FDA 510(k) Premarket Notification

K974484 ISOLATED ELECTROSURGICAL PROBES AND DEVICES-SALINETRODE

CLEARED

Feb 10, 1998
76 days to decision

K974484 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k974484/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 26, 1997
Decision date	Feb 10, 1998
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ximed/Prosure/Injectx**
Location	San Jose, CA, US
Contact	ASHVIN DESAI
510(k) history	9 submissions · 9 cleared · 1995-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k974484/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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