

**K933624 VITALMAX 4000**Jul 19, 1994  
365 days to decisionK933624 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k933624/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jul 19, 1993
Decision date	Jul 19, 1994
Days to decision	365 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933624/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026