

**K970837 SOMNUS MODEL 6000 DISPOSABLE TISSUE  
COAGULATING ELECTRODE**Jan 9, 1998  
308 days to decisionK970837 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k970837/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 7, 1997
Decision date	Jan 9, 1998
Days to decision	308 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970837/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated July 3, 2026