FDA 510(k) Premarket Notification

K905108 CONTORNO(R), MODELS E AND ST

CLEARED

Feb 4, 1991
83 days to decision

K905108 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k905108/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 13, 1990
Decision date	Feb 4, 1991
Days to decision	83 days
Third-party review	No

APPLICANT

Company	**Palex Intl. SA**
Location	Spain, ES
Contact	MARTIN A JORDAN
510(k) history	5 submissions · 5 cleared · 1990-1991