



Objective Measurement of Walking Activity Using Wearable Technologies in People with Parkinson Disease: A Systematic Review

Mathias Baptiste Correno ^{1,2,3}, Clint Hansen ^{3,*}, Thomas Carlin ^{1,2}, and Nicolas Vuillerme ^{1,2,4}

- ¹ Laboratory AGEIS, Université Grenoble Alpes, 38000 Grenoble, France; mathias.correno@gmail.com (M.B.C.); carlin.thom@gmail.com (T.C.); nicolas.vuillerme@univ-grenoble-alpes.fr (N.V.)
- ² LabCom Telecom4Health, Orange Labs, Université Grenoble Alpes, CNRS, Inria, Grenoble INP-UGA, 38000 Grenoble, France
- ³ Department of Neurology, Universitätsklinikum Schleswig-Holstein, 24105 Kiel, Germany
- ⁴ Institut Universitaire de France, 75231 Paris, France
- * Correspondence: c.hansen@neurologie.uni-kiel.de

Abstract: Parkinson's disease (PD) is a complex neurodegenerative disease with a multitude of disease variations including motor and non-motor symptoms. Quality of life and symptom management may be improved with physical activity. Due to technological advancement, development of small new wearable devices recently emerged and allowed objective measurement of walking activity in daily life. This review was specifically designed to synthesize literature on objective walking activity measurements using wearable devices of patients with PD. Inclusion criteria included patients with a diagnosis of PD and exclusion criteria included studies using animal models or mixed syndromes. Participants were not required to undergo any type of intervention and the studies must have reported at least one output that quantifies daily walking activity. Three databases were systematically searched with no limitation on publication date. Twenty-six studies were eligible and included in the systematic review. The most frequently used device was the ActiGraph GT3X which was used in 10 studies. Duration of monitoring presented a range from 8 h to one year. Nevertheless, 11 studies measured walking activity during a 7-day period. On-body sensor wearing location differed throughout the included studies showing eight positions, with the waist, ankle, and wrist being the most frequently used locations. The main procedures consisted of measurement of walking hours during a 2-day period or more, equipped with a triaxial accelerometer at the dominant hip or ankle. It is also important for further research to take care of different factors such as the population, their pathology, the period, and the environment.

Keywords: Parkinson's disease; walking activity; wearable devices; monitoring strategy; systematic review

1. Introduction

Neurodegenerative diseases such as Parkinson's disease (PD) can lead to motor [1,2] and non-motor symptoms [3]. The latter also often occur in in the general elderly population but people with PD show a stronger decline in a number of cognitive domains when compared to age-matched healthy adults (e.g., executive, attentional, and visuospatial domains) [4]. Motor symptoms on the other hand express themselves as bradykinesia, rigidity, tremor, and eventually even affect the ability to walk or maintain balance [1,5]. The most common motor-related deficits are gait disorders [6,7] which can lead to a loss of independence and increase the incidence of falls [8]. In addition, people with PD suffer from impaired functional abilities [9–11], based on a reduced level of strength [11,12] and lower physical activity levels [13,14].



Citation: Correno, M.B.; Hansen, C.; Carlin, T.; Vuillerme, N. Objective Measurement of Walking Activity Using Wearable Technologies in People with Parkinson Disease: A Systematic Review. *Sensors* 2022, 22, 4551. https://doi.org/10.3390/ s22124551

Academic Editor: Brett Fling

Received: 18 May 2022 Accepted: 9 June 2022 Published: 16 June 2022

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). This systematic review focuses on the use of wearable technology as a method to monitor the relationship between walking in PD and between clinical rating scales (e.g., MDS-UPDRS III) [15]. A better understanding and representation of solutions for instrumented monitoring of walking activity in PD could help clinicians and researchers when designing interventions and trials. Consequently, this review aimed to identify and map available studies on the use of wearable technology for objectively measuring walking in people with PD.

2. Materials and Methods

This review complies with the PRISMA guidelines to provide an evidence-based minimum set of items for reporting in systematic reviews [16]. The use of wearable technology to detect walking activity is explored to map the state of evidence and to identify potential research gaps [17]. The protocol of this current review has been registered in the PROSPERO (CRD42020210866) prospective register of systematic reviews and published in July 2021 [18].

Detailed information about the eligibility criteria, population, measurement tools, experimental procedures, measured outcomes, data sources and search strategy, study selection, data extraction, and data synthesis can be found in [18]. Briefly, peer-reviewed scientific original articles on patients with PD were included in this review. Participants were not required to undergo any type of intervention and the wearable technology could contain any combination of electronic or spring-levered uni- or multiaxial accelerometer, gyroscope, magnetometer, or barometer. The experimental protocol could take place in a laboratory or in a free-living environment. The reported outcomes contained parameters quantifying daily walking activity (e.g., daily step count or distance travelled). After completion of the screening process, two reviewers (M.C. and C.H.) independently extracted the data from each included article including wearing location, technology used, and the methodology to capture daily walking activity.

3. Results

3.1. Study Selection

A total of 72 studies were identified. Additional content (n = 4) was included with this selection from other sources and the article selection process is detailed in a flowchart (Figure 1). After removing duplicates, a total of 60 studies were screened based on the abstracts. This first step excluded 50% of abstracts according to inclusion and exclusion criteria. Then, the thirty eligible articles remaining were assessed on full text. Four studies were excluded after full reading and finally twenty-six studies were eligible and included in the systematic review.

3.2. Study Characteristics

The earliest research dates back to 2004, and interest has sparked over the last years with no less than 14 studies published since 2017 (Figure 2). Fifty percent of the included studies originates from the USA followed by Sweden and the United Kingdom (Figure 3).

The objective of the 26 studies ranged from observational (n = 18, 69.2%) [19–36] to interventional studies (n = 6, 23.1%) [37–42] including pre- and post-measurements or the comparison of walking activity parameters with a control group. Two studies can be considered as proof of concept [28,43]. Six main objectives have been pursued in the included studies:

- 1. To evaluate ambulatory activity [21,23,26,29–31,33,35–37];
- 2. To assess the effect of a training program [27,38,39,41,42,44];
- 3. To examine the relationship between two variables [20,22,24,25,28,32];
- 4. To compare accuracy of wearable sensors according to on-body location [40] and environment [34];
- 5. To compare PD parameters assessed with the ActiGraph GT3X+ (AGT3X) accelerometer and processed with two different filter settings [40];



6. To investigate the reliability and validity of a device [43].

Figure 1. Flowchart of study selection. A total of 26 studies were included according to the eligible criteria in the synthesis.



Figure 2. Number of published studies per year (*n*).



Figure 3. Number of eligible papers published per country (*n*).

3.3. Sample Characteristics

The 26 studies combined a sample size of 1263 people with PD and healthy controls (n = 317). Table A1 in Appendix A, reporting participant's characteristics, shows mean age ranging from 54.9 [30] to 73.4 [23] years old, mean height (160 [29]–172 cm [19]), a mean weight (69.2 [36]–77 kg [19]), and a mean BMI (24.2 [21]–28.07 [42]) among studies reporting these anthropometric data. Among all participants, 584 females and 679 males were included in this review. Disease severity was assessed using Hoehn and Yahr (n = 20, 76.9%) [19,20,22–24,26–31,33–38,41,42,44] and MDS-UPDRS Part III (n = 18, 69.2%) [19,20,22,24–27,29,31,32,34–39,41,42]. If reported, disease duration ranged from 0.8 [37] to 12.5 years [30] (Table A2).

