

**K791052 ENDOSCOPIC INSTRUMENT**Aug 3, 1979  
58 days to decisionK791052 · Product code: **HET** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k791052/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Jun 6, 1979
Decision date	Aug 3, 1979
Days to decision	58 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k791052/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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