

K932817 HM-880 IMar 9, 1994
273 days to decisionK932817 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k932817/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 9, 1993
Decision date	Mar 9, 1994
Days to decision	273 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Hill-Med, Inc.
Location	Miami, FL, US
Contact	ERNESTO ACKERMAN
510(k) history	10 submissions · 10 cleared · 1993-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932817/> 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 26, 2026