

K895379 LOW ENERGY DC - DEFIBRILLATOR (INCLUDING PADDLES)Feb 26, 1990
180 days to decisionK895379 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k895379/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 30, 1989
Decision date	Feb 26, 1990
Days to decision	180 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895379/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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