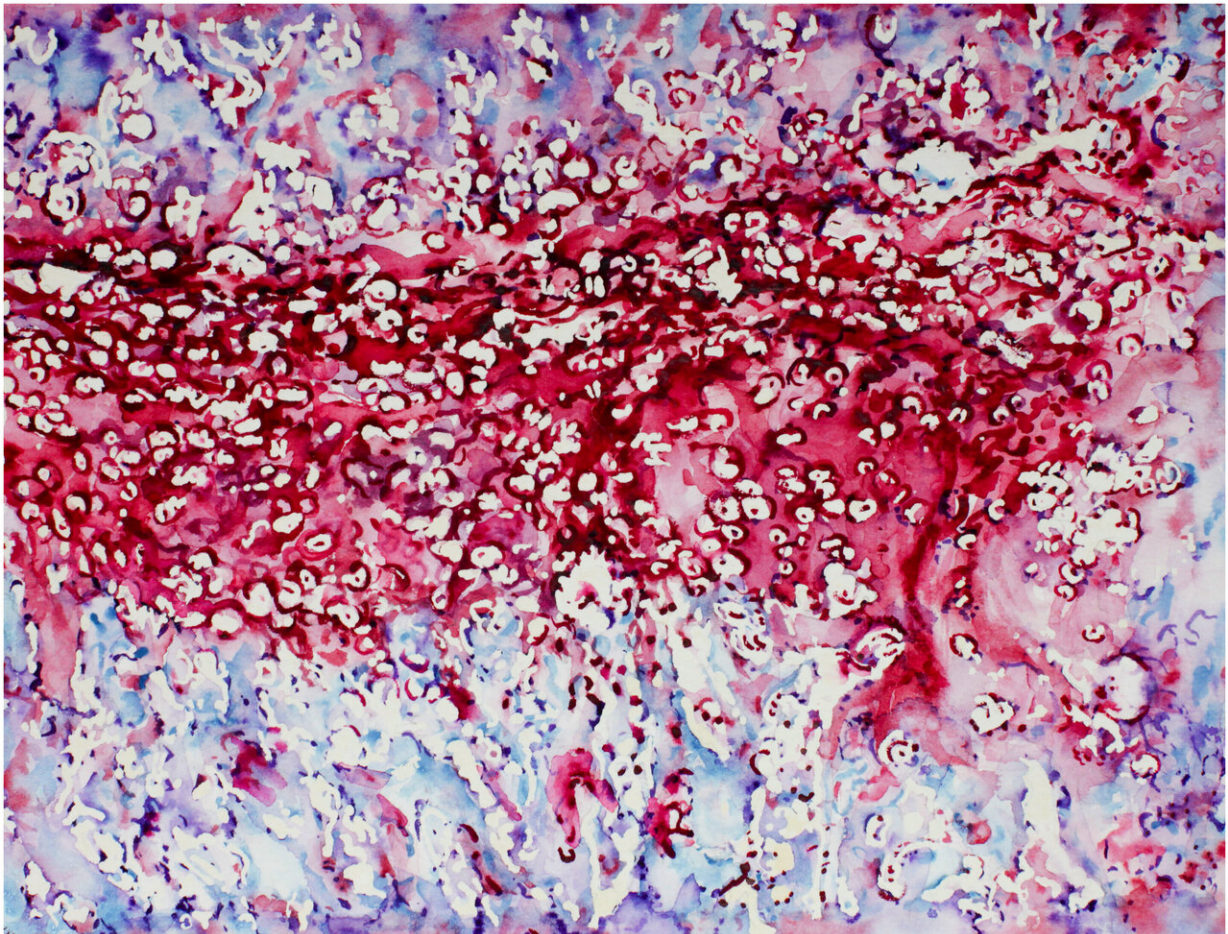


# Addition of growth factors to unique system helps new bone formation

August 28 2019

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Watercolor painting of endochondral ossification featuring both cartilage and bone. Credit: Joel Boerckel

