

K913625 ELECTROSCOPE MONITOR, EM-1Dec 30, 1991
137 days to decisionK913625 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k913625/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 15, 1991
Decision date	Dec 30, 1991
Days to decision	137 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Electroscope, Inc.
Location	Boulder, CO, US
Contact	ROGER C ODELL
510(k) history	6 submissions · 6 cleared · 1991-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913625/> 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 July 3, 2026