

K880170 MODEL 4900 ND:YAG LASER FOR PULMONARY OBSTRUCTIONSMar 24, 1988
69 days to decisionK880170 · Product code: **LLO** · Anesthesiology
Source: <https://www.510kdatabase.net/k880170/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Neodymium:yag, Pulmonary Surgery (LLO)
Date received	Jan 15, 1988
Decision date	Mar 24, 1988
Days to decision	69 days
Third-party review	No

APPLICANT

Company	Cooper Lasersonics, Inc.
Location	Orangeburg, NY, US
Contact	CHARLES L ROSE
Website	https://www.cooperhealthcare.com
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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