FDA 510(k) Premarket Notification

# K853903 ENDO-LASE CD40 CO2 FOR GYNECOLOGY

**CLEARED**

Feb 21, 1986
154 days to decision

K853903 · Product code: **HHR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k853903/

## SUBMISSION DETAILS

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|---|---|
| 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 |