

K905752 ARGON BEAM COMBO LAPAROSCOPIC ELECTRODEJul 22, 1991
210 days to decisionK905752 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k905752/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 24, 1990
Decision date	Jul 22, 1991
Days to decision	210 days
Third-party review	No

APPLICANT

Company	Beacon Laboratories, Inc.
Location	Westminster, CO, US
Contact	RICHARD P FLEENOR
510(k) history	18 submissions · 18 cleared · 1989-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k905752/>; 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 June 28, 2026