FDA 510(k) Premarket Notification

K042909 STELLARTECH COAGULATION SYSTEM, MODELS 1100C-115A & 1100C-230A

CLEARED

Nov 12, 2004
22 days to decision

K042909 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k042909/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 21, 2004
Decision date	Nov 12, 2004
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stellartech Research Corp.**
Location	Sunnyvale, CA, US
Contact	JAMES R SANTOS
510(k) history	11 submissions · 11 cleared · 2000-2006

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k042909/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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