

**K950856 ZOLL PD2000 PACEMAKER/ADVISORY
DEFIBRILLATOR, MODEL PD2000, ZOLL D2000 ADVISORY
DEFIBRILLATOR, MODEL D2000**Dec 29, 1995
305 days to decisionK950856 · Product code: **DRO** · Cardiovascular
Source: <https://www.510kdatabase.net/k950856/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Feb 27, 1995
Decision date	Dec 29, 1995
Days to decision	305 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k950856/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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