FDA 510(k) Premarket Notification

K232592 CITRELOCK® DUO

CLEARED

Sep 20, 2023
26 days to decision

K232592 · 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