

**K990372 MODULAR INSTRUMENT SYSTEM FOR MINIMAL
INVASIVE SURGERY**Oct 14, 1999
248 days to decisionK990372 · Product code: **HET** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k990372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Feb 8, 1999
Decision date	Oct 14, 1999
Days to decision	248 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Dr. Fritz GmbH
Location	Baldwin Park, CA, US
Contact	JAMES CHITTY
510(k) history	1 submissions · 1 cleared · 1999-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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