

**K013623 CLEARLIGHT PHOTOTHERAPY SYSTEM, MODEL CL  
420**Aug 16, 2002  
284 days to decisionK013623 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k013623/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 5, 2001
Decision date	Aug 16, 2002
Days to decision	284 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Curelight, Ltd.</b>
Location	Washington, DC, US
Contact	JONATHAN S KAHAN
510(k) history	6 submissions · 6 cleared · 2002-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013623/> 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 3, 2026