

K932392 PHX TECHNOLOGIES CORP ELECTROSURGICAL PROBESNov 22, 1993
188 days to decisionK932392 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k932392/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 18, 1993
Decision date	Nov 22, 1993
Days to decision	188 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Phx Technologies Corp.
Location	Denton, TX, US
Contact	JAMES F CHAPEL
510(k) history	49 submissions · 49 cleared · 1993-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932392/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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