

**K010175 MICROLIGHT 830 LASER SYSTEM**Feb 6, 2002  
384 days to decisionK010175 · Product code: **NHN** · Physical MedicineSource: <https://www.510kdatabase.net/k010175/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)  |
| Submission type       | Traditional   |
| Device classification | Powered Light Based Laser Non-thermal Instrument With Non-heating Effect For Adjunctive Use In Pain Therapy (NHN) |
| Date received         | Jan 18, 2001  |
| Decision date         | Feb 6, 2002   |
| Days to decision      | 384 days  |
| Third-party review    | No  |
| Summary / Statement   | Summary   |

**APPLICANT**

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|----------------|--|
| Company        | <b>Microlight Corporation of America</b> |
| Location       | Missouri City, TX, US                    |
| Contact        | MICHAEL BARBOUR                          |
| 510(k) history | 3 submissions · 3 cleared · 2002-2008    |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010175/> 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 24, 2026