

**K141557 HERNIATOME PERCUTANEOUS DISECTOMY DEVICE  
- CERVICAL, HERNIATOME PERCUTANEOUS DISECTOMY  
DEVICE - LUMBAR**Dec 8, 2014  
179 days to decisionK141557 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k141557/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jun 12, 2014
Decision date	Dec 8, 2014
Days to decision	179 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gallini Medical Devices, Srl</b>
Location	South Euclid, OH, US
Contact	ARTHUR S GODDARD
510(k) history	1 submissions · 1 cleared · 2014-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141557/> 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