

**K023004 LFRE INDIRECT LARYNGOSCOPE**Jan 28, 2003  
141 days to decisionK023004 · Product code: **EOB** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k023004/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)          |
| Submission type       | Traditional                                 |
| Device classification | Nasopharyngoscope (flexible Or Rigid) (EOB) |
| Date received         | Sep 9, 2002                                 |
| Decision date         | Jan 28, 2003                                |
| Days to decision      | 141 days                                    |
| Third-party review    | No  |
| Summary / Statement   | Summary                                     |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Astralite Corp.</b>                |
| Location       | Diamond Springs, CA, US               |
| Contact        | DORIAN SWARTZ                         |
| 510(k) history | 2 submissions · 2 cleared · 1989-2003 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023004/> 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 July 5, 2026