

**K851139 THUMPER CARDIOPULMONARY RESUSCITATOR  
1005**May 15, 1985  
55 days to decisionK851139 · Product code: **DRM** · Cardiovascular  
Source: <https://www.510kdatabase.net/k851139/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Mar 21, 1985
Decision date	May 15, 1985
Days to decision	55 days
Third-party review	No

**APPLICANT**

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Company	<b>Michigan Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	JIM MAATMAN
510(k) history	7 submissions · 7 cleared · 1981-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k851139/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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