

**K020657 ACCULASER PRO LOW LEVEL LASER THERAPY  
DEVICE**Jul 29, 2002  
151 days to decisionK020657 · Product code: **NHN** · Physical Medicine  
Source: <https://www.510kdatabase.net/k020657/>**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	Feb 28, 2002
Decision date	Jul 29, 2002
Days to decision	151 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Acculaser, Inc.</b>
Location	Washington, DC, US
Contact	PATSY J TRISLER
510(k) history	2 submissions · 2 cleared · 2002-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020657/>; 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 29, 2026