FDA 510(k) Premarket Notification

K013139 STELLARTECH COAGULATION SYSTEM

CLEARED

Dec 18, 2001
90 days to decision

K013139 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k013139/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 19, 2001
Decision date	Dec 18, 2001
Days to decision	90 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