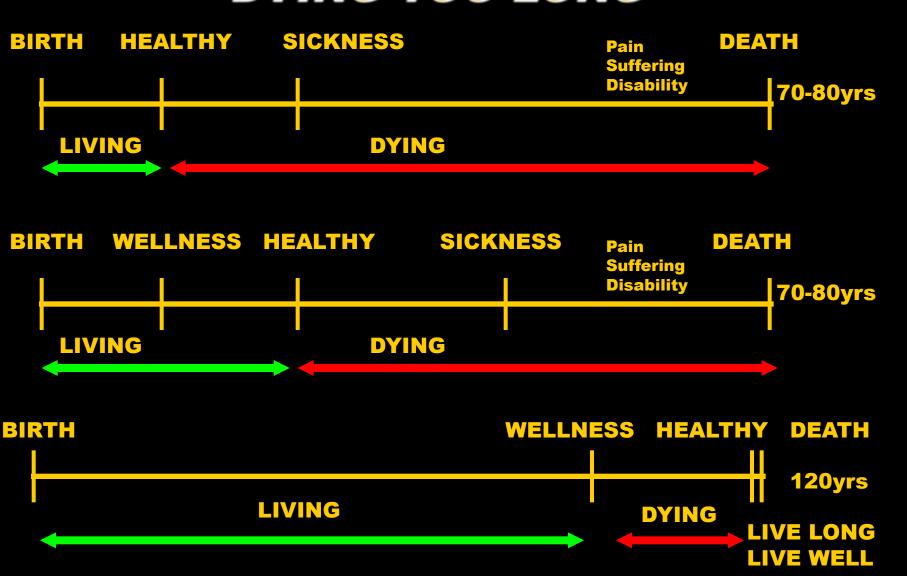
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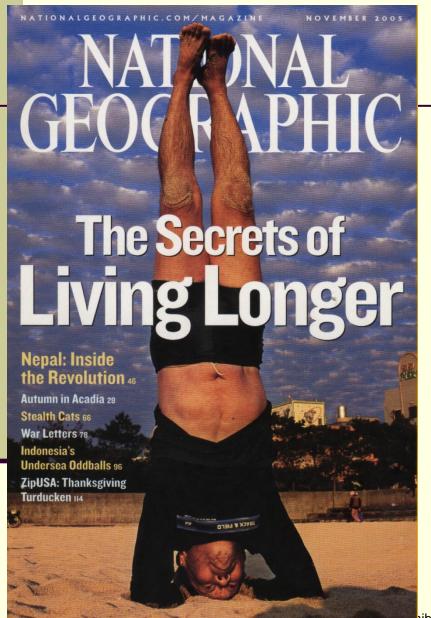


