

**K852808 CR26, CR26A, CR26B, CR26C(CARDIAC DC DEFIBRILLATO)**Sep 16, 1985  
76 days to decisionK852808 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k852808/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jul 2, 1985
Decision date	Sep 16, 1985
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardiac Recorders, Ltd.</b>
Location	London, GB
Contact	BEVIS
510(k) history	8 submissions · 8 cleared · 1985-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852808/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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