FDA 510(k) Premarket Notification

K921727 CORE MONITORING SYSTEM-SP1475/6275/5607/5708/4575

CLEARED

May 10, 1994
760 days to decision

K921727 · 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