

**K810410 STERILE PACKAGING**Mar 5, 1981  
15 days to decisionK810410 · Product code: **DXY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k810410/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Feb 18, 1981
Decision date	Mar 5, 1981
Days to decision	15 days
Third-party review	No

**APPLICANT**

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Company	<b>Telectronics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990

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

Device record: <https://www.510kdatabase.net/k810410/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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