

K833879 CARDIOPULMONARY RESUSCITATORDec 27, 1983
49 days to decisionK833879 · Product code: **DRM** · Cardiovascular
Source: <https://www.510kdatabase.net/k833879/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Nov 8, 1983
Decision date	Dec 27, 1983
Days to decision	49 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k833879/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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