

K914164 VITALMAX 840 SERIESApr 9, 1993
570 days to decisionK914164 · Product code: **BXN** · Anesthesiology
Source: <https://www.510kdatabase.net/k914164/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stimulator, Nerve, Battery-powered (BXN) |
| Date received | Sep 17, 1991 |
| Decision date | Apr 9, 1993 |
| Days to decision | 570 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---|
| Company | Pace Tech Medical Monitors, Inc. |
| Location | Clearwater, FL, US |
| 510(k) history | 3 submissions · 3 cleared · 1993-1998 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914164/>; Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026