FDA 510(k) Premarket Notification

K881010 CO2 RIGID LASER ENDOSCOPE PROBE SERIES 792/LAPARO

CLEARED

Apr 21, 1988
42 days to decision

K881010 · Product code: **HET** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k881010/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Mar 10, 1988
Decision date	Apr 21, 1988
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Sharplan Lasers, Inc.**
Location	Allendale, NJ, US
Contact	STEPHEN DALTON
510(k) history	78 submissions · 78 cleared · 1986-1997