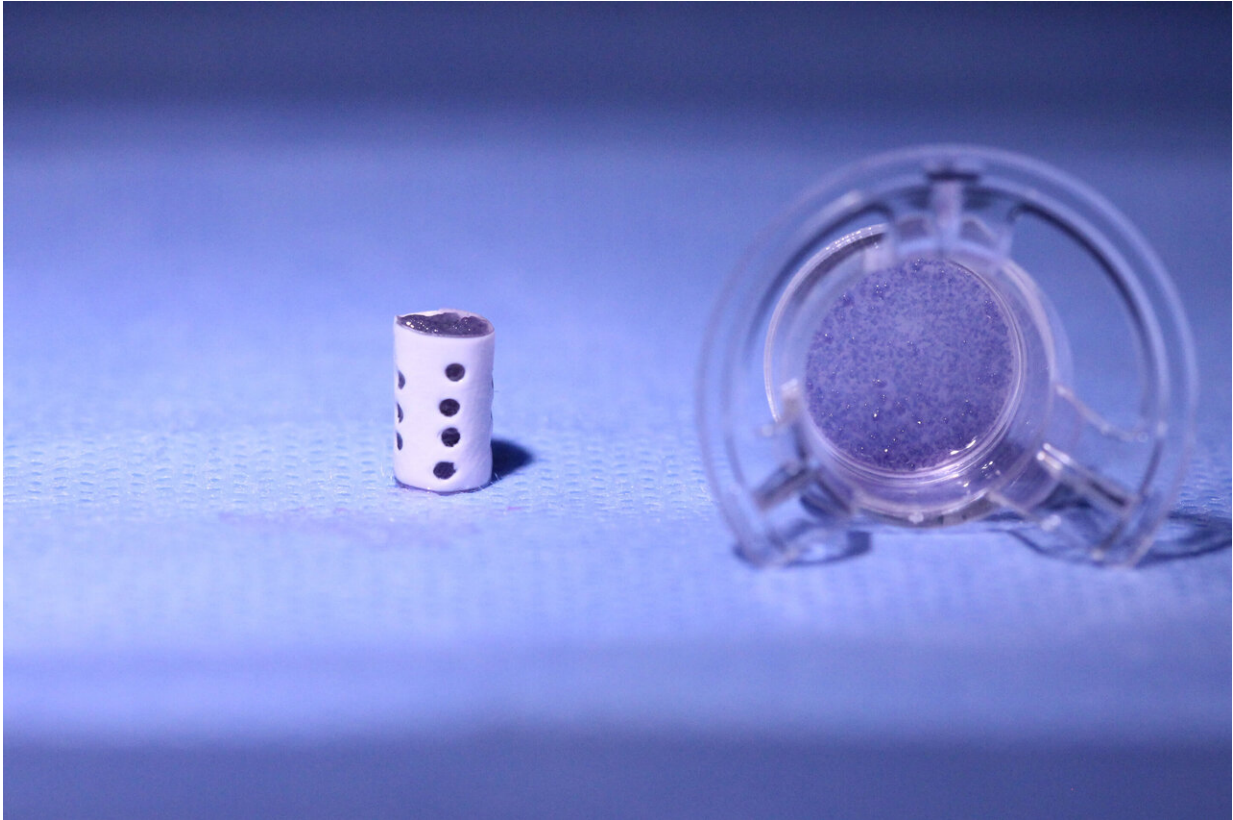
The development of new bone can be a multistep process: first, stem cells differentiate into cartilage cells. Next, the cartilage cells become bone cells. But that's not all: the cells must experience some mechanical stresses during the transformation in order to transform efficiently from stem cells to bone cells.

Researchers at the University of Illinois at Chicago, in collaboration with colleagues at the University of Pennsylvania and Case Western University, have been working on a unique system that delivers stem cells to sites of [bone](#) defects and uses flexible fixators—pins and rods that are used to hold bones together—to provide tunable amounts of mechanical stress.

Now, these researchers, led by Eben Alsberg, the Richard and Loan Hill Professor of Bioengineering and Orthopaedics in the UIC College of Engineering, and Joel Boerckel, assistant professor of orthopaedic surgery at the University of Pennsylvania, have come up with a way to provide timed release of two growth factors that mimic the bone formation process that occurs during embryonic bone development in a rat model.

