FDA 510(k) Premarket Notification

K832876 LIFEPAK 7P

CLEARED

Jan 30, 1984
157 days to decision

K832876 · Product code: **DRO** · Cardiovascular
Source: https://www.510kdatabase.net/k832876/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Aug 26, 1983
Decision date	Jan 30, 1984
Days to decision	157 days
Third-party review	No

APPLICANT

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