Immortality: The probable future of human evolution

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Introduction

Much the same as life, death is a natural phenomenon. Human have longed to be immortals and this is reflected in the beliefs of most, if not all, religions. In this article, brief overview of some of the immortal biological systems, both at the cellular and organismal levels are highlighted. Assumptions of the author on immortality and the probable future of human evolution are also discussed.

Key words: Cancer, HeLa, immortality, mortality, telomerase, Turritopsis dohrnii.
tales. The influence of superheroes and mytho-
llogical figures on our psychological construct is
so immense that very often one desires to be
immortal. It is amazing to learn that many young
beings wished to become the Marvel superhero,
Wolverine, of X-Men, perhaps because of his
regenerative ability. Come on, if a bullet or an
atom bomb can’t terminate your existence on
Earth, how cool would that be? But, young read-
ers of this article must be warned with much re-
gret that he is only a beloved fictitious character
of the entertainment industry. Oh! Don’t be dis-
heartened so hastily. You are about to read ac-
counts of non-fictitious biological immortals,
indeed, the real deal of life. Let us begin without
a backbone.

The Transparent Immortals

The seahorse (*Hippocampus* sp.) looks like a
limbless horse (*Equus ferus*), but a jellyfish does
not look anything like members of the class Pi-
sces. Perhaps, many of us have heard and seen a
ejellyfish, in real life or at least on papers or digital
graphics. They are free swimming aquatic ani-
mals found in every ocean, from the surface to
the depth of seas. They belong to the phylum
Cnidaria and have a somewhat gelatinous dome-
shaped body with varying numbers of protrud-
ing posterior tentacles. Their transparent body
color somehow looks harmless. They look harm-
less alright, but looks can often be very decep-
tive! Their tentacles can be used for capturing
prey or as a defense against invading predators
by discharging toxins that cause excruciating
pain to the victims.  

The life cycle of a typical jellyfish may be di-
vided into two phases, sexual and asexual. In the
sexual phase they are motile and in the asexual
phase, they are not. The sperm and egg from
solitary, sexual adult jellyfishes called medusae
(sing. medusa) fertilized and developed into an
embryo that further developed into planula
larva. The planula larvae formed a colonial polyp.
The solitary sexual medusa are formed asexually
from the colonial polyp. These medusae have

Figure 1 | A seahorse (*Hippocampus*).

Figure 2 | Life cycle of jellyfish.
limited lifespan and usually die shortly after fertilization. However, a not so typical jellyfish species called *Turritopsis dohrnii* (formerly classified as *Turritopsis nutricula*) have the ability to skip the “death” part of their lifecycle. Of course they can die the old fashion ways, like being eaten by other predators. But when they faced physical damage, starvation or stress, they can revert back to their baby form, i.e. polyp stage, by a process called transdifferentiation, and start their life cycle all over again, with no change in their genetic composition, hence, they are essentially the same individual. In transdifferentiation, mature somatic cells get transformed into another whole new type of cells. In simple terms, it’s like the liver cells becoming the skin cells. This amazing life reversal ability of this particular jellyfish has been observed in both wild and the laboratory, so, technically they are immortals.

If you were offered the choice, to revert back to baby, just moments before you inhale your last breath, will you take it? I might, if I am allowed to keep my diaries.

**Illustrious Lady**

This ordinary lady is astounding and immortal, figuratively in a way. She may be considered both dead and alive (this sounds like Schrödinger’s cat!). Let me elucidate the point. If we consider life of an individual at the organismal level, she is indeed, very dead. But if we consider it at the cellular level, she is still very much alive. She, in fact her cells, do not require any introduction among biologists. But for those of us who might still be in the dark let me introduce to you Mrs. Henrietta Lacks, more famously known as HeLa, of the HeLa cell line.

