

K831241 LASER LAPARAXOPEJun 10, 1983
56 days to decisionK831241 · Product code: **HET** · Obstetrics & GynecologySource: <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

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Device record: <https://www.510kdatabase.net/k831241/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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