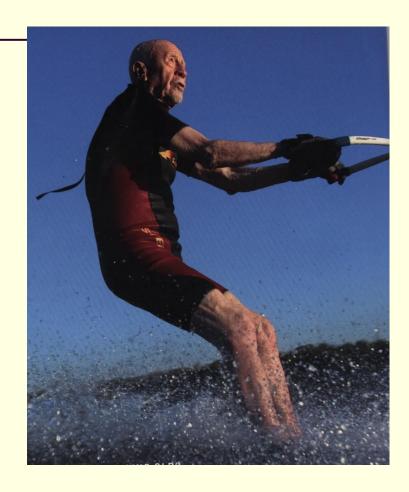




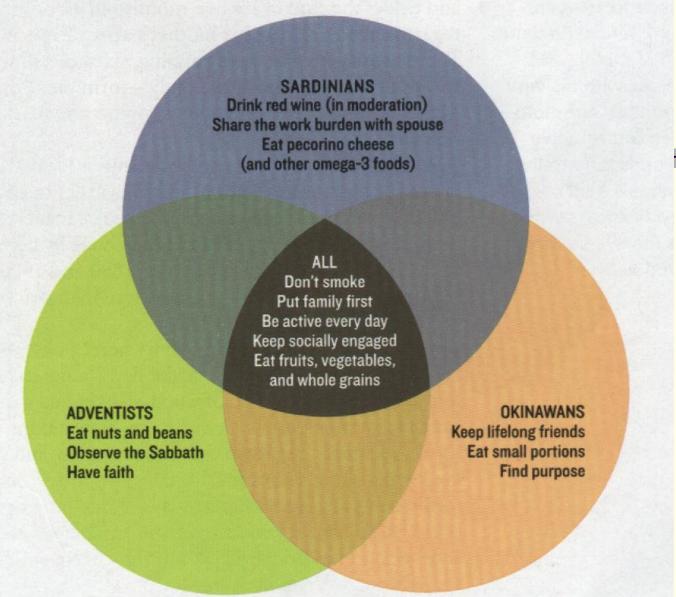
PEOPLE ARE LIVING TOO SHORT AND DYING TOO LONG







ש טו. Kajbans Singh March 2005



HOW THEY LIVE LONGER

Super seniors in three widely separated regions share a number of key habits, despite many differences in backgrounds and beliefs.



THE SCIENCE BEHIND RADICAL LIFE EXTENSION

FANTASTIC VOYAGE

LIVE LONG ENOUGH TO LIVE FOREVER

RAY KURZWEIL

TERRY GROSSMAN
M.D.

- Ray Kurzweil
- Famed Inventor & Computer Scientist
- Top MIT graduate Lemelson-MIT prize
- 1st Reading m/c for the blind
- 11 Doctorate's
- Fantastic Voyage



President Clinton presented Ray Kurzweil with the National Medal of Technology, the nation's highest honor for technological achievement.

"FIRST BRIDGE

- Living a healthy lifestyle to keep people fit enough to cross the second bridge
- Staying focus and discipline on health regimes with key emphasis on WELLNESS

• By keeping your body's internal biochemistry stable and healthy, allowing your body to commit its energy and resources into cell repair and rejuvenation from the inside, you will stay young and healthy.

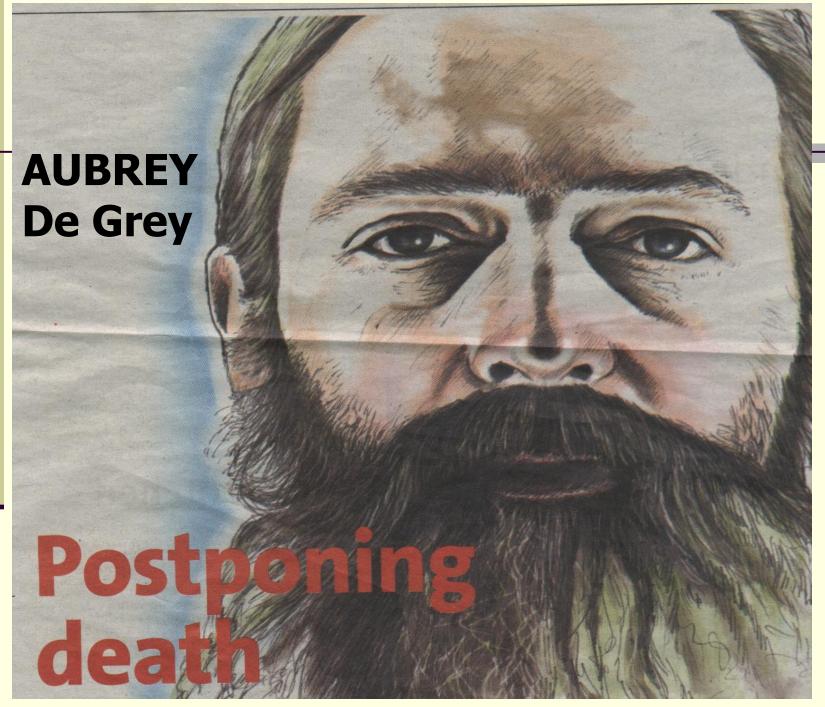
"SECOND BRIDGE

- A Biotechnological revolution
- Humanity is on the verge of controlling how genes express themselves & ultimately changing the genes
- Could block disease causing genes and introduce new ones that would slow or STOP the ageing process

"THIRD BRIDGE

NANOTECHNOLOGY and ARTIFICIAL INTELLIGENCE REVOLUTION!

