

K240090 Argo Knotless GENESYS AnchorMar 5, 2024
53 days to decisionK240090 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k240090/> 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 13, 2026