

K232592 CITRELOCK® DUOSep 20, 2023
26 days to decisionK232592 · Product code: **MBI** · Orthopedic
Source: <https://www.510kdatabase.net/k232592/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Aug 25, 2023
Decision date	Sep 20, 2023
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Acuitive Technologies, Inc.
Location	Alendale, NJ, US
Contact	Matthew Poggie
510(k) history	6 submissions · 6 cleared · 2021-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232592/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 25, 2026