FDA 510(k) Premarket Notification

K863303 MODEL 617D

CLEARED

Sep 5, 1986
11 days to decision

K863303 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k863303/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 25, 1986
Decision date	Sep 5, 1986
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
Contact	RON R DUCK
510(k) history	164 submissions · 160 cleared · 1976-2003