FDA 510(k) Premarket Notification

K831241 LASER LAPARAXOPE

CLEARED

Jun 10, 1983
56 days to decision

K831241 · Product code: **HET** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k831241/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Apr 15, 1983
Decision date	Jun 10, 1983
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Eder Instrument Co, Inc.**
Location	Walker, MI, US
510(k) history	10 submissions · 10 cleared · 1979-1984