

K811909 CARDIOPULMUNARY RESUSCITATORJul 16, 1981
15 days to decisionK811909 · Product code: **DRM** · CardiovascularSource: <https://www.510kdatabase.net/k811909/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Jul 1, 1981
Decision date	Jul 16, 1981
Days to decision	15 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/k811909/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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