FDA 510(k) Premarket Notification

K833879 CARDIOPULMONARY RESUSCITATOR

CLEARED

Dec 27, 1983
49 days to decision

K833879 · 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