

**K881711 VICOR MODELS 410B & 410D AND MODIFIED 410A  
PACEMA.**May 26, 1988  
36 days to decisionK881711 · Product code: **DXY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k881711/>**SUBMISSION DETAILS**

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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**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881711/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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