

**K940718 INNERVISION INC. LAPAROSCOPIC HAND INSTRUMENTS**Jul 22, 1994  
154 days to decisionK940718 · Product code: **HET** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k940718/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Feb 18, 1994
Decision date	Jul 22, 1994
Days to decision	154 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Innervision, Inc.</b>
Location	Memphis, TN, US
Contact	FRANK M LEWIS
510(k) history	6 submissions · 6 cleared · 1994-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k940718/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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