- Will deliver the nanobots that work like repaving crews in our blood streams and brains
- These intelligent machines will destroy disease, rebuild organs and wipe out known limits on human intelligence



26 Features

In.Tech

The battle against __ageing begins



THEORIES OF AGEING

- Social theories
- Biological theories

Biological Theories

- Programmed aging theory (Cellular aging)- aging genes
- Cross link theory Rate of DNA repair decreases as the cells age (Telomerase connection)
- Theories of Random deterioration Free radical reactions (anti-oxidants)
- 4. Immunologic theory
- 5. Stress theory
- 6. The neuro endocrine theory hormones
- 7. Glycosalation theory Metabolic syndrome.



Personal Health Pyramid



Stem Cells

Bioidentical Hormones & Metabolic Syndrome

Free Radicals
& Immune System

Nutrition, Exercise and Lifestyle Choices

Purpose, Goals, Dreams, Mind-Body

(DNA) Personal Genetic Profile

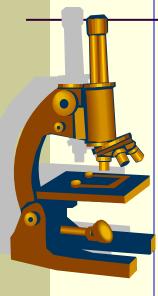
Understanding The Human Body

WELLNESS

SICKNESS

ORGANS

HUMAN BODY

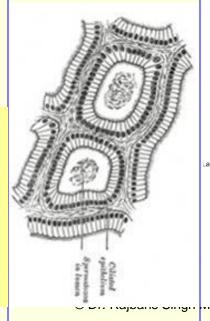


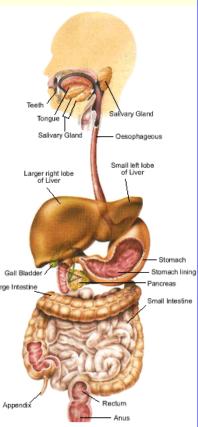
CELL

MOLECULE

loading...









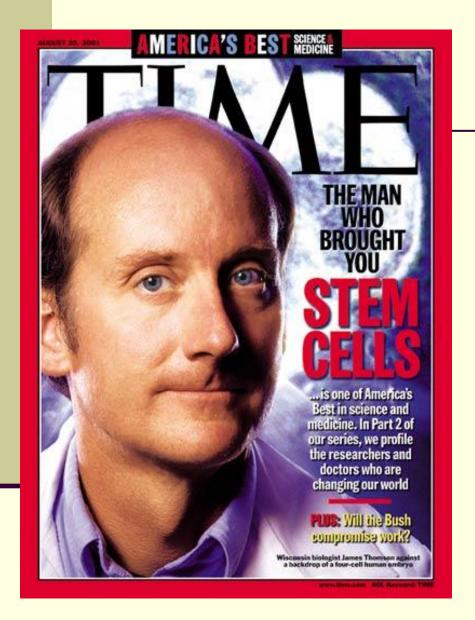
Top Five Biotechnologies

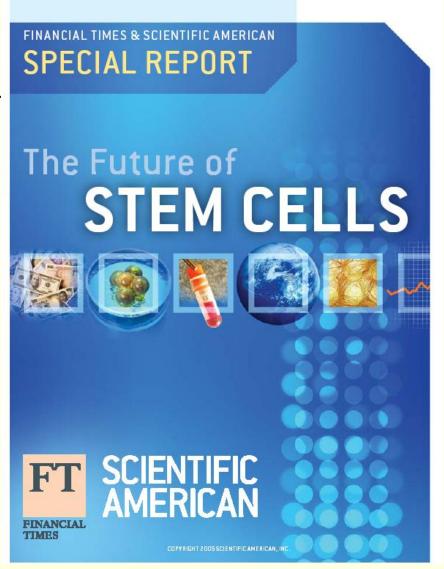
- 1. Stem Cells
- 2. Cloning
- 3. Nanotechnology
- 4. Artificial organs
- Nerve impulse continuity (brain/spinal cord)











What are Stem Cells?

