FDA 510(k) Premarket Notification

K880432 VICOR PACER, MODEL 410A

CLEARED

Feb 22, 1988
25 days to decision

K880432 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k880432/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jan 28, 1988
Decision date	Feb 22, 1988
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Tpl-Cordis, Inc.**
Location	Suffield, CT, US
Contact	WILLIAM C NEALON
510(k) history	6 submissions · 6 cleared · 1987-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k880432/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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