FDA 510(k) Premarket Notification

K934016 POWERCHARGER

CLEARED

Oct 18, 1994
426 days to decision

K934016 · Product code: **DTE** · Cardiovascular
Source: https://www.510kdatabase.net/k934016/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Aug 18, 1993
Decision date	Oct 18, 1994
Days to decision	426 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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