

K791417 IMPLANTABLE PULSE GENERATOR PACEMAKERSep 4, 1979
39 days to decisionK791417 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k791417/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jul 27, 1979
Decision date	Sep 4, 1979
Days to decision	39 days
Third-party review	No

APPLICANT

Company	Intermedics, Inc.
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996

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Device record: <https://www.510kdatabase.net/k791417/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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