FDA 510(k) Premarket Notification

K930243 FLEXIBLE EXTENDER

CLEARED

Mar 11, 1994
416 days to decision

K930243 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k930243/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 19, 1993
Decision date	Mar 11, 1994
Days to decision	416 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Beacon Laboratories, Inc.**
Location	Westminster, CO, US
Contact	RICHARD P FLEENOR
510(k) history	18 submissions · 18 cleared · 1989-1994