

K940750 INNERVISION,INC. LAPAROSCOPIC ELECTRODESAug 23, 1994
186 days to decisionK940750 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k940750/>**SUBMISSION DETAILS**

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| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrosurgical, Cutting & Coagulation & Accessories (GEI) |
| Date received | Feb 18, 1994 |
| Decision date | Aug 23, 1994 |
| Days to decision | 186 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Innervision, Inc. |
| Location | Memphis, TN, US |
| Contact | FRANK M LEWIS |
| 510(k) history | 6 submissions · 6 cleared · 1994-1999 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k940750/> 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 21, 2026