FDA 510(k) Premarket Notification

K860063 ND:YAG LASER SINGLE USE FIBER DELIVERY SYSTEM

CLEARED

Feb 5, 1986
30 days to decision

K860063 · Product code: **LNK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k860063/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser For Gastro-urology Use (LNK)
Date received	Jan 6, 1986
Decision date	Feb 5, 1986
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Laser Media**
Location	Hauppauge, NY, US
Contact	ROBERT A KAPLAN
510(k) history	12 submissions · 12 cleared · 1985-1987