3.4. Type of Sensors

The most frequently used wearable device was the AGT3X [19–23,36–40] used in 10 studies (38.5%). StepWatch Activity Monitors (SAM) were used in seven studies (26.9%): the StepWatch (n = 3, 11.5%) [24,25,43], the StepWatch 3 (n = 3, 11.5%) [26,27,41], and the StepWatch 4 (n = 1, 3.8%) [28]. ActivPAL (n = 1, 3.8%) [29,30] and FitBit Charge HR (n = 1, 3.8%) [31,32] were both used in two studies. Finally, the following wearable devices were employed only in one study each: ActiTrainer (n = 1, 3.8%) [33], Axivity AX3 (n = 1, 3.8%) [42], BioStampRC (n = 1, 3.8%) [34], FitBit Zip (n = 1, 3.8%) [44], Garmin Forerunner 405 GPS Watch (n = 1, 3.8%) [33], and PAMSys (n = 1, 3.8%) [35]. Table A3 gives a summary of study characteristics and provides information on device, wearing location, and length of monitoring.

Figure 4 illustrates proportion of sensors. The most commonly used sensors were triaxial accelerometers (n = 16, 61.5%) [19–23,31–40,42], followed by microprocessor linked devices which were used in seven studies (26.9%) [24–28,41,43]. Furthermore, two (7.7%) studies conducted assessments with uniaxial accelerometers [29,30]. Two studies assessed daily physical activity with a wireless activity tracker (n = 1, 3.8%) [44] and a Global Position System (n = 1, 3.8%) [33].



Figure 4. Proportion type of wearable devices used in the included studies (*n*; %).

3.5. Outcomes of Interest

In this review, 24 studies linked to the use of wearable devices in the context of walking activity were included. According to inclusion criteria, all studies somehow measured walking activity. The included studies have reported various outcomes such as steps per day, gait speed, and time spent walking but also more general outcomes such as sedentary time, active time, activity counts, walking activity, light physical activity, moderate to vigorous physical activity, and high intensity physical activity.

Outcomes of interest are presented in Table A4. Participants with PD had a large range of 2022 to 10,639 steps/day (Figure A1). Eight studies evaluated gait speed of the population [19,26,31,34–36,38,43], including self-selected speed and maximal speed [26,31].

Figure 5 allows a better understanding of the outcomes measured. Sedentary behavior was explored in 11 studies (42.3%) [19,21,23,26,31–35,39,40], and measured in min·day⁻¹ or in min·hours⁻¹ of wear time. Finally, active time is an outcome of interest due to the important number of articles which evaluated active time (n = 21, 80.8%) [19–24,26,27,29–37,39,40,42,44]. Active time was identified by: active time, activity counts, walking activity, light physical activity, moderate to vigorous physical activity, high intensity physical activity, or brisk walking.



Figure 5. Outcomes measured in included studies (*n*).

3.6. Monitoring Protocol

Duration of monitoring (Figure 6) was very incongruent and presented a range from eight hours [33] to one year [26]. However, 11 studies (42.2%) measured walking activity for a 7-day period [19,21,22,27–30,40–43].





The device location changed within the included studies, resulting in eight different on-body locations. The device on-body location is shown in Figure 7, high-lighting three main device locations: waist (n = 10, 38.5%) [19,21–23,33,37–40,44], ankle (n = 6, 23.1%) [29,30,43,45–47], and wrist (n = 5, 19.2%) [19,31–33,36]. Five studies (19.2%) [32–34,36,41] did not provide details about the laterality of the device location.

In the panel of included studies, 12 recorded physical activity also during the night [19,23,27,30–32,34–36,40,42,43]. Other studies kept their focus only on daytime hours [20–22,24–26,28,33,37–39,41]. Two studies did not provide detailed information about the wearing time [29,44].

Only one study [35] mentioned declared specifically to have recorded weekdays, while four studies [24,25,33,34] did not specify whether weekdays or the weekend was measured. Only two studies indicated clearly if they measured weekdays or weekend days and presented results for each period [23,37].

Three studies compared baseline to final measures [26,30,38]. Five studies evaluated PD patients and healthy controls [31–34,43], while PD patients were also compared to other groups: ActiGraph wearers vs. non-ActiGraph wearers [22], mHealth vs. active control [41], and older fallers vs. fallers with mild cognitive impairment [42]. Nine studies (34.6%) reported the measured outcomes without additional comparison to other groups, conditions, or outcomes [21,23–25,27,28,37,40,44].



Figure 7. Location and percentage of devices according to method description.

4. Discussion

The aim of this systematic literature review was to document the use of wearable technology for objective measurement of walking activity in people with PD [18].

Based on the included studies, triaxial accelerometers were the most represented wearable devices [19–23,31–40,42], followed by microprocessor-linked devices [24–28,41,43]. Both devices allow for an investigation of walking as well as physical activity measures depending on the underlying software/algorithm packages. At this point it has to be mentioned that some of the stated devices only provide raw sensor data, i.e., accelerations that need custom post-processing to compute the desired parameters. Systems such as the AGT3X sensor or the FitBit devices are extensively used to measure step numbers and physical activity intensity even though the parameters are not necessarily validated in the studied cohort. As an example, Riel et al. (2016) validated the ActiGraph step count for post-stroke survivors at specific walking velocities [45], but not yet for patients with PD [46]. Nonetheless, the devices prove beneficial when estimating steps under freeliving conditions [47], similarly to the FitBit at least in healthy female adults [48]. Overall, commercial wearable devices are accurate within their application specification [49] and it is necessary to understand for which clinical cohort the devices are actually validated and if they can be used in the clinical setting. Furthermore, the implementation of wearables in clinical settings requires formation and training of health professionals and patients when using wearable activity trackers [50,51].

While some wearable devices may be of preference for research, Bodine and Gemperle (2003) highlighted that the function of any wearable tool must outweigh any physical or social discomfort felt when wearing it [52]. This directly influences the sensor placement, which as a consequence affects the reliability and validity of the sensor outcomes. Kim et al. (2019) compared the number of steps recorded by an AGT3X attached at the wrist (least affected hand) and one attached at the waist (right hip) in a cohort of PD patients. At moderate speed (1.05–1.3 m/s), results showed an overestimation of daily step count for the wrist and an underestimation for the device recording at waist level. Similarly, in the laboratory environment, waist-worn sensors showed higher accuracy compared to wrist worn sensors, but all activity monitors underestimated the number of steps [53].

Yet, most of our walking activity happens outside the laboratory and the included studies provide a range from eight hours [33] to one year [26] for the monitoring period. While none of the studies justified their monitoring period, Cavanaugh et al. (2012) provided evidence that walking activity of PD patients did not differ if you measure longitudinal data over the course of one year, compared to shorter monitoring periods. Their findings are supported by Paul et al. (2016) who determined that two consecutive days of monitoring are sufficient to estimate daily activity reliably during a representative week in people with PD [54]. Moreover, the amount of ambulatory activity was greater on weekdays than weekends in this study which is supported by Christiansen et al. (2017) who found a significant difference for number of steps between weekdays and weekend days. This does not only hold true for PD patients but also for adolescent girls who present a greater activity and greater moderate vigorous intensities on weekdays than on weekends [55]. The concept of weekend warriors and couch potatoes is well established and depending on the employment, educational level, and household income, activity behaviors and patterns during the weekend differ, which is supported by [56,57].

