FDA 510(k) Premarket Notification

K822961 PORTA FIB III

CLEARED

Nov 1, 1982
27 days to decision

K822961 · Product code: **LDD** · Cardiovascular
Source: 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