

**K051220 REPROCESSED DEFLECTABLE
ELECTROPHYSIOLOGY DIAGNOSTIC CATHETERS**Jun 14, 2005
33 days to decisionK051220 · Product code: **NLH** · Cardiovascular
Source: <https://www.510kdatabase.net/k051220/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Recording, Electrode, Reprocessed (NLH)
Date received	May 12, 2005
Decision date	Jun 14, 2005
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051220/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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