

**K163650 ORTHOLOC® 3Di Ankle Fusion Plating System Line
Extension**Apr 24, 2017
122 days to decisionK163650 · Product code: **HRS** · Orthopedic
Source: <https://www.510kdatabase.net/k163650/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Dec 23, 2016
Decision date	Apr 24, 2017
Days to decision	122 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k163650/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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