FDA 510(k) Premarket Notification

K002280 PAD PRO PEDIATRIC ELECTRODE, MODEL PAD PRO 2602

CLEARED

Feb 5, 2001
194 days to decision

K002280 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k002280/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 26, 2000
Decision date	Feb 5, 2001
Days to decision	194 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medical Devices, Inc.**
Location	Mchenry, IL, US
Contact	WARREN R WALTERS
510(k) history	49 submissions · 47 cleared · 1977-2001