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Original Research

Can Commercially Available Pedometers Be Used For Physical Activity Monitoring In Patients With COPD Following Exacerbations?

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Abstract

Background: Commercially available pedometers have been used as tools to measure endpoints in studies evaluating physical activity promotion programs. However, their accuracy in patients recovering from COPD exacerbations is unknown. The objectives of this study were to 1) assess the relative accuracy of different commercially available pedometers in healthy volunteers and 2) evaluate the accuracy of the top-performing commercially available pedometer in patients recovering from COPD exacerbations following hospital discharge.

Methods: Twelve healthy volunteers wore 2 pedometers, 2 smartphones with pedometer apps and an accelerometer for 15 minutes of indoor activity. The top-performing device in healthy volunteers was evaluated in 4 patients recovering from COPD exacerbations following hospital discharge during 6 minutes of walking performed at home. Bland-Altman plots were employed to evaluate accuracy of each device compared with direct observation (the reference standard).

Results: In healthy volunteers, the mean percent error compared to direct observation of the various devices ranged from -49% to +1%. The mean percent error [95% confidence interval (CI)] of the top-performing device in healthy volunteers, the Fitbit Zip[®], was +1% [-33 to +35%], significantly lower than that of the accelerometer (-13% [-56 to +29%], $p=0.01$). The mean percent error [95% CI] for the Fitbit Zip[®] in patients recovering from COPD exacerbations was -3% [-7 to +12%].

Conclusions: The accuracy of commercially available pedometers in healthy volunteers is highly variable. The top-performing pedometer in our study, the Fitbit Zip[®], accurately measures step counts in both healthy volunteers and patients recovering from COPD exacerbations.

Abbreviations: chronic obstructive pulmonary disease, **COPD**; body mass index, **BMI**; forced expiratory volume in 1 second, **FEV₁**; 6-minute walk test, **6MWT**; standard deviation, **SD**; confidence interval, **CI**; modified Medical Research Council, **mMRC**; vector magnitude units, **VMU**
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Introduction

Deconditioning is common in patients with chronic obstructive pulmonary disease (COPD) exacerbations and leads to excessive fatigue, respiratory symptoms with exertion and reduction in physical activity, which together result in further deconditioning.^{1,2} Limited physical activity is also associated with decreased quality of life, more severe dyspnea and increased risk of exacerbations, hospitalizations (including 30-day rehospitalizations), and death.³⁻⁹ Patients recovering from COPD exacerbations have been shown to have even lower levels of physical activity compared to patients with stable COPD.^{10,11} Clinical trials have demonstrated that pulmonary rehabilitation programs in clinical settings that include early mobilization to increase physical activity improve functional status and clinical outcomes following COPD exacerbations.¹² However, pulmonary rehabilitation programs in hospitals or ambulatory care settings are often inaccessible to patients following COPD exacerbations.¹³ Moreover, there is a need for validated tools to promote physical activity that can be used outside of pulmonary rehabilitation and other clinical settings.

Pedometers have been shown to be effective tools in promoting physical activity in healthy adults when combined with goal setting, feedback and self-management education.¹⁴ Similar programs using commercially-available pedometers designed for patients with stable COPD have also resulted in increased levels of physical activity and improved quality of life.^{15,16} However, physical activity promotion programs have not been extensively evaluated in patients recovering from COPD exacerbations, a population at particularly high risk of adverse outcomes.^{17,18}

The accuracy of commercially available pedometers (including smartphone applications: *apps*) in relation to accelerometers has not been evaluated in comparison

with direct observation in either healthy adults or patients recovering from COPD exacerbations. As patients with COPD may have differences in gait, including decreased walk intensity, cadence and speed, compared with healthy adults,¹⁹ it is crucial that these commercially available pedometers be evaluated for accuracy before being used in physical activity promotion programs.

The objectives of this study were two-fold: 1) to compare, in healthy volunteers, the relative accuracy of commercially available pedometers to both direct observation and an accelerometer commonly used in clinical research, and 2) to evaluate the accuracy of the top-performing commercially available pedometer in patients recovering from COPD exacerbations. The results of this study could aid in selecting a device with sufficient accuracy for objectively monitoring physical activity in patients recovering from COPD exacerbations.

Methods

We conducted 2 sub-studies, each addressing one of the study objectives.

