FDA 510(k) Premarket Notification

K902035 FLEXIBLE EXTENDER

CLEARED

Jul 19, 1990
77 days to decision

K902035 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k902035/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 3, 1990
Decision date	Jul 19, 1990
Days to decision	77 days
Third-party review	No

APPLICANT

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