

**K780717 EXTERNAL TEST PULSE GENERATOR**Jun 30, 1978  
64 days to decisionK780717 · Product code: **DTC** · CardiovascularSource: <https://www.510kdatabase.net/k780717/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function (DTC)
Date received	Apr 27, 1978
Decision date	Jun 30, 1978
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitatron Medical BV</b>
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1976-1986

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

Device record: <https://www.510kdatabase.net/k780717/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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