Stem Cell Introduction

Mark H Smith

Queens, New York 11418, USA mark20082009@gmail.com

Abstract: The definition of stem cell is "an unspecialized cell that gives rise to a specific specialized cell, such as a blood cell". Stem Cell is the original of life. All cells come from stem cells. Serving as a repair system for the living body, the stem cells can divide without limit to replenish other cells as long as the living body is still alive. When a stem cell divides, each new cell has the potential to either remain a stem cell situation or become another type of cell with a more specialized function, such as a muscle cell, a red blood cell, a bone cell, a nerve cell, or a brain cell. Stem cell research is a typical and important topic of life science.

[Smith MH. Stem Cell Introduction. Stem Cell 2012;3(1):250-252] (ISSN 1545-4570). http://www.sciencepub.net/stem. 7

Key words: stem cell; life; gene; DNA; protein

The stem cell is the origin of an organism's life that has the potential to develop into many different types of cells in life bodies. In many tissues stem cells serve as a sort of internal repair system, dividing essentially without limit to replenish other cells as long as the person or animal is still alive. When a stem cell divides, each new cell has the potential either to remain a stem cell or become another type of cell with a more specialized function, such as a red blood cell or a brain cell.

Stem cells have two important characteristics: (1) They are unspecialized cells capable of renewing themselves through cell division, sometimes after long periods of inactivity; (2) Under certain physiologic or experimental conditions, they can be induced to become differentiated adult cells with special functions. In some organs, such as the bone marrow, stem cells regularly divide to repair and replace worn out or damaged tissues. In other organs, however, such as the pancreas and the heart, stem cells only divide under special conditions. Stem cells can be used in the clinical medicine to treat patients with a variety of diseases (Daar, 2003). Serving as a repair system for the living body, the stem cells can divide without limit to replenish other cells as long as the living body is still alive. When a stem cell divides, each new cell has the potential to either remain a stem cell situition or become another type of cell with a more specialized function, such as a muscle cell, a red blood cell, a bone cell, a nerve cell, or a brain cell. Stem cell research is a tipical and important topic of life science.

In 2003, it was found that foetal stem cells had the ability to multiply without limit and never grow old (Hawkes, 2003), which may make it possible to create foetal stem cells from adult cells. The gene Nanog is a regulator that controls the operation of many other genes. It operates only in embryonic stem cells, which are pluripotent. Nanog's role is to maintain stem cells and to make them grow. Nanog is a master gene that may make stem cells immortal.

Embryonic stem cells can go on dividing forever. This means that a culture of stem cells can be kept alive for transplantation into patients where they will diversify into necessary cells. For this to be possible, it needs to know how it is that stem cells can either divide without limit, or choose instead to differentiate into specialised cells. Nanog appears to be the key. Nanog does not disappear in adult cells, but it lies dormant. This means that if a way could be found to reactivate it, adult cells could be persuaded to become embryonic cells again. The next step is to work out how Nanog is switched on and off. To achieve that it may be necessary to continue working on embryonic stem cells and watching the process as it happens.

Many diseases are caused by the death of cells vital to the proper functioning of the organs. Heart failure, for example, is often caused by damage to the muscles caused by a blood clot. Stem cells injected into the heart could recreate the heart muscle.

Type 1 diabetes is caused by the destruction of the pancreatic cells that make insulin. These cells might be reintroduced as stem cells.

Parkinson's disease is caused by a loss of cells. In animal experiments stem cells have been shown to reduce symptoms of the disease.

Some of the most notable findings are as follows: (1) organ-specific stem-cell growth and differentiation are stimulated during the reparative phase following transient injury; (2) some organs contain resident marrow-derived stem cells, and their differentiation potential may only be expressed during repair; (3) the metanephric mesenchyme contains pluripotent and self-renewing stem cells; (4) epithelial-tomesenchymal transition generates renal fibroblasts, etc (Oliver, 2004).

Stem cell researches are developing very fast. As an example in Korea, after the claims of the first cloning of patient-specific stem cells using somatic nuclear transfer as published in Science in 2004 and 2005, former Seoul National University Professor Woo Suk Hwang became a national hero of Korea and an international celebrity. His academic reputation was down after January 11, 2006 when a ninemember investigation panel at Seoul National University reported that his data were intentionally fabricated. In spite of these negative effectives from both scientific and non-scientific communities, stem cell research in Korea is bouncing back with better focus and balance as evidenced below. First, the Korean government has been reassuring by vowing to continue to support stem cell research in Korea with a long-term spending plan of \$454 million over the next 10 years. This represents an even higher level of funding than before the Hwang scandal. Second, nongovernment sectors are also continuing to push their plans to support stem cell science. Third, despite concerns and unfounded worries, Korean biologists are performing exceedingly well and continue to publish their results in prestigious international journals (Kwang-Soo Kim. 2007).

