

**K791147 ARCO PROGRAMMABLE PACEMAKER SYSTEM**Aug 3, 1979  
46 days to decisionK791147 · Product code: **DXY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k791147/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jun 18, 1979
Decision date	Aug 3, 1979
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Arco Medical Products Co.</b>
Location	Walker, MI, US
510(k) history	21 submissions · 20 cleared · 1976-1980

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Device record: <https://www.510kdatabase.net/k791147/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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