Together, the stem cells, flexible fixators and timed release of growth factors could one day help heal large bone defects or breaks in people. Their results are published in the journal *Science Advances*.

"What we are doing, in effect, is delivering the right growth factors at the right time to encourage the development of bone from stem cells in the same way that it happens during natural bone healing and development," said Alsberg, who is a corresponding author on the paper along with Boerckel.



Images of mesenchymal condensations. Credit: Joel Boerckel

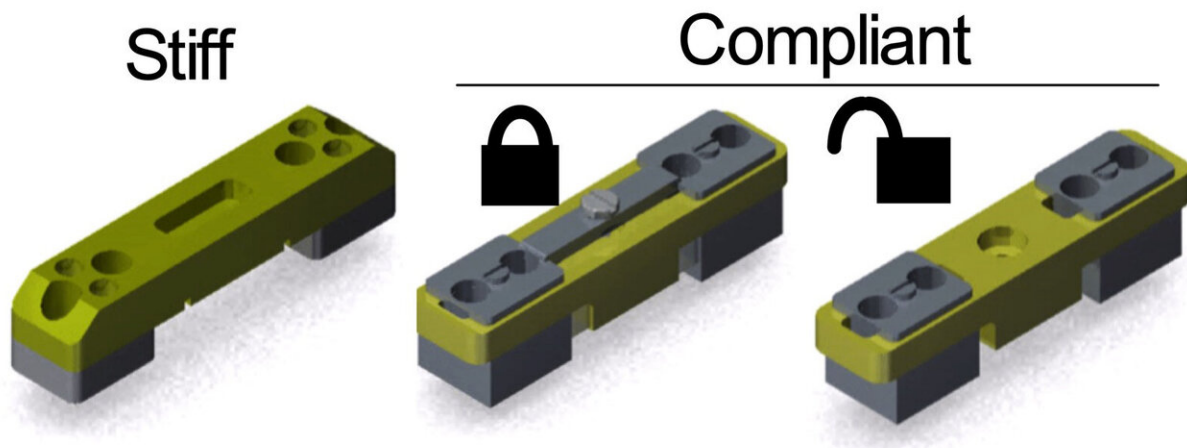
In previous work, Alsberg's group at UIC and colleagues at the University of Pennsylvania developed the flexible fixators and stem cell "condensates," which are masses of stem cells connected to one another so that they can be moved as sheets or plugs. The condensates allow the stem cells to be placed in specific areas of the body, such as within bone gaps or defects without the risk of floating away, as is often the case when stem [cells](#) are delivered to sites in the body through an injection of cell-containing liquids. When used together, the condensates and flexible fixators allowed for enhanced healing of bone defects in a rat model.

In the current research, Alsberg and colleagues added another layer to



their system: they incorporated multiple growth factors into the condensates—one that helps transform [stem cells](#) into [cartilage cells](#) called transforming growth factor beta1 or TGF-beta1, and another that promotes the transition of this cartilage into bone called bone morphogenic protein 2 or BMP-2.

In rats with bone gaps in their femurs, the application of Alsberg's system, including the growth factors, helps encourage the growth of new bone with enhanced function at 12 weeks compared with rats where [growth factors](#) weren't included or only a single growth factor was included in the condensate/flexible fixator system.



Computer models of fixation plates for mechanical load transfer. Credit: Joel Boerckel

"The bone formation achieved was comparable to that which occurs when BMP-2 is soaked in a collagen sponge and applied to bone breaks, which is currently an FDA-approved tissue engineering product for

spinal fusion," Alsberg said. "The collagen product can result in bone-forming where it's not wanted, but in our system, bone only formed in the areas where we placed the growth factor-infused condensates."

**More information:** S. Herberg et al., "Combinatorial morphogenetic and mechanical cues to mimic bone development for defect repair," *Science Advances* (2019). DOI: [10.1126/sciadv.aax2476](https://doi.org/10.1126/sciadv.aax2476) , [advances.sciencemag.org/content/5/8/eaax2476](https://advances.sciencemag.org/content/5/8/eaax2476)

Provided by University of Illinois at Chicago

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