

**K873158 MODIFIED MULTICOR AND OMNI-STANICOR  
PACEMAKERS**Jan 7, 1988  
150 days to decisionK873158 · Product code: **DXY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k873158/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 10, 1987
Decision date	Jan 7, 1988
Days to decision	150 days
Third-party review	No

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k873158/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 7, 2026