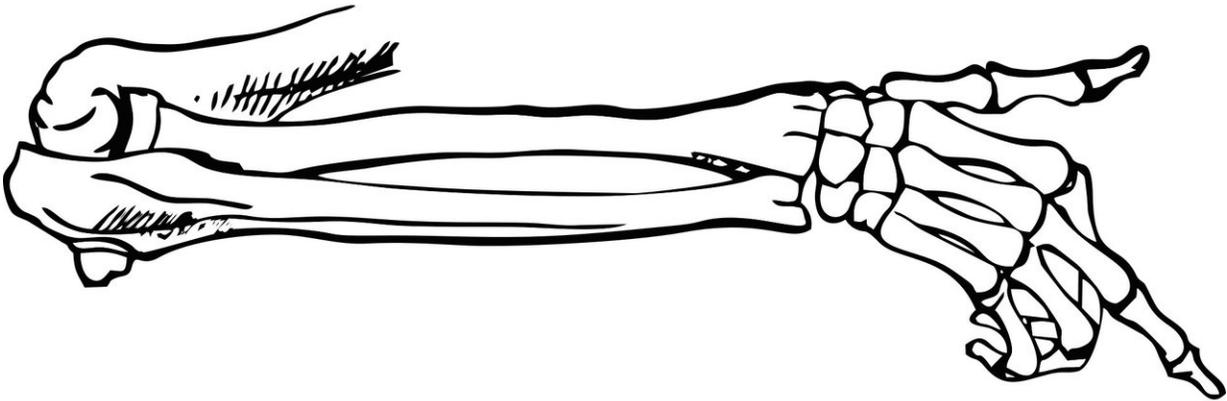


New mechanism of bone growth discovered

February 27 2019



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In a paper published in the journal *Nature*, an international research team led by researchers at Karolinska Institutet in Sweden reports that bone growth in mice takes place in accordance with the same principles as when new cells are produced in blood, skin and other tissue. This contradicts the previous understanding that bone growth depends on a finite number of gradually consumed progenitor cells. If the new findings also apply to humans, they could make an important contribution to the treatment of children with growth disorders.

The growth of children's bones depends on growth plates (physes) situated close to the ends of all long bones in the body. These plates consist of [cartilage cells](#), chondrocytes, that form a kind of scaffold supporting the formation of new bone tissue, and that are themselves generated from stem-cell like [progenitor cells](#) called chondroprogenitors.

For long bones to grow properly, chondrocytes must be generated constantly throughout the growth period. The general view in the field has been that there is a limited number of progenitor [cells](#) that are formed during [embryonic development](#) and then consumed for [bone growth](#) until they run out and we stop growing. In an attempt to ascertain whether or not this is true, researchers at Karolinska Institutet decided to study the formation of chondrocytes in mice.

"What we found was that small 'clones' of cells were generated from the same progenitor cells during embryonic development, which is in line with the current view," says research group leader Andrei Chagin, docent at the Department of Physiology and Pharmacology, Karolinska Institutet. "But after birth, there were dramatic changes in cell dynamics and large, stable clones were formed that proved to be a consequence of how the chondroprogenitors had acquired the ability to regenerate."

Such progenitor cell behaviour is typical for tissue that constantly produces new cells, such as skin, blood and intestinal tissue. For such tissue types, it has been shown that the progenitor cells are situated in a very specific micro-environment, a stem cell niche, which generates the necessary cells (e.g. skin and blood cells) but also enables the progenitor cells to renew themselves. If the niche is disrupted or dysfunctional, the progenitor cells become depleted and the tissue is damaged.

The researchers have now shown that there is a stem cell niche in growth plates, too, at least in mice, and that bone growth ceases if this local micro-environment is disrupted, implying that bone growth follows a

completely different principle from what scientists previously believed.

"If it turns out that humans also have this [growth mechanism](#), it could lead to a significant reassessment of numerous therapeutic approaches used for children with growth disorders," says Dr. Chagin. "The mechanism could also explain some previously puzzling phenomena, such as the unlimited growth seen in patients with certain genetic mutations."

More information: A radical switch in clonality reveals a stem cell niche in the epiphyseal growth plate. *Nature*, 27 February 2019. [DOI: 10.1038/s41586-019-0989-6](https://doi.org/10.1038/s41586-019-0989-6)

Provided by Karolinska Institutet

Citation: New mechanism of bone growth discovered (2019, February 27) retrieved 16 April 2024 from <https://medicalxpress.com/news/2019-02-mechanism-bone-growth.html>

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