FDA 510(k) Premarket Notification

K910334 ELECTROCAUTERY PROBE

CLEARED

Feb 21, 1991
27 days to decision

K910334 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k910334/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 25, 1991
Decision date	Feb 21, 1991
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Wiltek Medical, Inc.**
Location	Winston-Salem, NC, US
Contact	JON S WILSON
510(k) history	21 submissions · 21 cleared · 1989-1999