

**K760676 LIFEPAK 5 CARDIOSCOPE-RECORDER MODULE**Oct 13, 1976  
23 days to decisionK760676 · Product code: **DSF** · CardiovascularSource: <https://www.510kdatabase.net/k760676/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Sep 20, 1976
Decision date	Oct 13, 1976
Days to decision	23 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k760676/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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