She was also known by the name Helen Lane or Helen Larson, but her real name is in fact, Henrietta Lacks. Amazingly, her birth name is Loretta Pleasant, but no one knows how she became Henrietta Lacks. By today’s terminology, she will be referred to as an African-American, or simply Black. On 29th January, 1951, after experiencing abnormal vaginal bleeding and a feeling of “knot” in her womb for some time, Henrietta went to John Hopkins Hospital (Baltimore, Maryland, USA) gynecology ward where she was examined by Dr. Howard Jones and was diagnosed with epidermoid carcinoma of the cervix. Dr. Lawrence Wharton Jr. took samples from her and sent them to Dr. George Gey for tissue culture. Gey had spent around thirty years of his professional life trying to grow human cancer cells outside the body but failed repeatedly. After the second day of inoculating Henrietta’s cells into the culture tubes, their laboratory attendant Mary Kubicek found that the tubes containing HeLa cells were overcrowded and had to make subcultures. The amazing part of this story is that Henrietta had never been given credit for her important contribution to cell line culture and she had no knowledge of her contribution till her death on 4th October, 1951. Even her relatives, for a long period, had no awareness of the con-
tributions of HeLa cells to the scientific community and eventually humanity. Only after twenty five years Henrietta's husband, Day Lacks received a call from Dr. Susan Hsu, a geneticist from Baltimore and thus, learnt about the existence of his wife's cells. She, more precisely her cells, participated in many inconceivable scientific experiments and expeditions. Believe it or not she went to space; was in a nuclear explosion; infected with polio by Jonas Salk in search of its vaccine; participated in AIDS study, gene mapping, etc. HeLa cells were the first human cells to be successfully cloned in 1955. After more than half a century of Henrietta's death, more than 50 million metric tons of her cells have been grown and continue to occupy countless laboratories throughout the world. Her contribution to science and humanity is enormous, may be more than any individual human being that ever roam the Earth.

Oh Crab!

One of the most bewildering and common disease that killed billions throughout the ages is cancer. The word cancer is a latin word, a literal translation of which is “crab”. The Father of Medicine, Hippocrates after investigating a particular form of illness gave a Greek name for that disease, karkinos. Aulus Cornelius Celsius literally latinized karkinos and called it cancer. A standard definition of cancer might be “a disease in which a group of abnormal cells grow uncontrollably by disregarding the normal rules of cell division resulting in more or less immortal cells”. Like all other diseases its cure has been pursuit, but in this case, not with definite success. There are treatments that kill cancerous cells and extending the lives of the victims. But a perfect cure has never been found and whether or not it will be found is a million dollar question.

Here is a crazy thought. What if cancer is a good disease? In your head chemical reactions took place and neurons released neurotransmitters from pre-synaptic to post-synaptic cells and asked “A good disease? Are you crazy?” It is most explicable if you are startled and thought that I am crazy, the warning sign is right up there. Allow me, once again to elaborate. Medically speaking, absolutely, cancer is a disease that kills. But, evolutionarily speaking it seems to be an error in a trial of immortality. It seems nature has found a way to attain immortality at the cellular level, I assume that sometimes in the future nature will find a way to attain immortality at an individual level. It might just be nature rolling the dice; a gambling for life.

I hope an addition of a few lines of my conception would not hamper the scientific sentiment of this article. If we think of cells as individual beings, our concept of cancer as nature's way of attaining immortality may be more well-defined. Maybe this story will enlighten you, or at least amused you. One day a fine young cell-lady residing in the antrum of the stomach fortu-
nately discovered the ability to ‘not’ die and then started a cellular gossip. As human (an assemblage of cells) loves gossip, so do the cells. So, the genius cell-lady who had the Nobel prize winning discovery told every cells in her community, in this case the stomach, her trick of evading death. So, all of them decided not to die. Cells not dying is in fact the definition of cancer and voila, we have a stomach cancer. It is a remarkable quality of living organisms to try and survive even when survival means destruction to our environment. Likewise, the cancer cells cannot comply with their surroundings, their improvement causes havoc to their environments. Actually, we may and should not blame them for acting so out of line because even we, human, fail to comprehend the disaster we generated to our environments and yet we claimed to climb highest in the evolutionary tree. Maybe we are not so far along. It may be fortunate for the cells to attain immortality but definitely not for the persons owning the immortal cells. It is rightly a cellular fortune, but an individual’s damnation.

My notion is that cancer is a disease and an improvement. If we could be choosey and eliminate certain key characters of cancerous cells, like their ability to divide incessantly and their abnormal cellular membrane and stick only to their ability to evade apoptotic signals, we may be able to create an organism that neither die nor age. Evolution is often regarded as a non-reversible process, so, if cancer is a step in cellular evolution as I have assumed, there may never be a perfect cure of cancer, forgive me if I am too pessimistic. Of course killing the cells is a different story. If all cancerous cells died, there may be no potential way for the organism to survive, especially in high stage cancer where majority of the body’s cells are cancerous.

**Tail of Life**

The life cycle of most known organisms involve aging or senescence. Taking human as an example, after a certain age, say fifty, symptoms of aging start to appear. This include weakness of the bones and muscles, wrinkles, gray hairs, decrease visual acuity, etc. It is obvious that after sometime, our cells lose their ability to divide, a point called Hayflick limit. The answer why cells reach the Hayflick limit lie in phenotypic blueprint.