However, in PD patients, the situation is more complicated. While overweight/obesity are common [58], age, gender, education, disease duration, Hoehn and Yahr stage, UPDRS-II and UPDRS-III scores, and dosage of levodopa do not correlate with physical activity [59].

In addition, behavioral and environmental factors are known to affect physical performance. A relationship between sleep and physical activity exists, suggesting that sleep quality could deteriorate walking activity [60]. Dog owners also showed greater walking activity compared to people not owning a dog [61,62]. However, walking activity decreased significantly with increasing wind speed, precipitation, and humidity [63].

To conclude, the selection of an appropriate sensor ultimately depends on the purpose of the study, methodological considerations, and the population characteristics [64].

Even though this systematic review highlights major findings to evaluate walking activity, these findings must be interpreted with caution. Only the outcome steps per day was consistent throughout studies, and all other reported outcomes were exclusively reported by the respective study. Future research could analyze these individual outcomes to improve our understanding of the role of the device when measuring walking activity. Data processing techniques vary greatly and make the comparison between studies rather difficult. In addition, this review already includes a large sample of PD patients; however, the variability of PD severity between each group increases the difficulty to compare the specified protocols.

5. Conclusions

To conclude, this systematic review documents the most frequently used wearable devices as well as data collection procedures and data processing in PD patients. Walking activity is mainly assessed during a 2-day period or more, using a triaxial accelerometer preferably located at the hip or ankle. These findings may be taken into account when evaluating walking activity in PD patients.

Author Contributions: Conceptualization, C.H., T.C. and N.V.; methodology, M.B.C., C.H., T.C. and N.V.; formal analysis, M.B.C., C.H. and N.V.; writing—original draft preparation, M.B.C., C.H., T.C. and N.V.; visualization, M.B.C. and T.C.; supervision, C.H. and N.V.; funding acquisition, N.V. All authors have read and agreed to the published version of the manuscript.

Funding: This research was partly funded by the French National Research Agency, within the framework of the "Investissements d'avenir" program (ANR-10-AIRT-05 and ANR-15-IDEX-02). The APC was funded by the French National Research Agency, within the framework of the "Investissements d'avenir" program (ANR-10-AIRT-05). The sponsors had no involvement in study design, the collection, analysis, and interpretation of data, or in writing the manuscript. This work also forms part of a broader translational and interdisciplinary GaitAlps research program (N.V.).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: All data (articles) are available upon reasonable request to the corresponding author C.H.

Conflicts of Interest: The authors declare no conflict of interest. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript; or in the decision to publish the results.

Appendix A

Table A1. Characteristics of patients and anthropometric information.

| Study | Sample Size | Age, Years (SD) | Gender (n Male, %) | Height, cm (SD) | Weight, kg (SD) | BMI, kg/m ² (SD) |
|---------------------------------|--|--|--|--------------------|--------------------|--|
| Busse et al., 2004 [43] | 10 | 67.1 (8.2) | 7 (70.0%) | | | |
| Skidmore et al., 2008 [24] | 24 (26) | 70.0 (9.0) | 18 (69.2%) | | | |
| Xanthopoulos et al., 2008 [25] | 16 | 71 (11) | 10 (62.5%) | | | |
| Ford et al., 2010 [27] | 12 | 67.2 | 11 (91.7%) | | | |
| Cavanaugh et al., 2012 [26] | 33 | 67.06 (8.75) | 22 (66.7%) | | | |
| Rochester et al., 2012 [30] | 14 | 54.9 (9.5) | 9 (52.9%) | | | |
| Roland et al., 2012 [33] | Total = 15 Non-Frail = 5 Pre-Frail = 5 Frail = 5 | Total = $65.0 (9.0)$ NF = $69 (1)$ PF = $65 (10)$ F = $63 (11)$ | 0 (0.0%) | | | NF: 26.31 (5.6) PF: 22.95 (4.3) F: 25.06 (4.3) |
| Lord et al., 2013 [29] | 89 | 67.3 (9.9) | 62 (69.7%) | 160 (7) | | |
| Wallen et al., 2014 [40] | 66 | 73.1 (5.8) | 38 (57.6%) | 171.0 (8.8) | 74.8 (13.8) | 25.5 (3.8) |
| Conradsson et al., 2015 [38] | 47 | 72.9 (6.0) | 28 (59.8%) | 171.8 (9.2) | 75.8 (14.5) | |
| Toosizadeh et al., 2015 [35] | 15 | 71.2 (6.3) | 8 (53.0%) | 164.3 (10.9) | 74.9 (15.3) | 27.5 (6.5) |
| Wallen et al., 2015 [23] | 95 | 73.4 (5.7) | 53 (55.8%) | 171.4 (9.3) | 76.6 (14.2) | 25.8 (3.7) |
| Christiansen et al., 2017 [37] | 113 | 64.3 (8.6) | 63 (55.8%) | | | 26.9 (4.2) |
| Colón-Semenza et al., 2018 [44] | Total = 10 Peer Coach = 5 Peer Mentee = 5 | PC = 64.6 (4.04) PM = 63.4 (2.06) | PC = 3 (60%) PM = 3 (60%) | | | |
| Leavy et al., 2018 [20] | 49 | 75.0 (5.9) | 21 (49.1%) | | | 25.7 (3.5) |
| Porta et al., 2018 [36] | 18 | 68.0 (10.8) | 8 (44.4%) | 165.6 (7.9) | 69.2 (9.4) | |
| Ellis et al., 2019 [41] | Total = 51 mHealth = 26 | Total = 64.1 (9.5) mH = 64.8 (8.5) | Total = 28 (74.5%) mH = 15 (57.7) | | | |
| | Active Control = 25 | AC = 63.3 (10.6) | AC = 13 (52.0) | | | |
| Kim et al., 2019 [19] | 46 | 68.0 (7.9) | 32 (69.6%) | 172 (9.4) | 77 (15) | |
| Mantri et al., 2019 [22] | Total = 63 ActiGraph Wearer = 30 Non-ActiGraph Wearer = 33 | Total = 70 AW = 70 NAW = 70 | Total = 3 (95.2%) AW = 2 (93.3%) NAW = 1 (97.8%) | | | AW = 26.9 NAW = 26.5 |
| Pradhan and Kelly, 2019 [31] | 30 | 68.6 | 11 (36.7%) | | | |
| Prusynski et al., 2019 [32] | 25 | 69.0 (6.0) | | | | |
| Del Din et al., 2020 [42] | 128 | 71.68 (6.43) | 81 (83.0%) | | | 28.07 (3.62) |
| Adams et al., 2021 [34] | 17 | 66.4 (11.3) | 10 (58.8%) | | | |
| | Total = 110 | Total = 65 | Total = 62 (56.4%) | | | |
| Handlery et al., 2021 [39] | S+ = 74 | S+ = 61.5 | S+ = 45 (60.8%) | | | |
| | S-= 36 | S- = 70.5 | S- = 17 (47.2%) | | | |
| Leavy et al., 2021 [21] | 89 | 71.0 (6.0) | 48 (54.0%) | | | 24.2 (3.5) |
| Zajac et al., 2021 [28] | 69 | 67.5 (8.7) | 40 (58.0%) | | | |

