FDA 510(k) Premarket Notification

K030890 SWITCHBLADE

CLEARED

Apr 4, 2003
14 days to decision

K030890 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k030890/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 21, 2003
Decision date	Apr 4, 2003
Days to decision	14 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Snowden Pencer, Inc.**
Location	Tucker, GA, US
Contact	DAVID J BOOTH
510(k) history	1 submissions · 1 cleared · 2003-2003