FDA 510(k) Premarket Notification

K992572 NTERO RF SLEEVE

CLEARED

Sep 24, 1999
53 days to decision

K992572 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k992572/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 2, 1999
Decision date	Sep 24, 1999
Days to decision	53 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ntero Surgical, Inc.**
Location	Palo Alto, CA, US
Contact	D. BOMMI BOMMANNAN
510(k) history	3 submissions · 3 cleared · 1999-2002