FDA 510(k) Premarket Notification

K881711 VICOR MODELS 410B & 410D AND MODIFIED 410A PACEMA.

CLEARED

May 26, 1988
36 days to decision

K881711 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k881711/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Apr 20, 1988
Decision date	May 26, 1988
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Telectronics and Cordis Pacing Systems**
Location	Miami, FL, US
Contact	DUANE A SCHULTZ
510(k) history	3 submissions · 3 cleared · 1988-1988