FDA 510(k) Premarket Notification

K120802 ORTHOLOC(TM) 3DI LOCKING SCREWS ORTHOLOC(TM) BONE SCREWS

CLEARED

Apr 2, 2012
17 days to decision

K120802 · Product code: **HRS** · Orthopedic
Source: https://www.510kdatabase.net/k120802/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Plate, Fixation, Bone (HRS)
Date received	Mar 16, 2012
Decision date	Apr 2, 2012
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Wrightmedicaltechnologyinc**
Location	Arlington, TN, US
Contact	SARAH HOLTGREWE
510(k) history	302 submissions · 291 cleared · 1993-2023