

K861921 LASER BRONCHOSCOPE SYSTEMJun 2, 1986
14 days to decisionK861921 · Product code: **LLO** · Anesthesiology
Source: <https://www.510kdatabase.net/k861921/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Neodymium:yag, Pulmonary Surgery (LLO)
Date received	May 19, 1986
Decision date	Jun 2, 1986
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Pilling Co.
Location	McHenry, IL, US
Contact	LELAND J WIMER
Website	https://www.pilling.com
510(k) history	39 submissions · 39 cleared · 1976-1994

Pilling Co. is a medical device manufacturer based in McHenry, US, specializing in surgical instrumentation and specialized medical devices. The company has received FDA 510(k) clearances from total submissions, with clearances spanning from 1976 to 1994. Pilling Co. focused primarily on General & Plastic Surgery devices, alongside offerings in Cardiovascular, Dental, Gastroenterology & Urology, and Anesthesiology categories. Notable cleared devices include laparoscopic instrumentation, microlaryngeal forceps, vascular instruments, and coronary perfusion catheters. This c...

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k861921/> 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 27, 2026