

K831859 PACEMAKER PROGRAMMER #2032Oct 27, 1983
139 days to decisionK831859 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k831859/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jun 10, 1983
Decision date	Oct 27, 1983
Days to decision	139 days
Third-party review	No

APPLICANT

Company	Vitatron Medical BV
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1976-1986

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Device record: <https://www.510kdatabase.net/k831859/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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