

HHS Public Access

Transl J Am Coll Sports Med. Author manuscript; available in PMC 2023 October 12.

Published in final edited form as:

Author manuscript

Transl J Am Coll Sports Med. 2022; 7(4): 1-7. doi:10.1249/tjx.00000000000215.

Systematic Review of Fitbit Charge 2 Validation Studies for Exercise Tracking

Crista Irwin¹, Rebecca Gary¹

¹Nell Hodgson Woodruff School of Nursing, Emory University, Atlanta, GA

Abstract

Context—There are research-grade devices that have been validated to measure either heart rate (HR) by electrocardiography (ECG) with a Polar chest strap, or step count with ACTiGraph accelerometer. However, wearable activity trackers that measure HR and steps concurrently have been tested against research-grade accelerometers and HR monitors with conflicting results. This review examines validation studies of the Fitbit Charge 2 (FBC2) for accuracy in measuring HR and step count and evaluates the device's reliability for use by researchers and clinicians.

Design—This registered review was conducted using the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. The robvis (risk-of-bias visualization) tool was used to assess the strength of each considered article.

Eligibility Criteria—Eligible articles published between 2018 and 2019 were identified using PubMed, CINHAL, Embase, Cochran, and World of Science databases and hand-searches. All articles were HR and/or step count validation studies for the FBC2 in adult ambulatory populations.

Study Selection—Eight articles were examined in accordance with the eligibility criteria alignment and agreement among the authors and research librarian.

Main Outcome Measures—Concordance correlation coefficients (CCC) were used to measure agreement between the tracker and criterion devices. Mean absolute percent error (MAPE) was used to average the individual absolute percent errors.

Results—Studies that measured CCC found agreement between the FBC2 and criterion devices ranged between 26% and 92% for HR monitoring, decreasing in accuracy as exercise intensity increased. Inversely, CCC increased from 38% to 99% for step count when exercise intensity increased. HR error between MAPE was 9.21% to 68% and showed more error as exercise intensity increased. Step measurement error MAPE was 12% for healthy persons aged 24–72 years but was reported at 46% in an older population with heart failure.

Corresponding author: Crista Irwin, MS, BSN, RN, cevan01@emory.edu, Nell Hodgson Woodruff School of Nursing, Emory University, 1520 Clifton Rd., Atlanta, GA 30322.

Conflicts of Interest and Sources of Funding

The authors of this project have no conflicts of interest to declare. This project was partially supported by the T32 training grant in Interventions to Improve Outcomes in Chronic Conditions (T32NR012715–06, Dunbar S, Song M (co-PIs)), and by the Healing Hearts and Mending Minds "Fitbrain" study (R-01 NR014963, Gary R and Waldrop D (co-PIs)).

Conclusions—Relative agreement with criterion and low-to-moderate MAPE were consistent in most studies reviewed and support validation of the FBC2 to accurately measure HR at low or moderate exercise intensities. However, more investigation controlling testing and measurement congruency is needed to validate step capabilities. The literature supports the validity of the FBC2 to accurately monitor HR, but for step count is inconclusive so the device may not be suitable for recommended use in all populations.

Keywords

Fitbit Charge 2; exercise tracker; heart rate; step count

Introduction

Background

Current universal guidelines recommend at least 150 minutes of weekly moderate intensity exercise of between 3–6 metabolic equivalent of task (MET) for persons across the health strata (1). Moderate intensity exercise can be achieved by brisk walking at a pace of 100 steps per minute or measured at 60–70% of heart rate (HR) maximum (2). Persons with chronic conditions who routinely participate in physical activity at moderate intensity for a minimum of 150 minutes per week exhibit significant improvements in cardiovascular function and reduction in inflammation (1,3–12). Incorporating exercise prescriptions into healthcare protocols show major benefits among a variety of health conditions including cardiovascular disease, depression, and HIV, as well as for preventative medicine (13). Providers need cost-effective, reliable wearable devices that accurately measure physical activity to assess and provide feedback to patients regarding their physical activity and to fully utilize the benefits of an exercise prescription.

