

**K912966 PEDIATRIC DISPOSABLE EXTERNAL  
DEFIBRILATION/ECG**Feb 21, 1992  
228 days to decisionK912966 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k912966/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jul 8, 1991
Decision date	Feb 21, 1992
Days to decision	228 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Physio-Control Corp.</b>
Location	Mchenry, IL, US
Contact	SHERRI L.POCOCK
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/k912966/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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