

K231278 Knotless Suture AnchorAug 1, 2023
90 days to decisionK231278 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k231278/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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