Current research-grade devices have been validated to measure either HR using electrocardiography (ECG) (14), step count using an ACTigraph accelerometer (15), or sensing HR pulse using a Polar chest strap (16), but to date no wearable device has been validated to measure both HR and steps concurrently. Exercise equipment manufacturers continuously develop multifunction activity trackers worn as watches that purport to accurately capture steps and HR; however, these wearable activity trackers have been tested against research-grade accelerometers and heart rate monitors with variable results. The Fitbit Charge 2 (FBC2) (Fitbit Inc., San Francisco, CA, USA) is a low-cost, wearable device that tracks steps and HR and has been studied in a variety of populations in home-based and healthcare settings. The purpose of this review is to examine validation studies of the FBC2 for accurately measuring HR and step count and to evaluate the device's reliability to determine whether the device can be recommended by healthcare providers for use by patients.

Features of the Fitbit Charge 2

Fitbit trackers use microelectronic triaxial accelerometer and proprietary algorithms to measure step gait and distance and continuous light-emitting diode (LED) lighting to measure pulse continually. They are multifunctional, wrist-worn devices that not only measure steps and HR but include a multitude of user-friendly features. The device must

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be wirelessly connected via Bluetooth to a network-connected mobile phone. Through this connectivity, the device may receive text and call notifications. The device may be connected to the owner's contact list to develop community support networks for exercise motivation with special permission. The device software often sends supportive messages to encourage movement throughout the day or once the owner achieves personal activity goals set by him/herself. The software package also includes workout videos that can be accessed on the mobile phone application. The device also functions as a watch and has timer, mileage, relaxation, and stopwatch features.

Limitations of the Fitbit Charge 2

Because the device requires smart-phone and internet access and has an average price of \$150, the FBC2 may be a difficult option for low-income populations. Many basic and low-functioning mobile phones lack the capability to support the Fitbit application. The watch is rechargeable and includes a charging cord, which is easily misplaced and/or broken, and needs 6–7 hours charging time which lasts for approximately 3–4 days. Consumers report frequent watchband and equipment failure after 12–18 months of use. Additionally, the manufacturer does not report whether the updated models, which are released about every 18 months, have been altered significantly and, therefore, require updated testing and validation.

Methods

Design

This review was registered on the International Prospective Register of Systematic Reviews (PROSPERO) and the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines were used in conducting the review and reporting the appropriate articles (17). Figure 1 illustrates the PRISMA flow diagram of included articles. Eight articles on the validation of FBC2 in adult ambulatory populations that were published between 2018 and 2019 were examined for this review. Seven of the articles were randomized control trials and one was a test of a single participant who wore multiple trackers. Articles were excluded if they did not assess the FBC2 model, did not assess HR or step counts, or if the population sample was under 18 years of age. The relatively small article publication time range is due in part to the speed at which commercially available wearable exercise tracker technology changes.

Strategy

The methods for this review included a search of PubMed, the Cumulative Index to Nursing and Allied Health Literature (CINAHL), Cochran, Embase, World of Science, hand searches, and assistance by Emory University's Health Sciences Librarian. Key words and combinations of the words used in the search were "Fitbit Charge 2," "exercise tracker," "activity tracker," "activity monitor," "heart rate," "steps," and "validation." Article inclusion criteria were HR and/or steps validation studies for the FBC2 in adult ambulatory populations.

Data Extraction

Each validation article included in this review was required to assess HR and step count accuracy. Articles were also included if they evaluated intra-reliability testing between the FBC2 trackers used in the testing as well as inter-reliability among other trackers assessed. Criterion measures for HR were based on the research-validated ECG and the Polar chest strap for ambulatory activities (14,15). Step count data was compared to the validated ACTiGraph wGT3X-BT (16).

To assess the tracker's accuracy, seven of the eight studies in this review explained the differences in the data collected by the FBC2 using the following measures (Table1): mean error, as in the difference between the criterion measure and the consumer device; mean absolute error, as in the average absolute distance between the data from the consumer and the criterion devices; mean percent error or relative error rate, as in the difference between the criterion measure and the consumer device, represented by a percentage; and mean absolute percent error (MAPE), as in the average of the individual absolute percent errors. MAPE analyzes individual overestimation and underestimation values taken by the device and, therefore, may be a more appropriate representation of the activity monitors when comparing studies.

