

K972350 MODEL 50 BIPOLAR COAGULATORAug 21, 1997
58 days to decisionK972350 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k972350/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 24, 1997
Decision date	Aug 21, 1997
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Boston Surgical Products, Inc.
Location	Kingston, MA, US
Contact	CHARLES VASSALLO
510(k) history	17 submissions · 17 cleared · 1994-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k972350/> 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 29, 2026