FDA 510(k) Premarket Notification

K895185 FIRST MEDIC MODEL 610P

CLEARED

Oct 27, 1989
77 days to decision

K895185 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k895185/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 11, 1989
Decision date	Oct 27, 1989
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**First Medical Devices Corp.**
Location	Bellevue, WA, US
Contact	BRUCE G HAGGAR
510(k) history	7 submissions · 7 cleared · 1986-1990