

K800937 R2 PLATE TYPE CABLE-ADAPTOR #S 175,176May 23, 1980
32 days to decisionK800937 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k800937/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Apr 21, 1980
Decision date	May 23, 1980
Days to decision	32 days
Third-party review	No

APPLICANT

Company	R2 Corp.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1980-1983

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Device record: <https://www.510kdatabase.net/k800937/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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