FDA 510(k) Premarket Notification

K833557 MEDILAS LASER

CLEARED

Feb 7, 1984
123 days to decision

K833557 · Product code: **LNK** · Gastroenterology & Urology
Source: 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