

**K012625 REPROCESSED UNIPOLAR
LAPAROSCOPIC/ENDOSCOPIC INSTRUMENTS**Oct 22, 2001
70 days to decisionK012625 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k012625/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 13, 2001
Decision date	Oct 22, 2001
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Alliance Medical Corp.
Location	Phoenix, AZ, US
Contact	DON SELVEY
510(k) history	36 submissions · 36 cleared · 2001-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012625/> 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 20, 2026