

**K791638 MODEL E2510 SURGISWITCH**Sep 12, 1979  
22 days to decisionK791638 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k791638/>**SUBMISSION DETAILS**

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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**

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

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

Device record: <https://www.510kdatabase.net/k791638/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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