

K012571 REPROCESSED HARMONICNov 7, 2001
90 days to decisionK012571 · Product code: **NLQ** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k012571/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single-use Reprocessed Ultrasonic Surgical Instruments (NLQ)
Date received	Aug 9, 2001
Decision date	Nov 7, 2001
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012571/> 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