FDA 510(k) Premarket Notification

CLEARED

K190333 Paradigm All-Suture Anchor 1.8mm, White/Black #2

Mar 18, 2019
32 days to decision

K190333 · Product code: **MBI** · Orthopedic
Source: https://www.510kdatabase.net/k190333/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Feb 14, 2019
Decision date	Mar 18, 2019
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k190333/ 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 16, 2026