

K832876 LIFEPAK 7PJan 30, 1984
157 days to decisionK832876 · Product code: **DRO** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k832876/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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