

K070930 REPROCESSED AUTOSUTURE GIA ENDOSCOPIC STAPLERSOct 18, 2007
198 days to decisionK070930 · Product code: **NLL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k070930/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable, Reprocessed (NLL)
Date received	Apr 3, 2007
Decision date	Oct 18, 2007
Days to decision	198 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sterilmed, Inc.
Location	Plymouth, MN, US
Contact	DENNIS TOUSSAINT
510(k) history	64 submissions · 64 cleared · 2001-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k070930/> Data sourced from FDA 510(k) public records (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. - Generated May 19, 2026