

**K133939 MODIFIED DISTAL RADIUS PLATING SYSTEM**

Apr 1, 2014  
99 days to decision

K133939 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k133939/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Dec 23, 2013
Decision date	Apr 1, 2014
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomet</b>
Location	Warsaw, IN, US
Contact	SUZANA OTANO
Website	<a href="http://www.biomet.com/">http://www.biomet.com/</a>
510(k) history	2 submissions · 2 cleared · 2013-2014

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

Device record: <https://www.510kdatabase.net/k133939/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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