

**K901612 VITALMAX 2000 ECG MONITOR W/NON-INVASIVE
BLOOD-PRE**Apr 30, 1990
24 days to decisionK901612 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k901612/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Apr 6, 1990
Decision date	Apr 30, 1990
Days to decision	24 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k901612/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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