FDA 510(k) Premarket Notification

K911274 UNIPOLAR COAGULATOR PROBE SYSTEM

CLEARED

Jun 20, 1991
90 days to decision

K911274 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k911274/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 22, 1991
Decision date	Jun 20, 1991
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**American Surgical Instruments Corp.**
Location	Washington, DC, US
Contact	DORSEY III
510(k) history	6 submissions · 5 cleared · 1990-1993