

**K120802 ORTHOLOC(TM) 3DI LOCKING SCREWS
ORTHOLOC(TM) BONE SCREWS**Apr 2, 2012
17 days to decisionK120802 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120802/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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