

K063844 REPROCESSED HEART STABILIZERS AND POSITIONERSMar 15, 2007
79 days to decisionK063844 · Product code: **NQG** · Cardiovascular
Source: <https://www.510kdatabase.net/k063844/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stabilizer, Heart, Non-compression, Reprocessed (NQG)
Date received	Dec 26, 2006
Decision date	Mar 15, 2007
Days to decision	79 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/k063844/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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