

**K914458 ESCORT 300A SERIES DEFIBRILLATOR/PACER
OPTION**Dec 9, 1991
63 days to decisionK914458 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914458/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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