PC = peer coach. PM = peer mentee. mH = mobile health. AC = active control. S+ = more than 4200 steps/day. S- = less than 4220 steps/day. AW = ActiGraph wearer. NAW = non-ActiGraph wearer. NF = non-frail. PF = pre-frail. F = frail.

| Study | Duration Disease, Years (SD) | MDS-UPDRS III: Motor Examination Score (SD) | Hoehn and Yahr (SD) | Medications (LEDD in mg, SD) | |
|---------------------------------|--|--|---|---|--|
| Busse et al., 2004 [43] | | | | | |
| Skidmore et al., 2008 [24] | 7.5 (3.8) | On: 35 (10); Off: 28 (10) | 2 = 9 2.5 = 6 2(4 = 0) | | |
| Xanthonoulos et al 2008 [25] | 70(42) | 29 (11) | 3/4 = 9 NA | | |
| Ford et al., 2010 [27] | 7.0 (1.2) | 12.4 | $ \begin{array}{c} 1 = 4 \\ 1.5 = 1 \\ 2 = 4 \\ 2.5 = 1 \\ 3 = 2 \end{array} $ | | |
| Cavanaugh et al., 2012 [26] | 4.44 (4.21) | Baseline: 28.18 (8.56) One year: 28.52 (11.71) | Baseline: 2 = (1–3) One year: 2 = (1.5–3) | Baseline = 303.03 (294.38) One year = 423.49 (359.66) | |
| Rochester et al., 2012 [30] | 12.5 (6.4) | | 3 = 13 4 = 4 | Baseline = 1387.35 (415.9) 6 months follow-up = 1056.8 (293.0) | |
| Roland et al., 2012 [33] | | | NF = 1.83 (0.8) PF = 1.86 (0.6) F = 2.50 (0.4) | Yes | |
| Lord et al., 2013 [29] | | 25 (10.7) | 1 = 20 (22.5 %) 2 = 51 (57.3 %) 3 = 18 (20.2 %) | 174.6 (124.1) | |
| Wallen et al., 2014 [40] | | | NR | | |
| Conradsson et al., 2015 [38] | 6.0 (5.1) | 36 (10) | 2 = 20 (43%) 3 = 27 (57%) | 581 (295) | |
| Toosizadeh et al., 2015 [35] | 5.9 (5.3) | 34.8 (13.9) | 2.9 (0.9) | 517 (380) | |
| Wallen et al., 2015 [23] | 5.9 (5.0) | | 2 = 41 3 = 54 | | |
| Christiansen et al., 2017 [37] | 0.8 (0.9) | 21.1 (8.8) | 1 = 29 (25.6%) 2 = 84 (74.3%) | No medication | |
| Colón-Semenza et al., 2018 [44] | PC: 5.2 (1.24) PM: 6.2 (2.2) | | 1: PC = 3/PM = 1 2: PC = 1/PM = 3 3: PC = 1/PM = 1 | | |
| Leavy et al., 2018 [20] | 6 | 40 (10.9) | 2 = 22 (45%) 3 = 27 (55%) | 635 (306) | |
| Porta et al., 2018 [36] | 9.9 (6.0) | 17.8 (9.6) | 1.9 (0.4) | Levodopa: MAO-B inhibitors ($n = 11$), Rasagiline ($n = 8$), Safinamide ($n = 3$) | |
| Ellis et al., 2019 [41] | Total: 4.8 (3.1) mH: 5.9 (3.5) AC: 3.7 (2.1) | Tot = 29.6 (10.0) mH = 31.6 (10.7) AC = 27.6 (9.1) | 1: Tot = 1 (2%); mH = 1 (3.9); AC = 0 (0) 1.5: Tot = 2 (3.9%); mH = 1 (3.9); AC = 1 (4.0) 2: 38 (74.5%); mH = 20 (76.9); AC = 18 (72.0) 2.5: Tot = 7 (13.7%); mH = 4 (15.4); AC = 3 (12.0) 3: Tot = 3 (5.9%); mH = 0 (0); AC = 3 (12.0). | | |
| Kim et al., 2019 [19] | 7.6 (6.8) | 34.4 (13.2) | 1 = 4 (8.7%) 2 = 33 (71.7%) 3 = 7 (15.2%) 4 = 2 (4.3%) | Yes | |
| Mantri et al., 2019 [22] | AW: 4 NAW: 3 | AW = 16.5 NAW = 21 | AW = 2 NAW = 2 | | |
| Pradhan and Kelly, 2019 [31] | 7.8 (5.0) | 12.9 (10.3) | 1 = 18 2 = 12 | | |
| Prusynski et al., 2019 [32] | | 12 (9) | | | |
| Del Din et al., 2020 [42] | | 30.37 (16.96) | 2 = 48% 2.5 = 10% 3 = 42% | | |
| Adams et al., 2021 [34] | 4.8 (4.0) | 20.9 (7.9) | 1.9 (0.8) | | |
| Handlery et al., 2021 [39] | Total: 0 S+: 0 S-: 0 | Tot = 19 S+ = 18.5 S- = 22 | | Yes | |
| Leavy et al., 2021 [21] | 6.0 (4.3) | | | 580 (291) | |
| Zajac et al., 2021 [28] | | | 2 = 27 2.5 = 30 3 = 12 | | |

PC = peer coach. PM = peer mentee. mH = mobile health. AC = active control. S+ = more than 4200 steps/day. S- = less than 4220 steps/day. AW = ActiGraph wearer. NAW = non-ActiGraph wearer. NF = non-frail. PF = pre-frail. F = frail.

