

**K792223 PACESETTER MODEL 350**Nov 30, 1979  
25 days to decisionK792223 · Product code: **DTE** · CardiovascularSource: <https://www.510kdatabase.net/k792223/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Nov 5, 1979
Decision date	Nov 30, 1979
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Cynergy</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1978-1981

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

Device record: <https://www.510kdatabase.net/k792223/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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