FDA 510(k) Premarket Notification

K031987 POWERHEART AED G3, MODEL 9300

CLEARED

Jul 30, 2003
33 days to decision

K031987 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k031987/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jun 27, 2003
Decision date	Jul 30, 2003
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardiac Science**
Location	Edina, MN, US
Contact	KENNETH OLSON
510(k) history	1 submissions · 1 cleared · 2003-2003