| Study | Name of Sensor | Manufacturer | Type of Sensor | N of Sensor | Wearing Location | Side | Duration | Days Included | Wear Time |
|--------------------------------|--|--|---------------------------------------|-------------|--|------------------------|--------------------------------------|---------------|---|
| Busse et al., 2004 [43] | SAM | Cymatech, Seattle, WA, USA | Microprocessor-linked device | 1 | Right lower limb above the lateral malleolus | NR | 7 days \times 2 | WD-WED | Continuously. Except during water-related activities. |
| Skidmore et al., 2008 | SAM | Cyma Corporation, Mountlake Terrace, WA, USA | Microprocessor-linked device | 1 | Ankle | NR | 2 days | NR (WD) | Continuously. Except during water-related activities and sleep. |
| Xanthopoulos et al., 2008 [25] | SAM | Cyma Corporation, Mountlake Terrace, WA, USA | Microprocessor-linked device | 1 | Over the right lateral malleolus | NR | 2 days | NR (WD) | Continuously. Except during water-related activities and sleep. |
| Ford et al., 2010 [27] | SAM 3 | Orthocare Innovations, Mountlake Terrace, WA, USA | Microprocessor-linked device | 1 | Ankle | NR | 7 days | WD-WED | Continuously. All periods. |
| Cavanaugh et al., 2012 [26] | SAM 3 | Orthocare Innovations, Mountlake Terrace, WA, USA | Microprocessor-linked device | 1 | Ankle | NR | 1 year | WD-WED | Continuously. During customary activity, including exercise, waking hours. Except during water-related activities. |
| Rochester et al., 2012 [30] | ActivPAL | PAL Technologies, Glasgow, Scotland | Uniaxial accelerometer | 1 | NR | NR | 7 days | WD-WED | Continuously. Except during water-related activities. |
| Roland et al., 2012 [33] | (1) ActiTrainer (2) Garmin Forerunner 405 GPS watch | (1) ActiGraph, LLC, Fort Walton Beach, FL, USA (2) Garmin International Inc., Olathe, KS, USA | (1) Triaxial accelerometer (2) GPS | 2 | (1) Waist (2) Wrist | (1) Dominant (2) NR | 8 h | NR (WD) | Setup of daily physical activity monitors (accelerometer, GPS) were completed at the participant's home in the morning (8–10 a.m.). All PD participants were assessed between 1 and 2 h post anti-Parkinson's medication. The accelerometer, GPS, and physical activity logbook were collected approximately 7 h later (between 4–7 p.m.). |
| Lord et al., 2013 [29] | ActivPAL | PAL Technologies, Glasgow, Scotland | Uniaxial accelerometer | 1 | On the upper thigh | NR | 7 days | WD-WED | NR |
| Wallen et al., 2014 [40] | GT3X+ | ActiGraph, Pensacola, FL, USA | Triaxial accelerometer | 1 | Waist (at hip level above the anterior superior iliac spine) | NR | 3 days (min) and 7days d (max) | WD-WED | Continuously. Except during water-related activities and sleep. |
| Conradsson et al., 2015 [38] | GT3X+ | ActiGraph, Pensacola, FL, USA | Triaxial accelerometer | 1 | Waist | NR | 4 days (or more) | WD-WED | At least 9 h/day. |
| Toosizadeh et al., 2015 [35] | PAMSys | PAMSys, BioSensics, Boston, MA, USA | Triaxial accelerometer | 1 | Sternum | NA | 1 day | WD | Continuously. |
| Wallen et al., 2015 [23] | GT3X+ | ActiGraph, Pensacola, FL, USA | Triaxial accelerometer | 1 | Waist (at hip level) | NR | 7 days | WD-WED | Continuously. Except during water-related activities and sleeping. |

Table A3. Characteristics of sensors and conditions of data acquisition.

Type of Sensor N of Sensor Side Duration Days Included Wear Time Study Name of Sensor Manufacturer Wearing Location 10 h of valid wear time/day minimum, 90 min of ActiGraph, non-wear/day maximum. At Christiansen et al., 2017 [37] GT3X+ Triaxial accelerometer 1 Waist NR 10 days WD-WED least 3 weekdays and Pensacola, FL, USA 1 weekend day of valid wear time. Fitbit Inc., San Colón-Semenza et al., 2018 [44] FitBit Zip 1 Waist NR 8 weeks WD-WED NR Wireless activity tracker Francisco, CA, USA 4 days (min) ActiGraph, Leavy et al., 2018 [20] GT3X+ Triaxial accelerometer 1 NR NR and 7 days WD-WED <540 min/day. Pensacola, FL, USA (max) Continuously. 24 h/24 h. ActiGraph, Porta et al., 2018 [36] GT3X+ Triaxial accelerometer 1 Wrist Nondominant 3 months WD-WED Except during Pensacola, FL, USA water-related activities. Orthocare Continuously. During waking Innovations. Microprocessor-linked Least severe Ellis et al., 2019 [41] SAM 3 1 Leg $7 \text{ days} \times 4$ WD-WED hours. Except during Mountlake Terrace. device impairment water-related activities. WA, USA (1) Least ActiGraph, (1) Wrist 2 WD-WED Kim et al., 2019 [19] GT3X+ Triaxial accelerometer affected hand 7 days Continuously. 24 h/24 h. Pensacola, FL, USA (2) Waist (2) Right hip ActiGraph, Mantri et al., 2019 [22] GT3X+ Triaxial accelerometer 1 Waist NR 7 days WD-WED During walking hours. Pensacola, FL, USA Continuously. Except for the Fitbit Charge Fitbit Inc., San time needed to charge the Pradhan and Kelly, 2019 [31] Triaxial accelerometer 1 Wrist NR 14 days WD-WED HR Francisco, CA, USA device and during water-related activities. Continuously. Except during Fitbit Inc., San water-related activities and for Fitbit Charge Prusynski et al., 2019 [32] Triaxial accelerometer 1 Wrist Nondominant 14 days WD-WED HR Francisco, CA, USA the time needed to charge the device. Continuously. Participants Axivity AX3, Fifth lumbar were asked to continue their Del Din et al., 2020 [42] Axivity AX3 Triaxial accelerometer 1 NA 7 days WD-WED York, UK vertebra daily activities as usual and not to change their routine. (1) One on each anterior thigh. AMC10 Inc., BioStampRC 5 (2) One on each 44 h NR (WD) Adams et al., 2021 [34] Triaxial accelerometer Both sides Continuously. Cambridge, MA, USA anterior forearm. (3) One on the trunk ActiGraph, 1 WD-WED Handlery et al., 2021 [39] GT3X+ Triaxial accelerometer Waist NR 10 days \times 6 During waking hours. Pensacola, FL, USA ActiGraph, Continuously. 4 valid days min. 1 NR WD-WED Leavy et al., 2021 [21] GT3X+ Triaxial accelerometer Hip 7 days Pensacola, FL, USA 9 h or more of wear time/day. Orthocare Continuously. During all Innovations, Microprocessor-linked Above the SAM 4 1 WD-WED waking hours. Except during Zajac et al., 2021 [28] NR 7 days Mountlake Terrace, device lateral malleolus water-related activities. WA, USA

Table A3. Cont.

NR = not reported. WD = weekdays. WED = weekend days. SAM = StepWatch Activity Monitor.

