FDA 510(k) Premarket Notification

K000170 SECCA TUBULAR ELECTRODE DEVICE, MODEL A4000

CLEARED

Feb 14, 2000
27 days to decision

K000170 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k000170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 18, 2000
Decision date	Feb 14, 2000
Days to decision	27 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Conway Stuart Medical, Inc.**
Location	Sunnyvale, CA, US
Contact	THOMAS C WEHMAN
510(k) history	10 submissions · 10 cleared · 1998-2000