FDA 510(k) Premarket Notification

K924881 REFLEX(R) ESP

CLEARED

Jul 23, 1993
298 days to decision

K924881 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k924881/

SUBMISSION DETAILS

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

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