

K791639 MODEL E2515 ELECTROSURGICAL ACCESSORYSep 12, 1979
22 days to decisionK791639 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k791639/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 21, 1979
Decision date	Sep 12, 1979
Days to decision	22 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k791639/> 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 May 22, 2026