

**K813072 E2502 & E2502B LECTROSWITCH**Nov 10, 1981  
8 days to decisionK813072 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k813072/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 2, 1981
Decision date	Nov 10, 1981
Days to decision	8 days
Third-party review	No

**APPLICANT**

---

Company	<b>Valleylab, Inc.</b>
Location	Mchenry, IL, US
510(k) history	94 submissions · 93 cleared · 1976-2003

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k813072/> 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 May 22, 2026