

K171592 CrossFT™ Knotless Biocomposite Suture Anchor with Disposable DriverAug 29, 2017
90 days to decisionK171592 · Product code: **MAI** · Orthopedic
Source: <https://www.510kdatabase.net/k171592/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Biodegradable, Soft Tissue (MAI)
Date received	May 31, 2017
Decision date	Aug 29, 2017
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Conmed Corporation
Location	Utica, NY, US
Contact	Diana L. Nader-Martone
510(k) history	82 submissions · 82 cleared · 2004-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171592/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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