

K120204 REPROCESSED FEMORAL COMPRESSION DEVICEApr 3, 2012
71 days to decisionK120204 · Product code: **NMF** · Cardiovascular
Source: <https://www.510kdatabase.net/k120204/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular, Reprocessed (NMF)
Date received	Jan 23, 2012
Decision date	Apr 3, 2012
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120204/> 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