

**K850779 MODEL 4000 ND:YAG LASER FOR GENITOURINARY APPLICAT**Apr 2, 1985  
35 days to decisionK850779 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k850779/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 26, 1985
Decision date	Apr 2, 1985
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Cooper Lasersonics, Inc.</b>
Location	Orangeburg, NY, US
Contact	CHARLES L ROSE
Website	<a href="https://www.cooperhealthcare.com">https://www.cooperhealthcare.com</a>
510(k) history	54 submissions · 52 cleared · 1982-1988

Cooper Lasersonics, Inc. is a medical device manufacturer based in Orangeburg, US, specializing in laser surgical systems. The company has received FDA 510(k) clearances from total submissions since its first clearance in 1982. Cooper Lasersonics focused primarily on CO2 and Nd:YAG laser systems for surgical applications across multiple specialties, including general and plastic surgery, otolaryngology, gastroenterology, and neurosurgery. The company's regulatory activity concluded in 1988, making this a historical record of its contributions to surgical laser technology....

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