FDA 510(k) Premarket Notification

K872368 MODIFIED VERSA STIM ELECTRODE LINE

CLEARED

Sep 18, 1987
92 days to decision

K872368 · Product code: **DRO** · Cardiovascular
Source: https://www.510kdatabase.net/k872368/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Jun 18, 1987
Decision date	Sep 18, 1987
Days to decision	92 days
Third-party review	No

APPLICANT

Company	**Conmedcorp**
Location	Dayton, OH, US
Contact	WILLIAM W ABRAHAM
510(k) history	92 submissions · 92 cleared · 1981-2010