FDA 510(k) Premarket Notification

K914458 ESCORT 300A SERIES DEFIBRILLATOR/PACER OPTION

CLEARED

Dec 9, 1991
63 days to decision

K914458 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k914458/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 7, 1991
Decision date	Dec 9, 1991
Days to decision	63 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medical Data Electronics**
Location	Arleta, CA, US
Contact	CHIP HARLOW
510(k) history	27 submissions · 27 cleared · 1985-2002