Eukaryotes have chromosomes consisting of double stranded linear deoxyribonucleic acid (DNA). However, the end region of the chromosome is a single non-coding tandemly repeated sequence. This region of the chromosome is called the “telomere” and was independently discovered in the 1930s by two scientists, Barbara McClintock and Herman Muller. The human telomeric sequence is 5’-TTAGGG-3’ that vary from 2 to 50 kilo base pairs and a G-tail of 100-250 bases extending from double stranded to the single stranded region. This single stranded G-rich telomere acts with other proteins to form a T-loop structure serving a protection to the chromosome and maintaining genomic stability. Otherwise, the end termini of the chromosome may be recognized as double strand break. The telomeric sequence is added by a particular ribonucleoprotein complex called telomerase, identified by Elizabeth Blackburn and Carol Grieder.

During replication, DNA replicating machinery cannot copy the single stranded region of the telomere, therefore the telomeres shorten with each cell division, by 50-200 bp. Interestingly, telomerase activity is low or absent in most human cells. So, when the telomeres reach a threshold length because of repeated cell division, checkpoint-mediated cellular senescence is initiated, pausing cell division, the Hayflick limit. As a result, telomere length may be considered as a mitotic clock to limit cell proliferation, thus, reflecting the course of aging. It has been widely believed that reactivation of telomerase activity and eventual lengthening of telomere can result in immortalization. This make telomere or telomerase an important research area in humankind quest for immortality. Another school of thought, contrastingly, believed that there is no basis for a general telomere or telomerase involvement in the hypothesis of aging.

Anyhow, inappropriate activation of telom-
erase allows unregulated cell growth, like cancer. In at least 85% of human cancers, telomerase reactivation has been seen and many cancer drugs target telomerase, although it was recently described that another pathway, telomerase-independent alternative lengthening of telomere (ALT) pathway can maintain the length of telomeres. But this does not suggest that the activation of telomerase is all together bad. Insufficiency of telomerase can cause diseases like dyskeratosis congenita, aplastic anemia, and idiopathic pulmonary fibrosis (T3, T10, 54).10,24,25

Conclusion

Different organisms have different life cycles. The manner and duration remain relatively diverse. Some manage to live for over a century, while others persisted only a day or two. The oldest known living thing in the world, a gram-positive bacteria, Actinobacteria is estimated to be ~400,000 years old. Other species with long lasting lives include the great basin bristlecone spine (4,862 years), Antarctic sponge (1,550 years), Bowhead whales (210 years), etc.3 But are they immortals? I’m afraid not. Our interest is not on long lasting lives but everlasting lives, like the beliefs of most religions. But just because these long living organisms can somehow die, we should not simply stop being curious in finding their secrets. At least we have to start somewhere, like the time we learn to build an airplane before we fly to the moon. Our discussion is somehow too short as we have to exclude other interesting biological immortals. But to mention all would be too extended. It may not be very wrong to focus on long living organisms, which are among the simplest life forms called basal metazoans, a group that includes sponges, corals, jellyfish, comb jellies, hydras, and sea anemones in our search for immortality.26,27

At the macroscopic level, nature has a tendency to screw up, an effect called entropy in physics. But, on a microscopic scale, since the amount of energy involved is very small, it is very difficult to know for certainty that the entropy increases, thus, the entropy effect might altogether be negligible.28 This might explain why nature has find a way of immortality at the cellular level but not on a multicellular organismal level, because once we became an organism, nature screwed us to expire. To put a butter to this bread, cell lines, like the HeLa cells can be frozen and thawed without killing them, but our current scientific, technological and ethical advancement cannot yet determine the method to replicate this resurrecting technique on a multicellular organismal scale. Do not get me wrong here. A North American frog Rana sylvatica during hibernation have their body frozen and can live normal lives after being thawed. But in these case, only around 70% of water in their body is frozen.29 A 100% frozen body might present a different story.

I believe the core concept of every religion is to live forever. Some believe in infinite spiritual life after the termination of our physical beings, while others believe in reincarnation. Some religions even believe in the existence of a herb of immortality. Some scholars have identified this herb as Amanita muscaria, a.k.a. Fly Agaric, a.k.a. Maga.30 Our yearning of immortality, echoing through the beliefs of many, if not all religions and cultures throughout ages, may one fine day be accomplished through the advancement of science and technology, or by nature. By then, humans will no longer be humans but a whole new species, perhaps Homo infinitus, capable of evading death and no longer having to procreate (life must be so dull), because procreation after attaining immortality will only lead to population explosion. But this process of speciation may take millions or billions of years, an incomprehensible process to many because of the inability or ignorance to incorporate the time factor involved. This rationale may possibly be that our mind does not evolve to think for period as millions or billions of years. Sadly, we may not have the time to evolve to immortals. Looking at the effects we have imprinted on the only known planet that can sustain life, this anthropocene period may be our last signature.
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