FDA 510(k) Premarket Notification

K240090 Argo Knotless GENESYS Anchor

CLEARED

Mar 5, 2024
53 days to decision

K240090 · Product code: **MAI** · Orthopedic
Source: https://www.510kdatabase.net/k240090/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Biodegradable, Soft Tissue (MAI)
Date received	Jan 12, 2024
Decision date	Mar 5, 2024
Days to decision	53 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Conmed Corporation**
Location	Utica, NY, US
Contact	Dionne Sanders
510(k) history	82 submissions · 82 cleared · 2004-2026