

**K911159 PALPATION PROBE**Mar 29, 1991  
14 days to decisionK911159 · Product code: **GEN** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k911159/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, General & Plastic Surgery (GEN)
Date received	Mar 15, 1991
Decision date	Mar 29, 1991
Days to decision	14 days
Third-party review	No

**APPLICANT**

---

Company	<b>Northgate Technologies, Inc.</b>
Location	Arlington Heights, IL, US
Contact	SAMUELLA D EMRICH
510(k) history	55 submissions · 55 cleared · 1991-2021

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911159/>; 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 June 28, 2026