

K810677 MODELS 5976/5977 PACEMAKER PULSE GENERMay 27, 1981
75 days to decisionK810677 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k810677/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Mar 13, 1981
Decision date	May 27, 1981
Days to decision	75 days
Third-party review	No

APPLICANT

Company	Medtronic, Inc.
Location	Mounds View, MN, US
Contact	Rachel Brucker
Website	https://www.medtronic.com
510(k) history	209 submissions · 208 cleared · 1981-2026

Medtronic, Inc. is a global medical device manufacturer headquartered in Mounds View, United States. The company develops and markets a broad range of medical devices across multiple therapeutic areas. Medtronic maintains a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions since 1981. The company specializes primarily in Cardiovascular devices, which represent 82% of its submission portfolio. Recent clearances include coronary perfusion cannulae, intracoronary shunts, venous cannulae, guidewires, deflectable catheter systems,...

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