FDA 510(k) Premarket Notification

K140933 ALIVECOR HEART MONITOR

CLEARED

Aug 15, 2014
126 days to decision

K140933 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k140933/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Apr 11, 2014
Decision date	Aug 15, 2014
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**AliveCor, Inc.**
Location	San Francisco, CA, US
Contact	Albert Boniske
510(k) history	19 submissions · 19 cleared · 2012-2026