FDA 510(k) Premarket Notification

K102429 ORTHOLOC 3DI ANKLE PLATING SYSTEM, ORTHOLOC BONE SCREW

CLEARED

Nov 23, 2010
90 days to decision

K102429 · Product code: **HRS** · Orthopedic
Source: https://www.510kdatabase.net/k102429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Aug 25, 2010
Decision date	Nov 23, 2010
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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