FDA 510(k) Premarket Notification

K932392 PHX TECHNOLOGIES CORP ELECTROSURGICAL PROBES

CLEARED

Nov 22, 1993
188 days to decision

K932392 · Product code: **GEI** · General & Plastic Surgery
Source: 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