

K012004 EASY GUIDE ELECTROSURGICAL ACCESS DEVICESep 25, 2001
90 days to decisionK012004 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k012004/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 27, 2001
Decision date	Sep 25, 2001
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Senorx, Inc.
Location	Irvine, CA, US
Contact	AMY BOUCLY
510(k) history	30 submissions · 26 cleared · 2000-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012004/> 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