

**K854134 FIBEROPTIC COLLIMATING HANDPIECE**Feb 7, 1986  
120 days to decisionK854134 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k854134/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Oct 10, 1985
Decision date	Feb 7, 1986
Days to decision	120 days
Third-party review	No

**APPLICANT**

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Company	<b>Dawn Laser Technology, Inc.</b>
Location	Santa Barbara, CA, US
Contact	NYLE G UTTERBACK
510(k) history	1 submissions · 1 cleared · 1986-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854134/>; 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 8, 2026