

K860063 ND:YAG LASER SINGLE USE FIBER DELIVERY SYSTEMFeb 5, 1986
30 days to decisionK860063 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k860063/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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