

**K110739 ENTELLUS MEDICAL SINUS GUIDEWIRE**Jun 14, 2011  
89 days to decisionK110739 · Product code: **LRC** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k110739/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ent Manual Surgical (LRC)
Date received	Mar 17, 2011
Decision date	Jun 14, 2011
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Entellus Medical, Inc.</b>
Location	Maple Grove, MN, US
Contact	KAREN E PETERSON
510(k) history	27 submissions · 27 cleared · 2008-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110739/> 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 24, 2026