FDA 510(k) Premarket Notification

K931803 5200 ADULT SOLID GEL MULTI-FUNCTION ELECTRODES

CLEARED

Oct 8, 1993
179 days to decision

K931803 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k931803/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Apr 12, 1993
Decision date	Oct 8, 1993
Days to decision	179 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Zoll Medical Corp**
Location	Woburn, MA, US
Contact	BLAKE A CERULLO
510(k) history	33 submissions · 27 cleared · 1993-2014