

K853903 ENDO-LASE CD40 CO2 FOR GYNECOLOGYFeb 21, 1986
154 days to decisionK853903 · Product code: **HHR** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k853903/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Surgical, Gynecologic (HHR)
Date received	Sep 20, 1985
Decision date	Feb 21, 1986
Days to decision	154 days
Third-party review	No

APPLICANT

Company	Endo Lase, Inc.
Location	Walker, MI, US
Contact	JONATHAN S KAHAN
510(k) history	24 submissions · 24 cleared · 1984-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k853903/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026