

**K902429 MODIFIED HEARTSTART DEFIBRILLATION  
ELECTRODE**Sep 27, 1990  
128 days to decisionK902429 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k902429/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	May 22, 1990
Decision date	Sep 27, 1990
Days to decision	128 days
Third-party review	No

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902429/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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