FDA 510(k) Premarket Notification

K936075 REUSABLE, HAND CONTROLLED ESP

CLEARED

Jun 6, 1994
168 days to decision

K936075 · Product code: **GEI** · General & Plastic Surgery
Source: 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	**E & M Engineering, Inc.**
Location	Richmond, VA, US
Contact	DUNCAN FUNG
510(k) history	6 submissions · 6 cleared · 1993-1999