

K822961 PORTA FIB IIINov 1, 1982
27 days to decisionK822961 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k822961/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 5, 1982
Decision date	Nov 1, 1982
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Narco Scientific
Location	Walker, MI, US
510(k) history	30 submissions · 30 cleared · 1979-1983

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Device record: <https://www.510kdatabase.net/k822961/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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