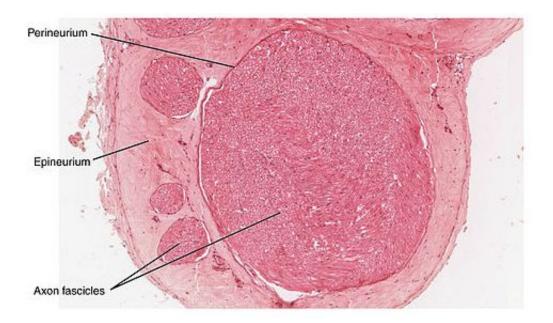


New way to repair nerves: Using exosomes to hijack cell-to-cell communication

September 15 2015



Cross-section of a nerve. Credit: OpenStax College/Wikipedia

Regenerative medicine using stem cells is an increasingly promising approach to treat many types of injury. Transplanted stem cells can differentiate into just about any other kind of cell, including neurons to potentially reconnect a severed spinal cord and repair paralysis.

A variety of agents have been shown to induce transplanted <u>stem cells</u> to differentiate into neurons. Tufts University biomedical engineers recently published the first report of a promising new way to induce



human mesenchymal stem cells (or hMSCs, which are derived from bone marrow) to differentiate into neuron-like cells: treating them with exosomes.

Exosomes are very small, hollow particles that are secreted from many types of cells. They contain functional proteins and genetic materials and serve as a vehicle for communication between cells. In the nervous system, exosomes guide the direction of <u>nerve</u> growth, control nerve connection and help regenerate peripheral nerves.

In a series of experiments reported in *PLOS ONE* in August, the Tufts researchers showed that exosomes from PC12 cells (neuron-like progenitor cells derived from rats) at various stages of their own differentiation could, in turn, cause hMSCs to become neuron-like cells. Exosomes had not previously been studied as a way to induce human stem cell differentiation.

The biomedical engineers also showed that the exosomes contain miRNAs—tiny pieces of RNA that regulate cell behavior and are known to play a role in neuronal differentiation. The researchers hypothesize that the exosomes caused the hMSCs to differentiate by delivering miRNA into the stem cells. The researchers plan future studies to determine the exact mechanism.

Synthetic Exosomes Could Avoid Need for Neural Progenitor Cells

"In combination with synthetic nanoparticles that my laboratory is developing, we may ultimately be able to use these identified miRNAs or proteins to make synthetic exosomes, thereby avoiding the need to use any kind of neural progenitor cell line to induce neuron growth," said the paper's senior and corresponding author Qiaobing Xu, assistant professor of biomedical engineering at Tufts School of Engineering.



Xu's work focuses on material science engineering, specifically nanoscience and its biomedical application: the development of new synthetic materials for the delivery of therapeutic proteins and genetic material. In 2015, he received a Faculty Early Career Development (CAREER) award from the National Science Foundation (NSF), funding research into a new way to deliver protein-based cancer-fighting drugs and other therapeutics directly into cells. He was named a Pew Scholar in biomedical sciences by the Pew Charitable Trusts in 2013. In addition to the School of Engineering, he has appointments in the School of Medicine and Sackler School of Graduate Biomedical Sciences at Tufts University.

More information: Takeda YS, Xu Q (2015) Neuronal Differentiation of Human Mesenchymal Stem Cells Using Exosomes Derived from Differentiating Neuronal Cells. *PLOS ONE* 10(8): e0135111. <u>DOI:</u> <u>10.1371/journal.pone.0135111</u>

Provided by Tufts University

Citation: New way to repair nerves: Using exosomes to hijack cell-to-cell communication (2015, September 15) retrieved 23 July 2023 from <u>https://medicalxpress.com/news/2015-09-nerves-exosomes-hijack-cell-to-cell.html</u>

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