

Fetal Movement of Individual Evolutionary Biology

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Summary: We are in preparing for the birth of a new science, Individual Evolutionary Biology.

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The stem cell is the ultimate medical weapon (Pluchino et al., 2020; Youssef et al., 2016; Pal, 2015). We have studied them for more than a half-century, but we still know only a little. Mainly, we have known there are naturally many stem cells in most tissues in an adult human body (Demarco et al., 2020), some stem cells stay quiescent and some are differentiating (Tumpel and Rudolph, 2019), the human body can potentially repair any tissue injury and regenerate any organs by utilizing the stem cell differentiation, we can reprogram a somatic cell to the original pluripotent state to get unlimited stem cells (Takahashi et al., 2007), we can potentially kill any cancers by utilizing the stem cell quiescence and apoptosis mechanism (Pan et al., 2018), the center of manipulating stem cells is the signals, which consists of a majority of chemical molecules (Tanabe, 2015) with some biophysical signals (Yim, 2012).

We are collecting more and more signals, but the progress is slow that we basically have only one mature stem cell therapy, hematopoietic stem cell transplantation (Tiwari et al., 2020, Jiang and Lian, 2020, Rangatchew et al., 2020, Grochowski et al., 2018,

Jevotovsky et al., 2018), and it is far away from manipulating autologous hematopoietic stem cells. To completely manipulate a stem cell, we need a signal dictionary about stem cells' every molecule, every stage, and every environment, to light the research in every niche.

To speed up stem cell research, we need some new tools and new methods along with the biological experiments. Computational biology is the hottest one among others. Based on big data, we can use the computer to break the material barriers and quickly get the optimal solution which is equivalent to completing hundreds of physical experiments (Bian and Cahan, 2015). One of the sharpest tools for computational biology is the quantum computer. In 2019, Google's quantum computer, with 53 qubits, completed a complex computation in 200 seconds, which would need the most powerful supercomputers 10,000 years to complete (Arute, 2019).

Another powerful tool for computational biology is Artificial Intelligence. The most powerful aspect of Artificial intelligence is its quickly self-evolving, maybe thousands of generations in a single day. Evolution, the central concept of biology, is also the central mission of Artificial intelligence.

With the combination of the quantum computer and artificial intelligence, we can expect much higher productivity in the next 50 years for stem cell science.

We want to stick to the stem cell science for future medicine because it gives us a promise of immortal life. Based on a complete signal knowledge about stem cells, we could freely manipulate all the stem cells in our body, timely repair all tissues, and

regenerate any organs in case of accidents. Theoretically, everyone could stay at the peak of life, around 18 years old, forever.

And the stem cell gives us another an even bigger promise of approaching individual perfect life. We can edit the gene in our stem cells (Trounson et al., 2019), and the new stem cells will make new body cells. In the long run, we will recreate our genes for the whole body to complete the self-evolution. Even Charles Darwin couldn't imagine this prospect because biological evolution always just talk about population evolution, can never talk about individual evolution. The advanced stem cell technology will unprecedentedly bring in individual evolution. This will be a real biological revolution, let's embrace this revolution.

References:

Arute F., Arya K., Babbush R., et al. (2019). Quantum supremacy using a programmable superconducting processor. *Nature* 574, 505–510.
<https://doi.org/10.1038/s41586-019-1666-5>

Bian Q., Cahan P. (2016). Computational tools for stem cell biology. *Trends in Biotechnology*. Volume 34, Issue 12. <https://doi.org/10.1016/j.tibtech.2016.05.010>

Demarco R.S., Clemot M., Jones D.L. (2020). The impact of aging on lipid-mediated regulation of adult stem cell behavior and tissue homeostasis. *Mechanisms of Ageing and Development*. Volume 189. <https://doi.org/10.1016/j.mad.2020.111278>

Grochowski C., Radzikowska E., Maciejewski R. (2018). Neural stem cell therapy—brief review. *Clinical Neurology and Neurosurgery*. Volume 173.
<https://doi.org/10.1016/j.clineuro.2018.07.013>

