FDA 510(k) Premarket Notification

K040453 AUTOPULSE RESUSCITATION SYSTEM, MODEL 100

CLEARED

Mar 11, 2004
17 days to decision

K040453 · Product code: **DRM** · Cardiovascular
Source: https://www.510kdatabase.net/k040453/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Compressor, Cardiac, External (DRM)
Date received	Feb 23, 2004
Decision date	Mar 11, 2004
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Revivant Corp.**
Location	Sunnyvale, CA, US
Contact	BOB H KATZ
510(k) history	4 submissions · 4 cleared · 2001-2004