

K001330 SOVEREIGN BIPOLAR INSTRUMENTSMay 30, 2000
33 days to decisionK001330 · Product code: **GEI** · General & Plastic SurgerySource: <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 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001330/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026