FDA 510(k) Premarket Notification

K001330 SOVEREIGN BIPOLAR INSTRUMENTS

CLEARED

May 30, 2000
33 days to decision

K001330 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k001330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 27, 2000
Decision date	May 30, 2000
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aesculap, Inc.**
Location	Burlingame, CA, US
Contact	LIA S JONES
510(k) history	207 submissions · 201 cleared · 1991-2025