

**K102429 ORTHOLOC 3DI ANKLE PLATING SYSTEM,  
ORTHOLOC BONE SCREW**

Nov 23, 2010  
90 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k102429/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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