FDA 510(k) Premarket Notification

K073079 THUMPER, MODEL: 1008

CLEARED

Feb 14, 2008
106 days to decision

K073079 · Product code: **DRM** · Cardiovascular
Source: https://www.510kdatabase.net/k073079/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Oct 31, 2007
Decision date	Feb 14, 2008
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Michigan Instruments, Inc.**
Location	Mchenry, IL, US
Contact	HOWARD M HOSTEIN
510(k) history	7 submissions · 7 cleared · 1981-2008