

**K073079 THUMPER, MODEL: 1008**Feb 14, 2008  
106 days to decisionK073079 · 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	<b>Michigan Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	HOWARD M HOSTEIN
510(k) history	7 submissions · 7 cleared · 1981-2008

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073079/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026