

**K121702 ORTHOCOR ACTIVE DEVICE ORTHOCOR GENERAL  
USE ACTIVE DEVICE**May 6, 2013  
332 days to decisionK121702 · Product code: **ILX** · Physical Medicine  
Source: <https://www.510kdatabase.net/k121702/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Diathermy, Shortwave, For Use Other Than Applying Therapeutic Deep Heat (ILX)
Date received	Jun 8, 2012
Decision date	May 6, 2013
Days to decision	332 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Orthocor Medical</b>
Location	Mccordsville, IN, US
Contact	DAVID ROTHKOPF
Website	<a href="http://www.orthocormedical.com">http://www.orthocormedical.com</a>
510(k) history	3 submissions · 3 cleared · 2009-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121702/> 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 June 9, 2026