

**K946080 PHYSIO CONTROL LIFEPAK 200 AUTO
DEFIBRILLATOR**Aug 2, 1995
232 days to decisionK946080 · Product code: **FCO** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k946080/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Box, Battery, Rechargeable (FCO)
Date received	Dec 13, 1994
Decision date	Aug 2, 1995
Days to decision	232 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Portable Power Systems, Inc.
Location	Castle Rock, CO, US
Contact	NORMAN A PREMOP
510(k) history	18 submissions · 18 cleared · 1992-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k946080/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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