Until recently, scientists primarily worked with two kinds of stem cells from animals and humans: <u>embryonic stem cells</u> and non-embryonic <u>somatic or</u> <u>adult stem cells</u>. In 2006, it was made another breakthrough by identifying conditions that would allow some specialized adult cells to be "reprogrammed" genetically to assume a stem celllike state, which is named <u>induced pluripotent stem</u> <u>cells (iPSCs)</u>.

In the 3- to 5-day-old embryo (<u>blastocyst</u>), the inner cells give rise to the entire body of the organism, including all of the many specialized cell types and organs such as the egg, sperm, skin, heart, lung, etc.

Stem cells offer new potentials for treating diseases such as diabetes, and heart disease. Adult stem cells typically generate the cell types of the tissue in which they reside. For example, a blood-forming adult stem cell in the bone marrow normally gives rise to the many types of blood cells.

References

1. Paul Woodard. SCDHAP Protocol: ematopoietic Stem Cell Transplantation (HSCT) for Patients with Sickle Cell Disease and Prior Stroke or Abnormal Transcranial Doppler Ultrasound (TCD) using Reduced Conditioning and T-Cell-Depleted Hematopoietic Stem Cells from Partially Matched Family Donors - Phase I Study.

http://www.stjude.org/protocols/0,2081,450_232 7_18472,00.html. 2007.

- Renee Madden. SCT521 (COG # ASCT0521) Protocol: Soluble Tumor Necrosis Factor Receptor: Enbrel (Etanercept) for the Treatment of Acute Non-Infectious Pulmonary Dysfunction (Idiopathic Pneumonia Syndrome) Following Allogeneic Stem Cell Transplantation. <u>http://www.stjude.org/protocols/0,2881,450_233</u> <u>3_5873,00.html</u>. 2007.
- 3. Kimberly Kasow. OPBMT2 Protocol: Allogeneic Hematopoietic Stem Cell Transplantation for Children Affected with Malignant Osteopetrosis - A Pilot Study. <u>http://www.stjude.org/protocols/0,2881,450_233</u> <u>1_17072,00.html</u>. 2007.
- Baud L, Haymann JP, Bellocq A, Fouqueray B. Contribution of stem cells to renal repair after ischemia/reperfusion. Bull Acad Natl Med. 2005;189(4):635-43.
- Bernard Lo, Patricia Zettler, Marcelle I. Cedars, Elena Gates, Arnold R. Kriegstein, Michelle Oberman, Renee Reijo Pera, Richard M. Wagner, Mary T. Wuerth, Leslie E. Wolf, Keith R. Yamamoto. A New Era in the Ethics of Human Embryonic Stem Cell Research. Stem Cells. <u>http://www.StemCells.com. http://stemcells.alphamedpress.org/cgi/reprint/20</u>05-0324v1.pdf. 2005.
- 6. Cantley LG. Adult stem cells in the repair of the injured renal tubule. Nat Clin Pract Nephrol. 2005;1(1):22-32.
- 7. Duffield JS, Bonventre JV. Kidney tubular epithelium is restored without replacement with bone marrow-derived cells during repair after ischemic injury. Kidney Int. 2005;68(5):1956-61.
- Duffield JS, Park KM, Hsiao LL, Kelley VR, Scadden DT, Ichimura T, Bonventre JV. Restoration of tubular epithelial cells during repair of the postischemic kidney occurs independently of bone marrow-derived stem cells. J Clin Invest. 2005;115(7):1743-55.
- Herrera MB, Bussolati B, Bruno S, Fonsato V, Romanazzi GM, Camussi G. Mesenchymal stem cells contribute to the renal repair of acute tubular epithelial injury. Int J Mol Med. 2004;14(6):1035-41.
- 10. Hishikawa K, Fujita T. Stem cells and kidney disease. Hypertens Res. 2006;29(10):745-9.
- 11. Humphreys BD, Duffield JD, Bonventre JV. Renal stem cells in recovery from acute kidney injury. Minerva Urol Nefrol. 2006;58(1):13-21.
- 12. Kwang-Soo Kim. Stem cell research continues in Korea beyond the Hwang scandal. Stem Cells. <u>http://www.StemCells.com.</u> <u>http://stemcells.alphamedpress.org/cgi/reprint/20</u> <u>07-0089v1.pdf</u>. 2007.

