FDA 510(k) Premarket Notification

K941970 FAST-PATCH

CLEARED

Dec 7, 1994
229 days to decision

K941970 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k941970/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Apr 22, 1994
Decision date	Dec 7, 1994
Days to decision	229 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Physio-Control Corp.**
Location	Mchenry, IL, US
Contact	SHERRI L POCOCK
510(k) history	80 submissions · 78 cleared · 1976-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k941970/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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