

**K933300 GEISTER SURGICAL INT'L MONOPOLAR
ENDOSCOPIC SURG.INST.**Sep 30, 1994
450 days to decisionK933300 · Product code: **KNF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k933300/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coagulator-cutter, Endoscopic, Unipolar (and Accessories) (KNF)
Date received	Jul 7, 1993
Decision date	Sep 30, 1994
Days to decision	450 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lyle Howard Corp.
Location	Lebanon, NJ, US
Contact	LYNETTE HOWARD
510(k) history	1 submissions · 1 cleared · 1994-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933300/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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