Sub-study 1:

We sought to assess the validity of different pedometers in healthy volunteers during indoor activities over 15 minutes, compared with direct observation. The results of this sub-study were intended to guide the selection of a commercially available device for sub-study 2. We did not provide specific instructions for physical activity to these healthy volunteers. A convenience sample of 12 healthy non-smoking adults (age 18 years or older) was recruited for this sub-study. Age and gender of the volunteers were recorded.

We selected a sample of low-cost (<\$150) wireless-enabled tri-axial (motion captured in the horizontal, lateral and vertical axes) pedometers, and smartphone-based apps that take advantage of tri-axial accelerometers built into modern smartphones (Table 1). We also measured physical activity using an accelerometer commonly used in clinical research (Actigraph® wGT3X-BT [ActiGraph, Pensacola, Florida]).²⁰ Retail prices of the various devices were obtained from online sources.²⁰⁻²⁴ The pedometers and accelerometer were used according to the manufacturers' recommendations and instructions. After the investigator attached the devices, each healthy volunteer was asked to carry out their usual activities

Table 1. Device Characteristics

Class	Device	Output	Position Worn	Retail Price
Smartphone Apps	Runtastic®	Step count, energy expenditure, distance walked	Pocket	\$0
	Moves®	Step count, energy expenditure, distance walked	Pocket	\$0
Pedometers	Fitbit Zip®	Step count, energy expenditure, distance walked	Hip	\$60
	Fitbit Force®	Step count, energy expenditure, distance walked, stairs climbed	Wrist	\$129
Accelerometer	Actigraph®	Step count, VMU, energy expenditure, body position	Hip / Wrist	\$225 + \$1495 for analysis software

VMU=vector magnitude units

indoors while under direct observation by a trained research assistant. Smartphones with pedometer apps (Moves® and Runtastic®) were carried in the front pants pockets, the Actigraph® and Fitbit Zip® were attached to the participant's waist, and the Fitbit Force® was worn on the participant's non-dominant wrist. This field study was conducted indoors to approximate the type of physical activity that may be observed in patients recovering from COPD exacerbations.⁵

Sub-study 2:

In sub-study 2 we used the device with the lowest error compared to direct observation established in sub-study 1. Sub-study 2 took place in the homes of patients recovering from COPD exacerbations within one month of hospital discharge. Patients were eligible to participate if they had a physician diagnosis of COPD exacerbation, were able to walk unaided (e.g., without a walker or cane) and provided written informed consent. Participants were instructed to walk for up to 6 minutes

to mimic the duration of activity promotion used in previous studies.^{25,26} We also collected age, gender, weight and height (for calculating body mass index [BMI]; kg/m²), post-bronchodilator percent predicted forced expiratory volume in 1 second (FEV₁), time walked during a 6-Minute Walk Test (6MWT, minutes), and resting Borg dyspnea and fatigue scores (0-10, higher scores indicate more severe dyspnea and fatigue, respectively).²⁷⁻²⁹ The study was approved by the institutional review board at the University of Illinois at Chicago.

Analyses

For both sub-studies, directly observed step counts were the reference standard. Descriptive statistics employed proportions, mean (standard deviations [SD]), or median (range), as appropriate. Bland-Altman analyses were performed to calculate the mean percent error (reference standard-calculated/reference standard*100%) and 95% confidence interval (CI) between step counts calculated by each pedometer versus the reference standard.³⁰ Paired t-tests were also used to compare the error for each device with the error of the Actigraph®. All analyses were performed using SAS® (Cary, North Carolina).

