

K802040 BATTERY FOR #229-01 PULSE GENERATORSep 9, 1980
29 days to decisionK802040 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k802040/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 11, 1980
Decision date	Sep 9, 1980
Days to decision	29 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/k802040/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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