FDA 510(k) Premarket Notification

K172778 CheckMyHeart Plus

CLEARED

Jul 5, 2018
294 days to decision

K172778 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k172778/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Sep 14, 2017
Decision date	Jul 5, 2018
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dailycare Biomedical, Inc.**
Location	Flagstaff, AZ, US
Contact	Ming Da Lee
510(k) history	5 submissions · 5 cleared · 2005-2018