Concordance correlation coefficient (CCC) 95% confidence intervals (CI) were used to describe strength of agreement between the devices in four studies (18–20). Interclass correlation coefficient (ICC) is used when comparing multiple devices to each other and to the criterion. Additionally, because precise data congruency collected between consumer devices is unlikely, Bland-Altman (BA) analysis was used by four studies to evaluate proximity to data measured by criterion devices (18,20,23,24).

Another article was included in this validation review because the authors tested the FBC2 tracker's accuracy for measuring cardiorespiratory fitness compared to maximal oxygen uptake (VO_{2max}) (25). Cardiorespiratory fitness (CRF), defined as the circulatory and respiratory systems' transport and utilization of oxygen to the skeletal muscles, is typically measured by maximal graded exercise testing on a treadmill and measured in units of mL·kg⁻¹·min⁻¹. Under strict laboratory protocols by a trained exercise physiologist using precise equipment, the gold standard for evaluating CRF levels is by VO_{2max} (26). Researchers have found that low levels of CRF measured by VO_{2max} treadmill testing have been associated with cardiovascular disease risk (25,27,28). The FBC2 purports to evaluate CRF using proprietary algorithms which include an individual user's age, weight, height, resting HR, and peak HR.

Articles were evaluated by two independent researchers and inclusion agreement was discussed in detail. Validation articles included in this review are listed and described in Tables 1 and 2.

Risk of bias assessment

Each study was evaluated for risk of bias (Table 2). Criteria used for the assessment included randomization bias, recruitment bias, protocol deviation bias including criterion tool bias, missing outcome bias, and reporting bias. The selected studies were consistently evaluated

as high quality based on these criteria with few concerns for risk of bias. Most of the concerns were regarding racially homogenous or small samples. One study included a single participant and most of the studies included majority white persons and healthy populations. The risk-of-bias visualization tool (robvis; Bristol, AC, UK) was used to create the risk of bias assessment (Fig. 2) (29).

Results

Accuracy testing

Of the eight studies found, five reported MAPE values of the FBC2 against the criterion for each study which are considered acceptable at <10%. The MAPE values for the studies evaluating HR accuracy include 9.21% (18), 10.79% (19), and 69% (24). Mean absolute percent error values investigating step count acuity were 12.36% (22), 46% for participants with heart failure (HF), and 12% among healthy controls (21). Another study measured the HR difference between the FBC2 and the criterion using relative error rate (RER) (light activity, 5.36%, to moderate activity, 9.3%) (20).

Intra-reliability testing

A 24-hour evaluation study found 91% CCC (95% CI 0.896, 0.914) agreement of the criterion with the FBC2 in a single participant (male, 29 years) (18). Another study cited 92% CCC (95% CI 0.92, 0.93) agreement of the criterion with the FBC2 in their randomized controlled trial including 20 participants (mean age 27.5 yr, standard deviation (SD) 6 yr; 55% female) while walking on a treadmill (19). Additionally, researchers collected data from 30 participants (mean age 23.5 yr, SD 3 yr; 50% female) and found that, as exercise intensity increased, agreement decreased (20). At very low HR, intensity ranging between 55 and 90 beats per minute (bpm), CCC agreement between the criterion and the FBC2 was 89% (95% CI 0.79, 0.95) (20). When HR ranged between 90 and 120 bpm, CCC agreement was moderate at 55%, (96% CI 0.28, 0.74) and CCC was poor at 26% (95% CI 0.01, 0.46) when HR ranged between 110 and 150 bpm (20). In contrast, step counting criterion reliability increased with treadmill speeds in a 3-day field study of 15 participants (mean age 65.5 yr, SD 12.6 yr; 40% female) with heart failure compared to 14 (mean age 43 yr, SD 18.9 yr; 64% female) healthy controls (21). CCC agreement of the step criterion with the FBC2 was 38% (95% CI 0.00, 0.67) at 2.4 km/h (slow walk), 82% (95% CI 0.68, 0.97) at 3.0 km/h (moderate walk), and 99% (95% CI 0.98, 1.0) at 3.6 km/h (brisk walk) (21).

