

**K014223 MODIFICATION TO GUARDWIRE PLUS TEMPORARY
OCCLUSION & ASPIRATION SYSTEM**Jan 25, 2002
30 days to decisionK014223 · Product code: **NFA** · Cardiovascular
Source: <https://www.510kdatabase.net/k014223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Temporary Coronary Saphenous Vein Bypass Graft For Embolic Protection (NFA)
Date received	Dec 26, 2001
Decision date	Jan 25, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medtronic Percusurge, Inc.
Location	Sunnyvale, CA, US
Contact	MATT MOON
510(k) history	2 submissions · 2 cleared · 2002-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k014223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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