Stem cells are highly plastic, multipotential cells with the capacity to self replicate; that is, they can generate all the cell types composing more specialised stem cells and then ultimately a functional organ or tissue, while maintaining a pool of cells like themselves for the lifetime of the individual.

Stem cells – The Renewal System

A number of recent studies collectively suggest that health is a balance between cellular loss taking place as part of normal process of tissue turnover or ageing and Bone Marrow based tissue regeneration.

Diseases would form out of an imbalance between tissue turnover and tissue repair.

Stem cells – The Renewal System

- Two aspects must be considered for health maintenance and disease prevention
- Enhancing tissue repair/renewal by supporting stem cell function
- Preventing conditions that might accelerate cellular loss by adopting lifestyle strategies that support health at the cellular level.

What are stem cells useful for

- The Four R's of Stem Cell Therapeutics
- Replace
- 2. Repair
- 3. Regenerate
- 4. Rejuvenate

These processes recapitulate/mimic natural events of developmental biology

What are stem cells useful for?

1. Replace – Many diseases such as Parkinson's Disease and juvenile diabetes result from the death or dysfunction of just one or a few cells. Therefore, stem cells can be used as replacement cells, offer "lifelong treatment".

What are stem cells useful for?

2. Repair – Stem cells can be used to repair diseased or damaged organs. By isolating stem cells in a laboratory, scientists could grow new heart cells to repair damage from heart attacks, new liver cells to treat hepatitis and new red blood and stromal cells for cancer patients after ablative radiotherapy.

What are Stem cell useful for?

- 3. Regenerate Embryonic or TR stem cells can be used to renew biological functions or damaged organs.
- 4. Rejuvenate Stem cells can to used to become new, healthy "younger" cells to rejuvenate, restore and repair older cells and ailing hearts, liver, brains and other organs.

Perhaps a 5th "R"?

Rebalance the body's biochemical processes definitely, obviating the need for classic, costly, repetitive pill-taking, injections and surgery.

Pilot Study

- Pilot study to determine the efficacy and safety of allogenic mesenchymal stem cells adminstration for healthy ageing.
- 50 million allogenic stem cells given 1st month and repeated after 6 months.
- Or 100 million cells once.
- Intravenous infusion over 30 minutes.

Pilot Study

- Measurements:
- 1. Quality of life Questionairre
- Body Composition
- Full blood profile including hormones and cancer markers.
- 4. Heart scan, U/S abdomen, mammogram
- Bone Density

Pilot Study

- Measurements were before 1st injection and 6 months after last injection.
- 50 patients underwent the therapy.
- Data collected from 27 patients.
- All improved in QOL questionnaire ?placebo
- 25 out of 27 had an increase in IGF1 between 30-40%.
- Almost all had same levels or decrease in cancer markers.

Study Protocol - Title

A Randomised, Double Blind, Multicentric, Placebo Controlled, Phase 11 Study Assessing the Safety and Efficacy of Intravenous Ex-Vivo Cultured Adult Allogeneic Mesenchymal Stem Cells in Improving Aging Parameters in Older Individuals.

Study Protocol

- A total of 72 patients are planned to be randomised at one study centre in Malaysia.
- Clinical trial duration will be for 2 years.
- Primary Objective- To assess the safety and tolerability following a single dose of allogenic mesenchymal stem cells administered by intravenous injection in improving the ageing parameters in older individuals.

Study Protocol

Secondary Objective – To explore the efficacy of allogenic mesenchymal stem cells in improving the ageing parameters in older individuals and to find the appropriate dose of allogenic mesenchymal stem cells.

