FDA 510(k) Premarket Notification

K914164 VITALMAX 840 SERIES

CLEARED

Apr 9, 1993
570 days to decision

K914164 · Product code: **BXN** · Anesthesiology
Source: https://www.510kdatabase.net/k914164/

SUBMISSION DETAILS

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