

**K123203 MAXLOCK EXTREME SYSTEM MODEL MXL**Dec 14, 2012  
63 days to decisionK123203 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k123203/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Oct 12, 2012
Decision date	Dec 14, 2012
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Orthohelix Surgical Designs, Inc.</b>
Location	Cleveland, OH, US
Contact	DEREK LEWIS
510(k) history	25 submissions · 25 cleared · 2005-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123203/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 17, 2026