

K000170 SECCA TUBULAR ELECTRODE DEVICE, MODEL A4000Feb 14, 2000
27 days to decisionK000170 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000170/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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