

**K922173 MODEL # 830 ST ADULT STIMULATION PADS  
(STERILE)**Jun 11, 1992  
34 days to decisionK922173 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k922173/>**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 8, 1992
Decision date	Jun 11, 1992
Days to decision	34 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cardiotronics, Inc.</b>
Location	West Carlsbad, CA, US
Contact	TIM J WAY
510(k) history	27 submissions · 27 cleared · 1988-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922173/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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