

**K862456 ND:YAG LASER 4000/6000/8000 ENDOMETRIAL
ABLATION**Aug 12, 1986
46 days to decisionK862456 · Product code: **HHR** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k862456/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Surgical, Gynecologic (HHR)
Date received	Jun 27, 1986
Decision date	Aug 12, 1986
Days to decision	46 days
Third-party review	No

APPLICANT

Company	Cooper Lasersonics, Inc.
Location	Orangeburg, NY, US
Contact	CHARLES 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....

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862456/> Data sourced from FDA 510(k) public records (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. - Generated May 21, 2026