FDA 510(k) Premarket Notification

K913263 VITALMAX 830 SERIES

CLEARED

Oct 10, 1991
79 days to decision

K913263 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k913263/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jul 23, 1991
Decision date	Oct 10, 1991
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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