FDA 510(k) Premarket Notification

K024060 MODIFICATION TO ASNIS III CANNULATED SCREW SYSTEM

CLEARED

Dec 20, 2002
11 days to decision

K024060 · Product code: **HWC** · Orthopedic
Source: https://www.510kdatabase.net/k024060/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Screw, Fixation, Bone (HWC)
Date received	Dec 9, 2002
Decision date	Dec 20, 2002
Days to decision	11 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Howmedica Osteonics Corp.**
Location	Allendale, NJ, US
Contact	KAREN ARIEMMA
510(k) history	288 submissions · 288 cleared · 1999-2020