Inter-reliability testing

In a 24-hour study of 20 (mean age 70.2 yr, SD 2.9 yr; 55% female) older healthy adults in Ireland, results showed >0.89 ICC strength of agreement between devices for step count evaluation (22). In addition, a study from Korea including 51 participants (mean age 44.4 yr, SD 16.6 yr; 53% male; 100% Asian) who were undergoing electrophysiological study and ablation to treat paroxysmal tachycardia or supraventricular tachycardia found >0.98 ICC strength of agreement between the FBC2 and the ECG for HR monitoring (23). Recorded baseline HR monitoring with the FBC2 was within \pm 5 beats per minute of the criterion ECG at 95% accuracy (23). However, device agreement of the FBC2 and criterion results by

Pearson correlations assessed in other studies in the review were incongruent measuring 0.23 (poor) and 0.94 (equivalent), respectively (19,24).

Cardiorespiratory fitness assessment

Researchers compared the VO_{2max} values obtained from the standard treadmill tests with the CRF values estimated by the FBC2 (25). In a sample of 65 healthy adults aged 18–45 yr (55% female), Bland-Altman analyses showed that the FBC2 CRF had a positive bias of 1.59 mL·kg⁻¹·min⁻¹ when compared to the treadmill testing at 15 s and a positive bias of 0.30 mL·kg⁻¹·min⁻¹ at 60 s with MAPE values <10% for each comparison (26).

Discussion

Heart rate validation

Four studies in this review assessed the FBC2 for HR accuracy validation. All except one study included healthy participants, aged 21–73 yr, and generally reported more accuracy at lower-intensity activity levels (18–23). Heart rate MAPE values while walking at low-to-moderate-intensity levels, at 9.21%, 10.79%, and 69%, reveal a wide interval of error results, with two of the three being similar (18,19,24). The RER reported by one group of researchers supports validation with their statistically moderate error rate at low-to-moderate walking intensity (light activity, 5.36%, to moderate activity, 9.3%) (20). Another study used pacing cycle length (PCL) data obtained during scheduled electrophysiological studies to evaluate the HR accuracy of the FBC2 (23). At 100 bpm, the FBC2 measured within \pm 5 bpm when compared to the ECG criterion at a rate of 93% accuracy with atrial pacing and 80% accuracy with ventricular pacing (23). However, the FBC2 device became significantly less accurate at higher bpm (23). HR and steps inherently fluctuate with intensity. These results are similar to the other studies reviewed.

Step count validation

In the Irish study of older adults, the FBC2 overestimated step count (MAPE 12.36%, approaching the acceptable range of <10%), but had vastly different results than the study comparing the older HF subjects (MAPE 46%) to younger healthy controls (MAPE 12%) (21,22). The explanation for why the MAPE values of the two healthy populations in the studies were similar while the MAPE values among the HF participants showed much higher error rates is unclear. However, alterations in gait and slower walking speed among the HF patients likely challenge the FBC2 to track steps reliably and may be a concern when using this device to track steps in populations with ambulation limitations or considerable exercise intolerance due to symptom severity.

FBC2 as HR monitor

Reliability results as determined by criterion agreement with the FBC2 reported in this review were markedly varied. Scores <0.50 indicate poor reliability, 0.50 to 0.75 moderate reliability, and >0.75 good reliability (30). Nelson et al. (18) and Reddy et al. (19) reported high CCC scores of >90%. However, Thomson et al. (20) showed decreasing reliability from 56% (moderate) to 26% (poor) as HR intensity increased. Pearson coefficient results from two studies revealed the widest reliability agreement strength discrepancy from

0.23 (weak) and 0.94 (equivalent) (19,24). Finally, Bland Altman analysis plots revealed HR underestimation measured by the FBC2 compared to criterion at all intensity levels (18,20,24). The differences of the results may be caused by erratic arm movements or misplacement of the tracker bands as the participants move and perspiration. These varied results make it difficult to reach a definitive conclusion regarding reliability across intensity levels, but support reliability at low-to-moderate exercise dose levels.

