

**K900451 VITALMAX 800 NON-INVASIVE VITAL SIGN  
MONITOR/PULSE**Jul 18, 1990  
169 days to decisionK900451 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k900451/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jan 30, 1990
Decision date	Jul 18, 1990
Days to decision	169 days
Third-party review	No

**APPLICANT**

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Company	<b>Pace Tech, Inc.</b>
Location	Clearwater, FL, US
Contact	MARY J BILGUTAY
510(k) history	18 submissions · 18 cleared · 1988-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900451/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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