FDA 510(k) Premarket Notification

K880591 LASEGUIDE 600A, 600B, 400A, 400B FOR OB-GYN USE

CLEARED

Jun 1, 1988
111 days to decision

K880591 · Product code: **HHR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k880591/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Surgical, Gynecologic (HHR)
Date received	Feb 11, 1988
Decision date	Jun 1, 1988
Days to decision	111 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