

**K844103 CARDIOPAC DEFIBRILLATOR 444**Jan 18, 1985  
87 days to decisionK844103 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k844103/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 23, 1984
Decision date	Jan 18, 1985
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>Birtcher Corp.</b>
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k844103/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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