

**K130503 PATHASSIST LED LIGHT FIBER**Jun 11, 2013  
104 days to decisionK130503 · Product code: **LRC** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k130503/>**SUBMISSION DETAILS**

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|                       |                                       |
|-----------------------|---------------------------------------|
| Decision              | Substantially Equivalent (Cleared)    |
| Submission type       | Traditional                           |
| Device classification | Instrument, Ent Manual Surgical (LRC) |
| Date received         | Feb 27, 2013                          |
| Decision date         | Jun 11, 2013                          |
| Days to decision      | 104 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/k130503/> 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