

K964602 DAVIS BAYONET ELECTRODESJan 9, 1997
52 days to decisionK964602 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k964602/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 18, 1996
Decision date	Jan 9, 1997
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k964602/> 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