

**K760670 PACEMAKERS, VITATRONS (SERIES 2000)**Nov 12, 1976  
56 days to decisionK760670 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k760670/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Sep 17, 1976
Decision date	Nov 12, 1976
Days to decision	56 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k760670/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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