FDA 510(k) Premarket Notification

K063602 AUTOPULSE RESUSCITATION SYSTEM MODEL100

CLEARED

Dec 21, 2006
17 days to decision

K063602 · Product code: **DRM** · Cardiovascular
Source: https://www.510kdatabase.net/k063602/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Compressor, Cardiac, External (DRM)
Date received	Dec 4, 2006
Decision date	Dec 21, 2006
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Zoll Circulation**
Location	Sunnyvale, CA, US
Contact	MARK PERKINS
510(k) history	5 submissions · 4 cleared · 2006-2014