

K801544 PORTA-FIB LPDJul 14, 1980
7 days to decisionK801544 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k801544/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jul 7, 1980
Decision date	Jul 14, 1980
Days to decision	7 days
Third-party review	No

APPLICANT

Company	Narco Bio-Systems
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1976-1986

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Device record: <https://www.510kdatabase.net/k801544/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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