FDA 510(k) Premarket Notification

K811909 CARDIOPULMUNARY RESUSCITATOR

CLEARED

Jul 16, 1981
15 days to decision

K811909 · Product code: **DRM** · Cardiovascular
Source: 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