

**K841232 PULSE GENERATOR 696, 696B & 696L**Dec 3, 1984  
256 days to decisionK841232 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k841232/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Mar 22, 1984
Decision date	Dec 3, 1984
Days to decision	256 days
Third-party review	No

**APPLICANT**

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Company	<b>Siemens Elema AB</b>
Location	Mchenry, IL, US
510(k) history	63 submissions · 60 cleared · 1978-2003

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

Device record: <https://www.510kdatabase.net/k841232/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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