

K902555 AMCATH COAGULATION PROBEJun 20, 1990
12 days to decisionK902555 · Product code: **HFG** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k902555/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 25, 2026