

**K945511 LIFEPAK 10 DEFIBRILLATOR/MONITOR/PACEMAKER,
AND LIFEPAK 11 CARDIAC MONITOR**Sep 26, 1995
321 days to decisionK945511 · Product code: **MPE** · Cardiovascular
Source: <https://www.510kdatabase.net/k945511/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Auxiliary Power Supply (acor Dc) For External Transcutaneous Cardiac Pacemaker (MPE)
Date received	Nov 9, 1994
Decision date	Sep 26, 1995
Days to decision	321 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Physio-Control Corp.
Location	Mchenry, IL, US
Contact	MICHAEL D WILLINGHAM
510(k) history	80 submissions · 78 cleared · 1976-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k945511/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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