

**K822513 MODEL #2000 HI-TEMP & #3000 LOW-TEMP**Sep 21, 1982  
32 days to decisionK822513 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k822513/>**SUBMISSION DETAILS**

---

|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                         |
| Submission type       | Traditional  |
| Device classification | Electrosurgical, Cutting & Coagulation & Accessories (GEI) |
| Date received         | Aug 20, 1982   |
| Decision date         | Sep 21, 1982   |
| Days to decision      | 32 days  |
| Third-party review    | No   |

**APPLICANT**

---

|                |  |
|----------------|--|
| Company        | <b>Surgicare Scientific Laboratory, Inc.</b> |
| Location       | Mchenry, IL, US                              |
| 510(k) history | 3 submissions · 3 cleared · 1982-1983        |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k822513/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 9, 2026