Steps/Day (SD) Study Comparison Gait Speed (m/s) Sedentary Time (SD) Active Time **Baseline (If Possible)** End HC Busse et al., 2004 [43] 3818 0.99 (0.16) Stage 2 = 5147 (1903) Walking activity (steps/h): HY 2 = Skidmore et al., 2008 [24] No comparison Stage 2.5 = 4087 (1286) 20.7 (9.0)/HY 2.5 = 12.9 (3.9)/HY 1 Stages 3/4 = 2708 (1155) 3-4 = 9.6 (4.5)Xanthopoulos et al., 2008 [25] No comparison 1 4378 (2057) Ford et al., 2010 [27] 8996 (3466) Walking activity (min/d): 322 (88) No comparison 1 Max speed: Sedentary time (%): MVPA (min/day): Baseline = 22.49 Cavanaugh et al., 2012 [26] Baseline vs. one year 10,261.15 (4332.56) 9159.44 (3534.21) Baseline = 1.77 (0.50)Baseline = 78.53 (6.93) (24.16)/End = 16.07 (18.68) End = 1.74(0.53)End = 80.14(5.90)**Activity count (n):** T1 = 315.3 Baseline vs. 6 months Rochester et al., 2012 [30] 2258.50 (1373) 2022.40 (1147.20) follow-up (42.9)/T2 = 229.5 (47.4)Active time (%): 32.20 (10.6) Sedentary time (%): 61.70 Roland et al., 2012 [33] HC 3476 (2814) MLTA questionnaire: 3052.3 (14.1)(1611.6) Total = 5452 (2501)Total time spent walking (%): HYI = 6302 (6302)Lord et al., 2013 [29] Control group 1 Total = 5.1 (2) / HYI = 5.8 (1.8) / HYIIHYII = 5335 (2716) = 5.1 (2.2) / HYIII = 4.4 (1.6)HYIII = 4840 (1851)LPA I (min/d): 138 (64) Sedentary time (min/d): Wallen et al., 2014 [40] No comparison 1 4730 (3210) LPA II (min/d): 31 (24) 612 (103) MVPA (min/d): 17 (22) Baseline: 1.19 (0.03) Baseline Conradsson et al., 2015 [38] 4842 (528) 5123 (545) Control group End: 1.28 (0.03) Sitting (%): 44.11 (16.39) Walking activity (%): 6.02 (3.83) Standing (%): 14.40 (7.79) Walking episodes (n): 381 (205) Toosizadeh et al., 2015 [35] Control group 1 4099 (2673) 0.66 (0.11) Lying (%): 35.36 22.01) Max steps (n): 189 (290) **LPA (min/d):** M4/7 = 140.6/3-5 WD = 143.0 / WE = 142Minimum of 4/7 d/week = Sedentary time (min/d): MPALS (min/d): M4/7 = 30.1/3-5 M4/7 = 588.94765 WD = 30.7/WE = 31.3Wallen et al., 2015 [23] No comparison 1 3-5 Weekdays = 4721 3-5 WD = 593.9 **MVPA (min/d):** M4/7 = 16.4/3-5Weekend days = 4888 WE = 584.5WD = 15.2/WE = 18.4Activity count (n): 135,721 (92 368) Total = 5362 (2890) Activity count (n): Total = 345,870 Christiansen et al., 2017 [37] No comparison 1 WD = 5573 (3144) (141,796)/WD = 353,196 WE = 4692 (2800)(154,472)/WE = 321,456 (141,173)Activity (min/week): Baseline = Colón-Semenza et al., 2018 [44] No comparison 5428 (2440) 7115 (1291) 199(95)/End = 282(83) LPA (min/d): 197 (72) Leavy et al., 2018 [20] Control group 1 3653 MVPA (min/d): 47 (34) Total = 10,639 MVPA (%): 06:00-12:00 = 06::00-12:00 = 4313Baseline = 1.18Porta et al., 2018 [36] Other studies 1 43.2/12:00-18:00 = 12:00-18:00 = 3437 End = 1.18 36.3/18:00-22:00 = 31.418:00-22:00 = 2889

Table A4. Main outcomes of the included studies.

Table A4. Cont.

| Study | Comparison | Steps/Day (SD) | | Cait Speed (m/s) | Sedentary Time (SD) | A stive Time |
|------------------------------|--|--------------------------------------|--|---|---|---|
| Study | Companson | Baseline (If Possible) | End | Gait Speed (IIVS) | Sedentary Time (SD) | Active 11me |
| Ellis et al., 2019 [41] | mHealth vs. active control | mH = 8478 (3699) AC = 8902 (2967) | mH = 8457 (3184) AC = 9028 (3366) | | | |
| Kim et al., 2019 [19] | Wrist vs. waist | / | Wrist = 9236 (3812) Waist = 5324 (2800) | Time spent walking < 1.04 (%): Wrist = 71 Waist = 95 Time spent walking 1.05–1.30 (%): Wrist = 29 Waist = 4 Time spent walking > 1.31 (%): Wrist = 0 Waist = 1 | Sedentary time (%): Wrist = 38 (13) Waist = 70 (11) | LPA (%): Wrist = 51 (9)/Waist = 28 (10) MVPA (%): Wrist = 11 (8)/Waist = 2 (2) Activity count (n): Wrist = 872 590 (349 148)/Waist = 186 491 (101 989) |
| Mantri et al., 2019 [22] | ActiGraph Wearers vs. Non-ActiGraph Wearers | / | 3615 | | | MVPA (min): 8.1 (Q = 2.2–23.2) |
| Pradhan and Kelly, 2019 [31] | НС | / | 6416.9 (2795.5) | Self-selected speed: 1.5 (0.3) Max speed: 2.2 (0.4) | Sedentary time (min/d)/(%): 803.7 (154.9)/78.9 (7.8) | LPA (%): 17.8 (6.6) MVPA (%): 2.7 (1.9) |
| Prusynski et al., 2019 [32] | НС | / | 5953 (2363) | | Sedentary time (min/d): 846 (122) | LPA (min): 172 (60) MVPA (min): 15 (11) HIPA (min): 19 (14) |
| Del Din et al., 2020 [42] | Older fallers vs. fallers with MCI | I1 = 8874 (4538) I2 = 9000 (4535) | I1 = 8654 (4638) I2 = 9468 (5576) | | | Total Walking Time per Day (min): Baseline: I1 = $117.3 (59.1)/I2 =$ 117.8 (62.3) End: I1 = $116.2 (65.1)/I2 =$ 127.9 (84.3) Percentage of Walking Time (%): Baseline: I1 = $8.2 (4.1)/I2 = 8.2 (4.3)$ End: I1 = $8.1 (4.5)/I2 = 8.9 (5.9)$ Activity count (act/d): Baseline: I1 = $233 (106)/I2 =$ 242 (110) End: I1 = $229 (116)/I2 = 254 (107)$ |
| Adams et al., 2021 [34] | НС | / | 4980 | 0.91 | Lying (h/d): 9.1 Sitting (h/d): 10.7 Standing (h/d): 3.3 | Walking activity (h/d): 0.9 (0.5–1.3) |
| Handlery et al., 2021 [39] | >4200 steps/day (S+) vs. <4200 steps/day (S–) | / | Tot = 4817 S+ = 6066 S- = 2852 | | Sedentary time (min/h of wear time): Tot = 33.5 S+ = 32.3 S- = 36 | Total daily MVPA (min): Total = $36.8/S+ = 49/S- = 11.1$ LPA (min/h of wear time): Total = $23.3 (5.5)/S+ = 23.5$ ($5.8)/S- = 22.9 (5.0)$ MVPA (min/h of wear time): Total = $2.8 (0.1-9.5)/S+ = 3.5$ ($1.3-9.5)/S- = 0.8 (0.1-3.4)$ Activity count (n): Total = $330,500$ ($109,670-820,247)/S+ = 376,152$ ($168,845-820,247)/S- = 239,875$ ($109,670-365,337$) |

| Table | A4. | Cont. |
|-------|-----|---------|
| Invit | | 001111. |

| Study | Comparison | Steps/Day (SD) | | Cait Speed (m/s) | Sodontary Time (SD) | A stive Time |
|-------------------------|---------------|------------------------|-----------------|-------------------|-------------------------------------|---|
| | Companison | Baseline (If Possible) | End | Gait Speed (iivs) | Sedentary Time (SD) | Acuve fillite |
| Leavy et al., 2021 [21] | No comparison | 1 | 5876 (3180) | | Sedentary time (min/d): 598 (92) | Brisk walking (min/d): 23.5 (Q = 5.4-42.2) LPA (min/d): 197 (72) MVPA (min/d): 47 (34) |
| Zajac et al., 2021 [28] | No comparison | / | 7606.2 (3625.8) | | | |

HC = healthy control. I = intervention. WD = weekdays. WE = weekend days. MLTA = Minnesota Leisure Time Activity. LPA = light physical activity. MVPA = moderate vigorous physical activity. HIPA = high intensity physical activity. mH = mHealth. AC = active control. HY = Hoehn and Yahr. MPALS = moderate-intensity lifestyle activities.



