

**K921727 CORE MONITORING SYSTEM-
SP1475/6275/5607/5708/4575**May 10, 1994
760 days to decisionK921727 · Product code: **KNF** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k921727/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coagulator-cutter, Endoscopic, Unipolar (and Accessories) (KNF)
Date received	Apr 10, 1992
Decision date	May 10, 1994
Days to decision	760 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Viggo-Spectramed
Location	Oxnard, CA, US
Contact	CHERYL L SHEA
510(k) history	1 submissions · 1 cleared · 1994-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921727/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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