FDA 510(k) Premarket Notification

K171029 eCareCoordinator

CLEARED

Jul 14, 2017
99 days to decision

K171029 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k171029/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Apr 6, 2017
Decision date	Jul 14, 2017
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Visicu, Inc.**
Location	Lake Forest, CA, US
Contact	Daniel R. Plonski
510(k) history	7 submissions · 7 cleared · 2000-2022