

**K840748 LITHIUM PACEMAKER 811-811L & 811XL**Aug 3, 1984  
162 days to decisionK840748 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k840748/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Feb 23, 1984
Decision date	Aug 3, 1984
Days to decision	162 days
Third-party review	No

**APPLICANT**

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Company	<b>Omikron Scientific, Ltd.</b>
Location	Walker, MI, US
510(k) history	4 submissions · 3 cleared · 1984-1991

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

Device record: <https://www.510kdatabase.net/k840748/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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