

**K820657 PULSE GENERATOR 678 B**Apr 15, 1982  
37 days to decisionK820657 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k820657/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Mar 9, 1982
Decision date	Apr 15, 1982
Days to decision	37 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/k820657/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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