Study Protocol

The primary end points are safety and tolerability, assessed by adverse events

The secondary endpoints are:

- 1. Clinical Evaluation
- 2. Hormones esp. IGF1
- 3. Body Composition
- 4. Bone Density
- 5. Ferrans and Powers QOL index generic version 111.

THE END OF DYING

By studying the genetics of aging, scientists are traversing the road to immortality

BY WILLIAM LAI

here is no mistaking the onset of aging. Fine wrinkles begin to appear on the face. Hair starts to turn gray. The unaided eye encounters problems focusing on nearby objects and eventually loses the ability to distinguish fine details. Men, especially, find it increasingly difficult to hear higher sound frequencies. The heart grows larger while the maximum breathing capacity of the lungs declines. The bladder progressively holds smaller amounts of liquid. Body fat migrates to hips and thighs in women and to the gut in men. As time passes, chronic illnesses like osteoporosis, arthritis and Alzheimer's disease strike with little warning.

Traditionally, scientists believed that aging was a biological process in which cells simply stopped dividing. Gerontologists, who specialize in aging, now know the explanation is not that simple. One theory links aging to cumulative cell-division errors. Another attributes it to the shrinkage of telomeres, the structures at the tip of chromosomes. A third hypothesis suggests a connection with the build-up of so-called junk DNA, genetic material that does not have any obvious use. Other researchers zero in on environmental factors such as high-calorie diets. A common thread runs through all these studies: the belief that in unlocking the secrets of aging, doctors can devise ways of retarding and even reversing

Genetics. Some light was shed on the mystery in 1996 when Chang-En Yu, Junko Oshima, Ying-Hui Fu and other researchers identified the first human gene that affects aging. The breakthrough came from studies of people with the rare condition known as Werner's Syndrome. As they enter adoles-

diverse in through the second of the second

MEASURE OF LIFE If a way could be found to repair shortened telomeres (orange structures), humans might live longer

A common thread runs through all these studies: the belief that doctors can retard and even reverse the process of aging cence, sufferers develop wrinkled skin, lose their hair and become susceptible to disease associated with the elderly such as cataracts, heart disease, diabetes and cancer. The researchers traced the abnormalities to mutations in the WRN gene, which controls the production of an enzyme belonging to a group known to play a crucial role in manipulating genetic material. The hope is that reengineering the defective WRN gene (it has been successfully cloned) could help cure Werner's Syndrome — and perhaps extend natural lifespans.

A California team led by Richard Lerner, president of the Scripps Research Institute, has embarked on another approach. The researchers theorize that the mechanism of human aging and its associated diseases can be traced to a gradual increase in cell-division errors in tissues

throughout the body. "This functional change begins slowly in middle age and increases gradually

with advancing age," wrote
Lemer in a recent issue of
the journal Science. The
theory is that those errors
gradually alter the way
key genes operate,
which in turn causes
the loss of tissue functions that result in
aging. Imagine your
cells as tiny gene factories where quality control of the manufacturing process naturally
declines with the passage
of time.

Lerner and his colleagues studied genes in actively dividing cells from young, middle-

aged and old patients, and those with progeria, another rare genetic disorder characterized by accelerated aging. The researchers found that 61 out of more than 6,000 genes examined showed consistent changes in expression - that is, there were slight differences in their characteristics depending on the age of the donor and whether the patient had progeria. Some of the 61 genes are linked to chronic diseases such as age-related breast cancer and arthritis, and defects in organs like the kidney. heart and ovaries. Lerner cautions that the research is not conclusive. Further studies are needed before the process of aging can be completely understood.

Other researchers focus on telomeres, which are found on the tips of chromosomes (see picture above). Every time a chromosome replicates itself, its telomeres shorten in length. The older an organism is, the shorter are its telomeres. Thomas Wan

ACTIVE 100

Come along and grow old with me, the best is yet to be.

Robert Frost

THANK YOU