

K822572 POLYHESIVE PATIENT RETURN ELECTRODESep 21, 1982
28 days to decisionK822572 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k822572/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 24, 1982
Decision date	Sep 21, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k822572/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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