

The FDA has approved ankle replacements, so why don't all insurance plans cover them?

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It's been a decade since the U.S. Food and Drug Administration approved the first total ankle-replacement system for patients with severe ankle arthritis.

But several insurance companies still deny coverage, Loyola University Health System orthopaedic surgeon Dr. Michael Pinzur writes in a FootForum commentary in *Foot & Ankle International*, the official journal of the American Orthopaedic Foot and Ankle Society.

"It seems curious that the FDA agrees with the [foot and ankle society] that total ankle replacement is a reasonable treatment option . . . while several insurance providers do not find ankle replacement as a reasonable treatment option for ankle arthritis," Pinzur writes in the June issue.

An ankle replacement is an option for certain patients who suffer severe osteoarthritis, rheumatoid arthritis or injury-related arthritis that does not respond to more conservative treatments. In such patients, arthritis has destroyed cartilage, so the ankle joint is bone-on-bone.

An ankle replacement is similar to a knee or hip replacement. An implant is attached to the bottom of the tibia (shinbone) and to the talus (the first large bone of the foot). The smooth plastic surface of the tibial implant rotates on the polished metal surface of the talar implant.

Since approving the first total ankle-replacement system in 1999, the



FDA has approved two other systems and given tentative approval to a third system. In 2003, the American Orthopaedic Foot and Ankle Society issued a statement that said a total ankle replacement "is a viable option for the treatment of ankle arthritis." And Medicare routinely covers ankle replacements.

Nevertheless, several insurance companies still deny coverage. They base their decision on a "meta-analysis" that concluded an ankle replacement was not a preferred treatment option. The meta-analysis compiled data from previous studies. It was sponsored by insurance companies and based on studies published in 2002 or earlier, Pinzur wrote.

"Should <u>insurance companies</u> make decisions on what treatments are appropriate and what treatments are deemed experimental?" Pinzur's commentary asks.

Source: Loyola University Health System (<u>news</u>: <u>web</u>)

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