FDA 510(k) Premarket Notification

K902555 AMCATH COAGULATION PROBE

CLEARED

Jun 20, 1990
12 days to decision

K902555 · Product code: **HFG** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k902555/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coagulator, Laparoscopic, Unipolar (and Accessories) (HFG)
Date received	Jun 8, 1990
Decision date	Jun 20, 1990
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Intl. Medical, Inc.**
Location	Mchenry, IL, US
Contact	PETER H WETTERMANN
510(k) history	17 submissions · 17 cleared · 1979-1997