

**K881010 CO2 RIGID LASER ENDOSCOPE PROBE SERIES
792/LAPARO**Apr 21, 1988
42 days to decisionK881010 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881010/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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