

**K042648 REPROCESSED COLD BIOPSY FORCEPS**Jul 20, 2005  
295 days to decisionK042648 · Product code: **NON** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k042648/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Non-electric, Reprocessed (NON)
Date received	Sep 28, 2004
Decision date	Jul 20, 2005
Days to decision	295 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Sterilmed, Inc.</b>
Location	Plymouth, MN, US
Contact	BRUCE LESTER
510(k) history	64 submissions · 64 cleared · 2001-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k042648/> 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 May 19, 2026