Figure A1. Wear time (in days) and mean daily step (*n* per day) of included studies.

References

- 1. Gelb, D.J.; Oliver, E.; Gilman, S. Diagnostic Criteria for Parkinson Disease. Arch. Neurol. 1999, 56, 33–39. [CrossRef]
- Poewe, W.; Seppi, K.; Tanner, C.M.; Halliday, G.M.; Brundin, P.; Volkmann, J.; Schrag, A.E.; Lang, A.E. Parkinson disease. *Nat. Rev. Dis. Primers* 2017, *3*, 17013. [CrossRef]
- Chaudhuri, K.R.; Healy, D.G.; Schapira, A.H. Non-motor symptoms of Parkinson's disease: Diagnosis and management. *Lancet Neurol.* 2006, *5*, 235–245. [CrossRef]
- 4. Warmerdam, E.; Romijnders, R.; Hansen, C.; Elshehabi, M.; Zimmermann, M.; Metzger, F.G.; von Thaler, A.K.; Berg, D.; Schmidt, G.; Maetzler, W. Arm swing responsiveness to dopaminergic medication in Parkinson's disease depends on task complexity. *NPJ Parkinson's Dis.* **2021**, *7*, 89. [CrossRef]
- 5. Lang, A.E.; Lozano, A.M. Parkinson's disease. Second of two parts. N. Engl. J. Med. 1998, 339, 1130–1143. [CrossRef]
- Bloem, B.R.; Beckley, D.J.; van Dijk, J.G.; Zwinderman, A.H.; Remler, M.P.; Roos, R.A.C. Influence of dopaminergic medication on automatic postural responses and balance impairment in Parkinson's disease. *Mov. Disord. Off. J. Mov. Disord. Soc.* 1996, 11, 509–521. [CrossRef]
- Rosqvist, K.; Horne, M.; Hagell, P.; Iwarsson, S.; Nilsson, M.H.; Odin, P. Levodopa effect and motor function in late stage Parkinson's disease. J. Parkinson's Dis. 2018, 8, 59–70. [CrossRef]
- 8. Hong, M.; Earhart, G.M. Effects of medication on turning deficits in individuals with Parkinson's disease. *J. Neurol. Phys. Ther. JNPT* **2010**, *34*, 11. [CrossRef]
- 9. Glendinning, D.S. A Rationale for Strength Training in Patients with Parkinson's Disease. J. Neurol. Phys. Ther. 1997, 21, 132–135.
- 10. Li, F.; Harmer, P.; Fitzgerald, K.; Eckstrom, E.; Stock, R.; Galver, J.; Maddalozzo, G.; Batya, S.S. Tai chi and postural stability in patients with Parkinson's disease. *N. Engl. J. Med.* **2012**, *366*, 511–519. [CrossRef]
- 11. Chung, C.L.H.; Thilarajah, S.; Tan, D. Effectiveness of resistance training on muscle strength and physical function in people with Parkinson's disease: A systematic review and meta-analysis. *Clin. Rehabil.* **2016**, *30*, 11–23. [CrossRef]
- 12. Bridgewater, K.J.; Sharpe, M.H. Trunk muscle training and early Parkinson's disease. *Physiother. Theory Pract.* **1997**, *13*, 139–153. [CrossRef]
- 13. Van Nimwegen, M.; Speelman, A.D.; Hofman-van Rossum, E.J.; Overeem, S.; Deeg, D.J.; Borm, G.F.; van der Horst, M.H.; Bloem, B.R.; Munneke, M. Physical inactivity in Parkinson's disease. *J. Neurol.* **2011**, *258*, 2214–2221. [CrossRef]
- 14. Speelman, A.D.; van de Warrenburg, B.P.; van Nimwegen, M.; Petzinger, G.M.; Munneke, M.; Bloem, B.R. How might physical activity benefit patients with Parkinson disease? *Nat. Rev. Neurol.* **2011**, *7*, 528–534. [CrossRef]
- 15. Galperin, I.; Hillel, I.; Del Din, S.; Bekkers, E.M.; Nieuwboer, A.; Abbruzzese, G.; Avanzino, L.; Nieuwhof, F.; Bloem, B.R.; Rochester, L.; et al. Associations between daily-living physical activity and laboratory-based assessments of motor severity in patients with falls and Parkinson's disease. *Parkinsonism Relat. Disord.* **2019**, *62*, 85–90. [CrossRef]
- 16. Moher, D.; Liberati, A.; Tetzlaff, J.; Altman, D.G.; Group, P. Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. *PLoS Med.* **2009**, *6*, e1000097. [CrossRef]
- Tricco, A.C.; Lillie, E.; Zarin, W.; O'Brien, K.K.; Colquhoun, H.; Levac, D.; Moher, D.; Peters, M.D.; Horsley, T.; Weeks, L.; et al. PRISMA extension for scoping reviews (PRISMA-ScR): Checklist and explanation. *Ann. Intern. Med.* 2018, 169, 467–473. [CrossRef]
- 18. Carlin, T.; Hansen, C.; Vuillerme, N. Objective Measurement of Walking Activity Using Wearable Technologies in People with Parkinson Disease: A Systematic Review Protocol. *Biomed. Hub* **2021**, *6*, 64–68. [CrossRef]
- 19. Kim, D.W.; Hassett, L.M.; Nguy, V.; Allen, N.E. A comparison of activity monitor data from devices worn on the wrist and the waist in people with Parkinson's disease. *Mov. Disord. Clin. Pract.* **2019**, *6*, 693–699. [CrossRef]
- 20. Leavy, B.; Löfgren, N.; Nilsson, M.; Franzén, E. Patient-reported and performance-based measures of walking in mild–moderate Parkinson's disease. *Brain Behav.* 2018, *8*, e01081. [CrossRef]
- Leavy, B.; Hagströmer, M.; Conradsson, D.M.; Franzén, E. Physical activity and perceived health in people with Parkinson disease during the first wave of COVID-19 pandemic: A cross-sectional study from Sweden. J. Neurol. Phys. Ther. 2021, 45, 266–272. [CrossRef]
- 22. Mantri, S.; Wood, S.; Duda, J.E.; Morley, J.F. Comparing self-reported and objective monitoring of physical activity in Parkinson disease. *Parkinsonism Relat. Disord.* 2019, 67, 56–59. [CrossRef]
- 23. Benka Wallén, M.; Franzén, E.; Nero, H.; Hagströmer, M. Levels and patterns of physical activity and sedentary behavior in elderly people with mild to moderate Parkinson disease. *Phys. Ther.* **2015**, *95*, 1135–1141. [CrossRef]
- 24. Skidmore, F.M.; Mackman, C.A.; Pav, B.; Shulman, L.M.; Garvan, C.; Macko, R.F.; Heilman, K.M. Daily ambulatory activity levels in idiopathic Parkinson disease. *J. Rehabil. Res. Dev.* **2008**, 45, 1343–1348. [CrossRef]
- 25. Xanthopoulos, P.; Heilman, K.M.; Drago, V.; Pardalos, P.; Foster, P.S.; Skidmore, F.M. An ambulatory persistence power curve: Motor planning affects ambulatory persistence in Parkinson's disease. *Neurosci. Lett.* **2008**, 448, 105–109. [CrossRef]
- Cavanaugh, J.T.; Ellis, T.D.; Earhart, G.M.; Ford, M.P.; Foreman, K.B.; Dibble, L.E. Capturing ambulatory activity decline in Parkinson disease. J. Neurol. Phys. Ther. JNPT 2012, 36, 51. [CrossRef]
- 27. Ford, M.P.; Malone, L.A.; Walker, H.C.; Nyikos, I.; Yelisetty, R.; Bickel, C.S. Step activity in persons with Parkinson's disease. *J. Phys. Act. Health* **2010**, *7*, 724–729. [CrossRef]

