

**K924881 REFLEX(R) ESP**Jul 23, 1993  
298 days to decisionK924881 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k924881/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 28, 1992
Decision date	Jul 23, 1993
Days to decision	298 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Richard-Allan Medical</b>
Location	Richland, MI, US
Contact	JULIE POWELL
510(k) history	8 submissions · 8 cleared · 1991-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924881/> 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