FDA 510(k) Premarket Notification

K851139 THUMPER CARDIOPULMONARY RESUSCITATOR 1005

CLEARED

May 15, 1985
55 days to decision

K851139 · Product code: **DRM** · Cardiovascular
Source: https://www.510kdatabase.net/k851139/

SUBMISSION DETAILS

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

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