FDA 510(k) Premarket Notification

K946080 PHYSIO CONTROL LIFEPAK 200 AUTO DEFIBRILLATOR

CLEARED

Aug 2, 1995
232 days to decision

K946080 · 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