

K880898 MULTICOR AND PACEMAKERSMay 6, 1988
70 days to decisionK880898 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k880898/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Feb 26, 1988
Decision date	May 6, 1988
Days to decision	70 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880898/>; Data sourced from FDA 510(k) public records (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. - Generated July 7, 2026