

K013641 SENORX BIOPSY DEVICE, DRIVER, CONTROL MODULE, INCLUDING ACCESSORIES (POWER CORD,CONNECTOR CORDS AND FOOTSWITCH), VACUUMS

Jan 29, 2002
85 days to decision

K013641 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k013641/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 5, 2001
Decision date	Jan 29, 2002
Days to decision	85 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.net

Device record: <https://www.510kdatabase.net/k013641/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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