Fitbit Charge 2 as step counter

Reliability of the FBC2 is in agreement with the step criterion, which is the opposite of the HR results. CCC agreement increased from 38% at lower speeds to 99% at a brisk walk (21). ICC results of >0.89 supports evidence for high agreement strength between the FBC2 step counter and actigraphy (22).

Cardiovascular fitness validation

Researchers reported that the FBC2 could be validated to evaluate CRF in relatively young, healthy persons, especially those with a high level of fitness (26). Nearly 92% of the total participants in this study were classified as having high CRF (26). Although the study found CRF agreement between the FBC2 and the Balke treadmill test among users with lower fitness levels, the low numbers in the "good" or "poor" fitness-leveled groups sampled in the study do not provide sufficient evidence of variation to determine validity nor is the sample representative of the general population who are typically less engaged in CRF activities. Validation studies are needed in populations with chronic conditions or who have ambulation challenges to further evaluate the CRF feature in the FBC2.

Study limitations

Validation consensus of the FBC2 is limited due to the studies' small sample sizes (n=1 to n=60) and non-standardized activity settings with some conducted in laboratories on treadmills and others in free-living conditions. Most of the HR examinations were only conducted using young and healthy subjects and may not be generalizable to populations who have chronic conditions or to older adults with other physical limitations. The study that compared HF subjects to healthy controls included far different age demographics (21). In addition, the review is limited by the small number of relevant studies available within a short time span which is due in part to the development speed of new technology. The Fitbit company released the FBC2 in 2016 and the FBC3 became available in 2018. The cost and research effort needed to perpetually study and validate new technology limits the viability of commercial wearable devices such as the FBC2 for research and use in primary care. Developers of commercial devices would benefit monetarily from strategic collaborations with healthcare researchers in producing devices that are technologically consistent and reliable. There is great potential for wide use of more accessible and affordable devices by healthcare providers worldwide.

Conclusion

Although the FBC2 has been validated for moderate HR and step count accuracy in some studies, more investigation controlling testing and measurement congruency is needed to

validate both HR and step capabilities. The literature supports the validity of the FBC2 to accurately monitor HR at low-to-moderate exercise intensities, but validation for step count is inconclusive and may not be suitable for recommended use by populations with gait speed or ambulation challenges.

Acknowledgements

The authors would like to acknowledge Sharon Leslie, the librarian for the Nell Hodgson Woodruff School of Nursing at Emory University, for her guidance in devising a strategic plan for this review. We also appreciate the support of Jeannine Cimiotti for her editorial and educational contributions. The results of the study do not constitute endorsement by ACSM.

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Low

				Risk o	of bias do	mains	-		
		D1	D1b	D2	D3	D4	D5	Overall	
	Hwang J, et at. (2019)	+	-	+	+	+	+	+	
	Klepin K, et at. (2019)	+	-	+	+	+	+	+	
	Nelson BW, et al (2019)	+	X	+	+	-	+	-	
Study	O'Driscoll R, et al. (2019)	+	-	+	+	+	+	+	
	Reddy RK, et al. (2018)	+	-	+	+	+	+	+	
	Tedesco S, et al (2019)	+	-	+	+	+	+	+	
	Thomson EA, et al (2019)	+	-	+	+	+	+	+	
	Vetrovsky T, et al. (2019)	+	+	+	+	+	+	+	
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- relation to timing of randomization. D2 : Bias due to deviations from intended intervention. D3 : Bias due to missing outcome data.
- D4 : Bias in measurement of the outcome.
- D5 : Bias in selection of the reported result.



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Table 1:

Summary of measurements

Study author/year	Criterion measure	HR	Steps	ccc	ICC	PC	BA	ME	MAE	MPE/RER	MAPE	Risk of Bias
Hwang J, et at. (2019)	ECG	Х			х		Х					TOW
Klepin K, et at. (2019)	VO _{2max}	х			х						Х	SOME CONCERNS
Nelson BW, et al (2019)	ECG	Х		х			Х		Х		Х	SOME CONCERNS
O'Driscoll R, et al. (2019)	Polar chest strap	Х				х	Х		Х		Х	TOW
Reddy RK, et al. (2018)	Polar chest strap	х		х		Х		х		Х	Х	TOW
Tedesco S, et al (2019)	ActiGraph wGT3X-BT and New-Lifestyles NL2000i		х		х			х		Х	Х	TOW
Thomson EA, et al (2019)	ECG	х		х			Х			Х		TOW
Vetrovsky T, et al. (2019)	ActiGraph wGT3X-BT		х	х							Х	TOW

