FDA 510(k) Premarket Notification

K231278 Knotless Suture Anchor

CLEARED

Aug 1, 2023
90 days to decision

K231278 · Product code: **MBI** · Orthopedic
Source: https://www.510kdatabase.net/k231278/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	May 3, 2023
Decision date	Aug 1, 2023
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Riverpoint Medical, LLC**
Location	Portland, OR, US
Contact	Bianca Silva de Sousa
510(k) history	11 submissions · 11 cleared · 2020-2026