FDA 510(k) Premarket Notification

K800937 R2 PLATE TYPE CABLE-ADAPTOR #S 175,176

CLEARED

May 23, 1980
32 days to decision

K800937 · Product code: **LDD** · Cardiovascular
Source: 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