BA, Bland-Altman analysis; CCC, concordance correlation coefficients; ECG, electrocardiography; ICC, interclass correlation; MAE, mean absolute error; MAPE, mean absolute percent error; ME, mean error; MPE/RER, mean percent error or relative error rate; PC, Pearson correlation; VO2max, maximal oxygen uptake.

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Table 2.

Summary of study findings

on/methods Summary of findinos	nts with history of paroxysmal iVT) were undergoing electrophysiological SVT and post ablation of SVT were	Ithy adults; study duration: 1-week; 3 GPS FBC2 60-second cardiorespiratory fitness I were worn continuously (CRF): highly associated with VO _{2max} ; MAPE 9.14% 9.14%	bject, healthy adult male, used to minimize FBC2 mean difference –3.47 bpm compared to ollected across sedentary, walking, CCC 91% compared to ECG over 24 hours; MAPE 9.21% walking	p; activities evaluated in a 1-day laboratory FBC2 HR = CM, but not consistent across sycling, mimicked ADLs, and sedentary activity levels; MAPE 31% walking incline; n	p; 2-day data collection; all healthy adults; FBC2 >error high intensity exercise; MAPE exercise, interval training, and ADLs 10.79%	hour free-living data collection: all older, All devices highly correlated (ICC>0.89), cluded a range of moderate to vigorous FBC2 overcounted steps, MAPE 12.36%	nents taken at intervals: rest (3 min), FBC2 error rate (3.9–13.5%). RER per activity level: light (5.36%), moderate (9.20%), overy.	main purpose was to evaluate step FBC2 healthy participants MAPE 12%; HF study mAPE 46%; >correlation low speeds on treadmill
Design/me	pared FBC2 to ECG; participants wit tventricular tachyarrhythmia (SVT) w r; baseline HR during induced SVT au ured	pared FBC2 to VO _{2max} ; all healthy ac ed 15-minute outdoor runs and were	pared FBC2 to ECG; single subject, I ble differences; 24-hour data collecte ing, activities of daily living (ADLs),	pared FBC2 to Polar chest strap; actives included running, walking, cycling itions with rest periods between	pared FBC2 to Polar chest strap: 2-da _{nax} testing included resistance exercis itions	pared FBC2 to ActiGraph; 24-hour fr hy adults; activities assessed included ing and sleeping conditions	pared FBC2 to ECG; measurements t ling (2 min), treadmill (every 3 min w ase until volitional fatigue), recovery.	pared FBC2 to accelerometer; main pracy of activity trackers in persons wi
Samule nonulation	N=51, mean age 44.4 (SD 16.6) yr, n=24 Com female, Korean study	N=60, mean age 31 (SD 7.3) yr, n=33 Com female, race not reported track	N=1, age 29 yr, white male Com varia runn	N=59, mean age 44.2 (SD 14.1) yr, n=41 Com female, European cond	$ \begin{array}{c} N=20, \mbox{ mean age } 27.5 \mbox{ (SD } 6.0) \mbox{ yr}, \mbox{ n=11} \\ \mbox{ females, n=17 } (85\%) \mbox{ white} \\ \mbox{ cond} \end{array} $	N=20, mean age 70.2 (SD 2.9) yr, n=11 Com female, all white (English/Irish) walk	N=30, mean age 23.5 (SD 3) yr, n=15 Com female, race not reported incre	Healthy control participants: n=15, mean Com age 65.5 (SD 12.6) yr, n=6 female. accu HF field-based study: n=14. mean age
Study author/year	Hwang J, et at. (2019)	Klepin K, et at. (2019)	Nelson BW, et al (2019)	O'Driscoll R, et al. (2019)	Reddy RK, et al. (2018)	Tedesco S, et al (2019)	Thomson EA, et al (2019)	Vetrovsky T, et al. (2019)