FDA 510(k) Premarket Notification

K911604 MDPHONE CELLULAR + RESCUE (DEFIB SYSTEM)

CLEARED

Sep 23, 1991
166 days to decision

K911604 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k911604/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Apr 10, 1991
Decision date	Sep 23, 1991
Days to decision	166 days
Third-party review	No

APPLICANT

Company	**Medphone Corp.**
Location	Washington, DC, US
Contact	GLENN KRAPF
510(k) history	2 submissions · 2 cleared · 1987-1991