FDA 510(k) Premarket Notification

K874680 LASEGUIDE 600A, 600B, 400A, AND 400B

CLEARED

Mar 23, 1988
131 days to decision

K874680 · Product code: **FCS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k874680/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Catheter, Fiberoptic, Glass, Ureteral (FCS)
Date received	Nov 13, 1987
Decision date	Mar 23, 1988
Days to decision	131 days
Third-party review	No

APPLICANT

Company	**Laser Peripherals, LLC**
Location	Hingham, MA, US
Contact	MICHAEL N BASEL
510(k) history	12 submissions · 12 cleared · 1988-2017