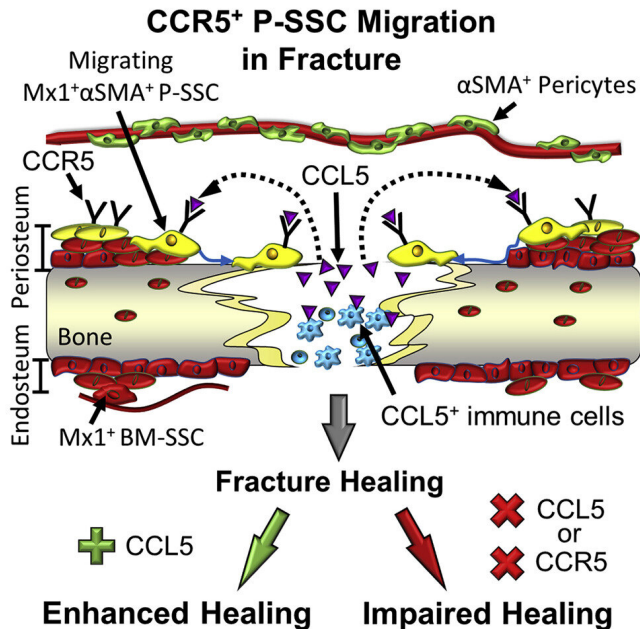


# New bone healing mechanism has potential therapeutic applications

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Periosteal bone stem cells migration and repair mechanism. Credit: Cell Stem Cell/the Park lab

Led by researchers at Baylor College of Medicine, a study published in the journal *Cell Stem Cell* reveals a new mechanism that contributes to adult bone maintenance and repair and opens the possibility of developing therapeutic strategies for improving bone healing.

"Adult bone repair relies on the activation of bone stem cells, which still remain poorly characterized," said corresponding author Dr. Dongsu Park, assistant professor of molecular and human genetics and of pathology and immunology at Baylor. "Bone stem cells have been found both in the [bone marrow](#) inside the bone and also in the periosteum—the outer layer of tissue—that envelopes the bone. Previous studies have shown that these two populations of stem cells, although they share many characteristics, also have unique

functions and specific regulatory mechanisms."

Of the two, periosteal stem cells are the least understood. It is known that they comprise a heterogeneous population of cells that can contribute to bone thickness, shaping and fracture repair, but scientists had not been able to distinguish between different subtypes of bone stem cells to study how their different functions are regulated.

In the current study, Park and his colleagues developed a method to identify different subpopulations of periosteal stem cells, define their contribution to bone fracture repair in live mouse models and identify specific factors that regulate their migration and proliferation under physiological conditions.

## Periosteal stem cells are major contributors to bone healing

The researchers discovered specific markers for periosteal stem cells in mouse models. The markers identified a distinct subset of stem cells that contributes to life-long adult bone regeneration.

"We also found that periosteal stem cells respond to mechanical injury by engaging in bone [healing](#)," Park said. "They are important for healing bone fractures in the adult mice and, interestingly, their contribution to bone regeneration is higher than that of bone marrow stem cells."

In addition, the researchers found that periosteal stem cells also respond to inflammatory molecules called chemokines, which are usually produced during bone injury. In particular, they responded to chemokine CCL5.

Periosteal stem cells have receptors—molecules on their cell surface—that bind to CCL5, which sends a signal to the cells to migrate toward the injured bone and repair it. Deleting the CCL5 gene in

mouse models resulted in marked defects in bone [repair](#) or delayed healing. When the researchers supplied CCL5 to CCL5-deficient mice, bone healing was accelerated.

The findings suggested potential therapeutic applications. For instance, in individuals with diabetes or osteoporosis in which bone healing is slow and may lead to other complications resulting from limited mobility, accelerating [bone healing](#) may reduce hospital stay and improve prognosis.

"Our findings contribute to a better understanding of how adult bones heal. We think this is one of the first studies to show that bone stem [cells](#) are heterogeneous and that different subtypes have unique properties regulated by specific mechanisms," Park said. "We have identified markers that enable us to tell bone stem cell subtypes apart and studied what each subtype contributes to bone health. Understanding how bone stem cell functions are regulated offers the possibility to develop novel therapeutic strategies to treat adult [bone](#) injuries."

**More information:** Laura C. Ortinau et al, Identification of Functionally Distinct Mx1+?SMA+ Periosteal Skeletal Stem Cells, *Cell Stem Cell* (2019). [DOI: 10.1016/j.stem.2019.11.003](https://doi.org/10.1016/j.stem.2019.11.003)

Provided by Baylor College of Medicine

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