

**K992572 NTERO RF SLEEVE**Sep 24, 1999  
53 days to decisionK992572 · Product code: **GEI** · General & Plastic SurgerySource: <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	<b>Ntero Surgical, Inc.</b>
Location	Palo Alto, CA, US
Contact	D. BOMMI BOMMANNAN
510(k) history	3 submissions · 3 cleared · 1999-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992572/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 3, 2026