Jevotovsky D.S., Alfonso A.R., Einhorn T.A., Chiu E.S. (2018). Osteoarthritis and stem cell therapy in humans: a systematic review. *Osteoarthritis and Cartilage*. Volume 26, Issue 6. <https://doi.org/10.1016/j.joca.2018.02.906>

Jiang Y., Lian X.L. (2020). Heart regeneration with human pluripotent stem cells: prospects and challenges. *Bioactive Materials*. Volume 5, Issue 1. <https://doi.org/10.1016/j.bioactmat.2020.01.003>

Pal L. (2015). Uterine stem cells—promise and possibilities. *Maturitas*. Volume 82, Issue 3. <https://doi.org/10.1016/j.maturitas.2015.07.018>

Pan S., Sun Y., Sui D., Yang T., Fu S., Wang J., Hui B., Xi R., He C., Zhang X. (2018). Lobaplatin promotes radiosensitivity, induces apoptosis, attenuates cancer stemness and inhibits proliferation through PI3K/AKT pathway in esophageal squamous cell carcinoma. *Biomedicine & Pharmacotherapy*. Volume 102. <https://doi.org/10.1016/j.biopha.2018.03.109>

Pluchino S., Smith J.A., Peruzzotti-Jametti L. (2020). Promises and limitations of neural stem cell therapies for progressive multiple sclerosis. *Trends in Molecular Medicine*. <https://doi.org/10.1016/j.molmed.2020.04.005>

Rangatchew F., Vester-Glowinski P., Rasmussen B.S., Haastrup E., Munthe-Fog L., Talman M.L., Bonde C., Drzewiecki K.T., Fischer-Nielsen A., Holmgaard R. (2020). Mesenchymal stem cell therapy of acute thermal burns: a systematic review of the effect on inflammation and wound healing. *Burns*. <https://doi.org/10.1016/j.burns.2020.04.012>

Takahashi K., Tanabe K., Ohnuki M., Narita M., Ichisaka T., Tomoda K., Yamanaka S. (2007). Induction of Pluripotent Stem Cells from Adult Human Fibroblasts by Defined Factors". *Cell*. 131 (5): 861–872. <https://doi: 10.1016/j.cell.2007.11.019>

Tanabe S. (2015). Signaling involved in stem cell reprogramming and differentiation. *World J Stem Cells*. 7(7):992-998. <http://doi.org/10.4252/wjsc.v7.i7.992>

Tiwari S., Khan S., Kumar V.S., Rajak R., Sultana A., Pasha S.A., Gauba D., Ghosh P., Khurana T., Kulkarni A., Reddy Y.P., Khan A.A., Sharma V.K. (2020). Efficacy and safety of neural stem cell therapy for spinal cord injury: a systematic literature review. *Therapies*. <https://doi.org/10.1016/j.therap.2020.06.011>

Trounson A., Boyd N.R., Boyd R.L. (2019). Toward a universal solution: editing compatibility into pluripotent stem cells. *Cell Stem Cell*. Volume 24, Issue 4. <https://doi.org/10.1016/j.stem.2019.03.003>

Tumpel S., Rudolph K.L. (2019). Quiescence: good and bad of stem cell aging. *Trends in Cell Biology*. Volume 29, Issue 8. <https://doi.org/10.1016/j.tcb.2019.05.002>

Yim E.K., Sheetz M.P. (2012). Force-dependent cell signaling in stem cell differentiation. *Stem Cell Res Ther* 3, 41. <https://doi.org/10.1186/scrt132>

Youssef A.A., Ross E.G., Bolli R., Pepine C.J., Leeper N.J., Yang P.C. (2016). The promise and challenge of induced pluripotent stem cells for cardiovascular applications. Volume 1, Issue 6. <https://doi.org/10.1016/j.jacbts.2016.06.010>