- Lin F, Cordes K, Li L, Hood L, Couser WG, Shankland SJ, Igarashi P. Hematopoietic stem cells contribute to the regeneration of renal tubules after renal ischemia-reperfusion injury in mice. J Am Soc Nephrol. 2003;14(5):1188-99.
- Lin F. Stem cells in kidney regeneration following acute renal injury. Pediatr Res. 2006;59(4 Pt 2):74R-8R.
- 15. Morigi M, Benigni A, Remuzzi G, Imberti B. The regenerative potential of stem cells in acute renal failure. Cell Transplant. 2006;15 Suppl 1:S111-7.
- 16. Oliver JA. Adult renal stem cells and renal repair. Curr Opin Nephrol Hypertens. 2004;13(1):17-22.
- Yamashita S, Maeshima A, Nojima Y. Involvement of renal progenitor tubular cells in epithelial-to-mesenchymal transition in fibrotic rat kidneys. J Am Soc Nephrol. 2005;16(7):2044-51.
- Wing Leung. INFT2 Protocol: HLA -Nonidentical Stem Cell and Natural Killer Cell Transplantation for Children Less than 2 Years of Age with Hematologic Malignancies. <u>http://www.stjude.org/protocols/0,2881,450_233</u> 0_11129,00.html. 2007.
- 19. Bavister BD, Wolf DP, Brenner CA. Challenges of primate embryonic stem cell research. Cloning Stem Cells 2005;7(2):82-94.
- Bhatt RI, Brown MD, Hart CA, Gilmore P, Ramani VAC, George NJ, Clarke NW. Novel method for the isolation and characterisation of the putative prostatic stem cell. Cytometry A. 2003;54(2):89-99.
- Condorelli G, Peschle C. Stem cells for cardiac repair: state of the art. Front Biosci 2005;10:3143-50.
- 22. Daar AS, Sheremeta L. The science of stem cells: ethical, legal and social issues. Exp Clin Transplant. 2003;1(2):139-46.

- 23. Kashofer K, Bonnet D. Gene Therapy Progress and Prospects: Stem cell plasticity. Gene Ther. 2005 (Epub ahead of print).
- 24. Ma H. Technique of Animal Clone. Nature and Science 2004;2(1):29-35.
- 25. Stedman's Medical Dictionary. The American Heritage®. Houghton Mifflin Company. <u>http://dictionary.reference.com/search?q=stem%</u> 20cell. 2002.
- 26. Williams D. Stem cells in medical technology. Med Device Technol 2005;16(3):9-11.
- 27. Nigel Hawkes. Scientists find the secret of eternal life for stem cells. http://www.timesonline.co.uk/tol/news/uk/article 1137674.ece
- 28. <u>http://stemcells.nih.gov/staticresources/research/</u> protocols/BresaGen_hESC_manual_2.1.pdf
- 29. Mitalipova et al. Stem Cells. 2003;21(5):521-6.
- Ma H, Chen G (2005). Stem Cell. J Am Sci. 1(2):90-92. <u>http://www.sciencepub.net/american/0102/14-</u> mahongbao.pdf.
- 31. Ma H, Chenrg S (2007). Eternal Life and Stem Cell. Nat Sci 5(1):81-96. <u>http://www.sciencepub.net/nature/0501/10-0247-</u> mahongbao-eternal-ns.pdf.
- 32. Ma H, Chenrg S (2007). Review of Stem Cell Studies. Nat Sci 5(2):45-65. <u>http://www.sciencepub.net/nature/0502/09-0247-</u> mahongbao-stem-ns.pdf.
- 33. Yang Y, Ma H (2010). Germ Stem Cell. Stem Cell. 1(2):38-60].
 <u>http://www.sciencepub.net/stem/stem0102/07_1</u>
 348stem0102_38_60.pdf.
- 34. Pubmed. Stem Cell. <u>http://www.ncbi.nlm.nih.gov/pubmed/?term=ste</u> <u>m+cell</u>.
- 35. Wikipedia. Stem Cell. http://en.wikipedia.org/wiki/Stem_cell.

10/11/2011