FDA 510(k) Premarket Notification

K012004 EASY GUIDE ELECTROSURGICAL ACCESS DEVICE

CLEARED

Sep 25, 2001
90 days to decision

K012004 · Product code: **GEI** · General & Plastic Surgery
Source: 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