FDA 510(k) Premarket Notification

CLEARED

K170973 VitalWatch Software User Interface

Jun 2, 2017
60 days to decision

K170973 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k170973/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Apr 3, 2017
Decision date	Jun 2, 2017
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vitalconnect, Inc.**
Location	Campbell, CA, US
Contact	Kevin Potgieter
510(k) history	10 submissions · 10 cleared · 2014-2025