

K810154 LIFEPAK 7 ACUTE CARDIAC CARE SYSTEMMar 26, 1981
64 days to decisionK810154 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k810154/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jan 21, 1981
Decision date	Mar 26, 1981
Days to decision	64 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/k810154/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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