

**K853900 ENDO-LASE CD40 FOR NEUROSURGERY**Jun 18, 1986  
271 days to decisionK853900 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k853900/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 20, 1985
Decision date	Jun 18, 1986
Days to decision	271 days
Third-party review	No

**APPLICANT**

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Company	<b>Endo Lase, Inc.</b>
Location	Walker, MI, US
Contact	<b>&amp; HARTSON</b>
510(k) history	24 submissions · 24 cleared · 1984-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k853900/>; 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 27, 2026