

**K936075 REUSABLE, HAND CONTROLLED ESP**Jun 6, 1994  
168 days to decisionK936075 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k936075/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 20, 1993
Decision date	Jun 6, 1994
Days to decision	168 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>E &amp; M Engineering, Inc.</b>
Location	Richmond, VA, US
Contact	DUNCAN FUNG
510(k) history	6 submissions · 6 cleared · 1993-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k936075/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026