FDA 510(k) Premarket Notification

K880930 MDLS 4000,4900,6000,8000,8900 ND:YAG/1700 C02 LASE

CLEARED

May 31, 1988
88 days to decision

K880930 · Product code: **HHR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k880930/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Surgical, Gynecologic (HHR)
Date received	Mar 4, 1988
Decision date	May 31, 1988
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Copper Lasersonics, Inc.**
Location	Santa Clara, CA, US
Contact	CHARLES L ROSE
510(k) history	1 submissions · 1 cleared · 1988-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k880930/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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