FDA 510(k) Premarket Notification

K061606 DEFIBRILLATOR CABLE TESTER, MODEL DT2200

CLEARED

Jun 21, 2006
12 days to decision

K061606 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k061606/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jun 9, 2006
Decision date	Jun 21, 2006
Days to decision	12 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Medical Devices/Padpro, Inc.**
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	1 submissions · 1 cleared · 2006-2006