FDA 510(k) Premarket Notification

CLEARED

K211822 LifePath Remote Patient Monitoring Platform

Feb 3, 2022
234 days to decision

K211822 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k211822/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jun 14, 2021
Decision date	Feb 3, 2022
Days to decision	234 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Elastic Care Incorporated**
Location	Markham, CA
Contact	Ashok Kalle
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	**Regulatory Technology Services, LLC**
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov