FDA 510(k) Premarket Notification

K901302 BIPOLARD POLYPECTOMY SYSTEM (TM)

CLEARED

Jun 6, 1990
78 days to decision

K901302 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k901302/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 20, 1990
Decision date	Jun 6, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Medical Devices, Inc.**
Location	Mchenry, IL, US
Contact	SAMUEL DICKSTEIN
510(k) history	49 submissions · 47 cleared · 1977-2001