FDA 510(k) Premarket Notification

K850728 MODEL 4000 ND:YAG LASER FOR GASTROINTESTIONAL BLEE

CLEARED

Apr 8, 1985
45 days to decision

K850728 · Product code: **LNK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k850728/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser For Gastro-urology Use (LNK)
Date received	Feb 22, 1985
Decision date	Apr 8, 1985
Days to decision	45 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....