FDA 510(k) Premarket Notification

K874091 360D DC DEFIBRILLATOR

CLEARED

Nov 23, 1987
47 days to decision

K874091 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k874091/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 7, 1987
Decision date	Nov 23, 1987
Days to decision	47 days
Third-party review	No

APPLICANT

Company	**Medac, Inc.**
Location	Tualatin, OR, US
Contact	ROBERT M BOONSTRA
510(k) history	10 submissions · 10 cleared · 1987-1988