FDA 510(k) Premarket Notification

K993412 BITX PROBES

CLEARED

Mar 31, 2000
171 days to decision

K993412 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k993412/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 12, 1999
Decision date	Mar 31, 2000
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Everest Medical Corp.**
Location	Brooklyn Center, MN, US
Contact	FREDERICK G MADES
510(k) history	21 submissions · 21 cleared · 1987-2000

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k993412/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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