

**K900169 MODIFIED HEARTSTART 2000 DEFIBRILLATION
ELECTRODE**Jan 26, 1990
14 days to decisionK900169 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k900169/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jan 12, 1990
Decision date	Jan 26, 1990
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Laerdal Mfg. Corp.
Location	Portland, OR, US
Contact	L KARPOWICZ
510(k) history	5 submissions · 5 cleared · 1988-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900169/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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