Results

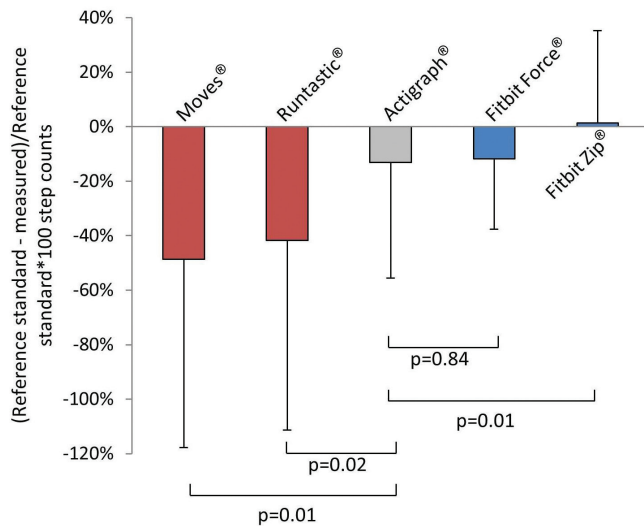
Sub-study 1

All 12 participants were under the age of 40 years and 7 were female. Assessed by direct observation, healthy volunteers walked a mean of 120 steps (SD = 53) over 15 minutes. The mean percent error for the various devices ranged from -49% to +1% (-62 to -2 steps) (Figure 1). The mean percent error [95% CI] of the accelerometer was -13% [-56 to +29%] or -20 steps [-62 to +21 steps]. The Fitbit Zip® had a significantly lower mean percent error (+1% [-33 to +35%]; -2 steps [-42 to +37 steps]) compared to the accelerometer ($p=0.01$). The Moves® and Runtastic® apps performed significantly worse than the accelerometer ($p<0.05$).

Sub-study 2

The 4 participants in this sub-study were men with a mean age of 69 years (SD=10) and mean BMI of 23kg/m² (SD=2). The mean % predicted post-bronchodilator FEV₁ for these participants was 33% (SD=5). The participants had a median resting Borg dyspnea and fatigue scores of 2 (range 0.5-4) and 1 (range 1-5), respectively. They walked a mean of 3.8 minutes

Figure 1. Relative Accuracy of the Different Commercially Available Pedometers



Bars represent standard deviation. Smartphones with pedometer apps (Moves® and Runtastic®) were carried in the front pants pockets, the Actigraph® and Fitbit Zip® were attached to the participant's waist, and the Fitbit Force® was worn on the participant's non-dominant wrist. Each of the healthy volunteers wore all devices concurrently.

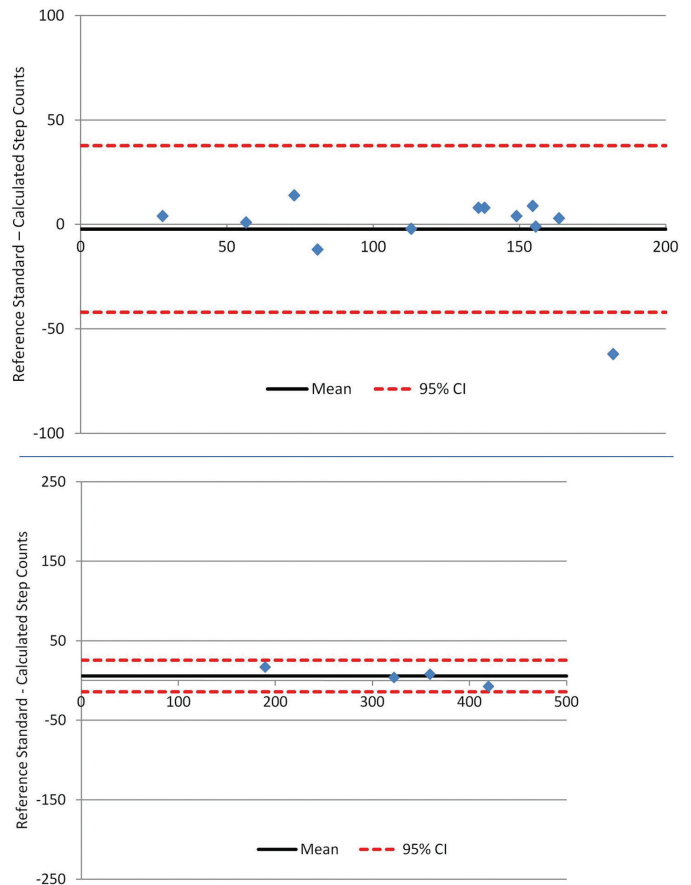
(SD=0.6), during which they walked a mean of 320 steps (SD=102) as measured by direct observation. The mean percent error (95% CI) for the Fitbit Zip® in this population was -3% (-7 to +12%) (Figure 2), or 6 steps (-14 to +25 steps).

