FDA 510(k) Premarket Notification

K942489 ELECTROSURGICAL CUTTING & COAGULATION DEVICE

CLEARED

Jul 25, 1994
61 days to decision

K942489 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k942489/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 25, 1994
Decision date	Jul 25, 1994
Days to decision	61 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Megadyne Medical Products, Inc.**
Location	Murray, UT, US
Contact	DREW D WEAVER
510(k) history	50 submissions · 48 cleared · 1990-2024