

**K970809 BENDABLE TISSUE COAGULATING ELECTRODE
WITH BENDING TOOL**Jun 18, 1997
105 days to decisionK970809 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k970809/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 5, 1997
Decision date	Jun 18, 1997
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Somnus Medical Technologies, Inc.
Location	Sunnyvale, CA, US
Contact	EVE CONNER
510(k) history	16 submissions · 16 cleared · 1996-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970809/>; 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 July 3, 2026