Discussion

Our results indicate that commercially available pedometers (including pedometer apps) have variable levels of step count accuracy when compared to direct observation and an accelerometer. Interestingly, one of the pedometers (the Fitbit Zip®), outperformed an accelerometer commonly used in clinical research studies (Actigraph®). When compared with direct observation, the Fitbit Zip® performed exceedingly well in both healthy volunteers and patients recovering from COPD exacerbations. The increased accuracy of the Fitbit Zip® in patients recovering from COPD exacerbations compared to the healthy volunteers is related to the design of the pedometer's proprietary algorithm; however, there may be other reasons for the observed findings as well.

Although previous studies have examined the validity of accelerometers,³¹ commercially available pedometers,³²⁻³⁴ and even a pedometer smartphone

Figure 2. Bland-Altman Plot Showing Accuracy of the Fitbit Zip® Compared to Direct Observation in (A) Healthy Volunteers and (B) Patients Recovering from COPD Exacerbations



app,³⁵ to our knowledge this is the first study to evaluate them concurrently and in comparison with direct observation. Furthermore, while a previous study in patients with stable COPD evaluated energy expenditure estimates using a commercially available device (Fitbit Ultra® [Fitbit Inc, San Francisco, California]) compared to an accelerometer (SenseWear Armband® [BodyMedia Inc, Pittsburgh, Pennsylvania]),³⁴ we are not aware of any studies examining the accuracy of step count measurements in patients recovering from COPD exacerbations.

Our current study focused on step counts, as lower daily step counts in patients with COPD are associated with a range of adverse surrogate clinical outcomes, including elevated inflammatory biomarkers (C-reactive protein and IL-6), lower functional capacity (6-minute walk distance), more severe dyspnea (modified

Medical Research Council [mMRC] score) and, higher risk of COPD-related hospitalizations or all-cause mortality.^{7,36,37} While these commercially available pedometers also measure other markers of physical activity, including energy expenditure, daily distance walked and active time, these other measures are less well studied in terms of their relation with important clinical outcomes, such as rehospitalizations or all-cause mortality.

Our study has a number of limitations. First, healthy volunteers in sub-study 1 were only monitored indoors for a short period of time (15 minutes) and their activities were not standardized. However, the goal of the sub-study was to characterize the step count accuracy of different pedometers that would approximate the level of physical activity that would likely be seen in patients recovering from COPD exacerbations in sub-study 2.⁵ Indeed, in sub study 2, we found that patients recovering from COPD exacerbations have very low walk times (mean of 3.8 minutes) during a 6MWT. Additional studies are needed to evaluate the accuracy of the top-performing pedometers over longer periods of observation for better characterization of these devices. Second, in sub-study 2 we only enrolled a small sample of patients recovering from COPD exacerbations following hospital discharge. Additional studies with larger numbers of patients recovering from COPD exacerbations are needed to confirm our results in more diverse populations.

The findings of our study have several implications. The high level of accuracy of the low-cost Fitbit Zip® (\$60) we examined in sub-study 2 opens the door for the development and evaluation of home-based physical activity promotion programs with objective measures of physical activity in patients recovering from COPD exacerbations following hospital discharge. Such programs are needed as the majority of the studies to date have focused on physical activity promotion in individuals with stable COPD, highlighting the potential significance of our findings. The wireless connectivity feature of current devices, including the Fitbit Zip®, allows real-time monitoring of physical activity performance as part of a home-based exercise program. Newer generation pedometers now also include heart

rate monitors and even pulse oximeters highlighting the potential for the development of even more sophisticated physical activity promotion programs that monitor heart rate and oxygen saturation.^{38,39} Additionally, the results of our study suggest that some accelerometers (Actigraph®) may have worse performance characteristics for assessing step counts than commercially available pedometers. Together, our findings highlight the importance of validating the performance of physical activity measures before they are deployed in specific patient populations.

In conclusion, commercially available pedometers have variable levels of accuracy. The Fitbit Zip® is a low-cost commercially available pedometer that outperformed a commonly used accelerometer when compared to direct observation in healthy volunteers. Moreover, the Fitbit Zip® performed exceedingly well compared to direct observation in patients recovering from COPD exacerbations. The results of this study could be used to inform the selection of devices for objective monitoring of physical activity in patients recovering from COPD exacerbation following hospital discharge.

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Declaration of Interest

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