FDA 510(k) Premarket Notification

K020288 PADPRO, MODEL 2603

CLEARED

Feb 27, 2002
30 days to decision

K020288 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k020288/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jan 28, 2002
Decision date	Feb 27, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Padpro, LLC**
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	5 submissions · 5 cleared · 2001-2002