

K833557 MEDILAS LASERFeb 7, 1984
123 days to decisionK833557 · Product code: **LNK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k833557/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser For Gastro-urology Use (LNK)
Date received	Oct 7, 1983
Decision date	Feb 7, 1984
Days to decision	123 days
Third-party review	No

APPLICANT

Company	Endo Lase, Inc.
Location	Walker, MI, US
510(k) history	24 submissions · 24 cleared · 1984-1986

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833557/> 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