- Zajac, J.A.; Cavanaugh, J.T.; Baker, T.; Colón-Semenza, C.; DeAngelis, T.R.; Duncan, R.P.; Fulford, D.; LaValley, M.; Nordahl, T.; Rawson, K.S.; et al. Are mobile persons with Parkinson disease necessarily more active? *J. Neurol. Phys. Ther.* 2021, 45, 259–265. [CrossRef]
- 29. Lord, S.; Godfrey, A.; Galna, B.; Mhiripiri, D.; Burn, D.; Rochester, L. Ambulatory activity in incident Parkinson's: More than meets the eye? *J. Neurol.* 2013, 260, 2964–2972. [CrossRef]
- 30. Rochester, L.; Chastin, S.F.M.; Lord, S.; Baker, K.; Burn, D.J. Understanding the impact of deep brain stimulation on ambulatory activity in advanced Parkinson's disease. *J. Neurol.* 2012, 259, 1081–1086. [CrossRef]
- 31. Pradhan, S.; Kelly, V.E. Quantifying physical activity in early Parkinson disease using a commercial activity monitor. *Parkinsonism Relat. Disord.* **2019**, *66*, 171–175. [CrossRef]
- 32. Prusynski, R.A.; Kelly, V.E.; Fogelberg, D.J.; Pradhan, S. The association between sleep deficits and sedentary behavior in people with mild Parkinson disease. *Disabil. Rehabil.* **2021**, 1–7. [CrossRef]
- 33. Roland, K.P.; MDCornett, K.; Theou, O.; Jakobi, J.M.; Jones, G.R. Physical activity across frailty phenotypes in females with Parkinson's disease. *J. Aging Res.* 2012, 2012, 468156. [CrossRef]
- 34. Adams, J.L.; Dinesh, K.; Snyder, C.W.; Xiong, M.; Tarolli, C.G.; Sharma, S.; Dorsey, E.; Sharma, G. A real-world study of wearable sensors in Parkinson's disease. *NPJ Parkinson's Dis* 2021, 7, 106. [CrossRef]
- Toosizadeh, N.; Mohler, J.; Lei, H.; Parvaneh, S.; Sherman, S.; Najafi, B. Motor performance assessment in Parkinson's disease: Association between objective in-clinic, objective in-home, and subjective/semi-objective measures. *PLoS ONE* 2015, 10, e0124763. [CrossRef]
- Porta, M.; Pilloni, G.; Pili, R.; Casula, C.; Murgia, M.; Cossu, G.; Pau, M. Association between objectively measured physical activity and gait patterns in people with Parkinson's disease: Results from a 3-month monitoring. *Parkinson's Dis.* 2018, 2018, 7806574. [CrossRef]
- 37. Christiansen, C.; Moore, C.; Schenkman, M.; Kluger, B.; Kohrt, W.; Delitto, A.; Berman, B.; Hall, D.; Josbeno, D.; Poon, C.; et al. Factors associated with ambulatory activity in de novo Parkinson disease. *J. Neurol. Phys. Ther. JNPT* **2017**, *41*, 93. [CrossRef]
- Conradsson, D.; Löfgren, N.; Nero, H.; Hagströmer, M.; Ståhle, A.; Lökk, J.; Franzén, E. The effects of highly challenging balance training in elderly with Parkinson's disease: A randomized controlled trial. *Neurorehabil. Neural Repair* 2015, 29, 827–836. [CrossRef]
- Handlery, R.; Stewart, J.C.; Pellegrini, C.; Monroe, C.; Hainline, G.; Flach, A.; Handlery, K.; Fritz, S. Physical Activity in De Novo Parkinson Disease: Daily Step Recommendation and Effects of Treadmill Exercise on Physical Activity. *Phys. Ther.* 2021, 101, pzab174. [CrossRef]
- 40. Wallén, M.B.; Nero, H.; Franzén, E.; Hagströmer, M. Comparison of two accelerometer filter settings in individuals with Parkinson's disease. *Physiol. Meas.* **2014**, *35*, 2287. [CrossRef]
- Ellis, T.D.; Cavanaugh, J.T.; DeAngelis, T.; Hendron, K.; Thomas, C.A.; Saint-Hilaire, M.; Pencina, K.; Latham, N.K. Comparative effectiveness of mHealth-supported exercise compared with exercise alone for people with Parkinson disease: Randomized controlled pilot study. *Phys. Ther.* 2019, 99, 203–216. [CrossRef] [PubMed]
- Del Din, S.; Galna, B.; Lord, S.; Nieuwboer, A.; Bekkers, E.M.; Pelosin, E.; Avanzino, L.; Bloem, B.R.; Olde Rikkert, M.G.; Nieuwhof, F.; et al. Falls risk in relation to activity.y exposure in high-risk older adults. *J. Gerontol. Ser. A* 2020, 75, 1198–1205. [CrossRef] [PubMed]
- 43. Busse, M.E.; Pearson, O.R.; van Deursen, R.; Wiles, C.M. Quantified measurement of activity provides insight into motor function and recovery in neurological disease. *J. Neurol. Neurosurg. Psychiatry* **2004**, *75*, 884–888. [CrossRef] [PubMed]
- 44. Colón-Semenza, C.; Latham, N.K.; Quintiliani, L.M.; Ellis, T.D. Peer coaching through mhealth targeting physical activity in people with Parkinson disease: Feasibility study. *JMIR mHealth uHealth* **2018**, *6*, e8074. [CrossRef] [PubMed]
- Riel, H.; Rathleff, C.R.; Kalstrup, P.M.; Madsen, N.K.; Pedersen, E.S.; Pape-Haugaard, L.B.; Villumsen, M. Comparison between Mother, ActiGraph wGT3X-BT, and a hand tally for measuring steps at various walking speeds under controlled conditions. *PeerJ* 2016, 4, e2799. [CrossRef] [PubMed]
- 46. Campos, C.; DePaul, V.G.; Knorr, S.; Wong, J.S.; Mansfield, A.; Patterson, K.K. Validity of the ActiGraph activity monitor for individuals who walk slowly post-stroke. *Top. Stroke Rehabil.* **2018**, *25*, 295–304. [CrossRef]
- 47. Henriksen, A.; Johansson, J.; Hartvigsen, G.; Grimsgaard, S.; Hopstock, L. Measuring physical activity using triaxial wrist worn polar activity trackers: A systematic review. *Int. J. Exerc. Sci.* **2020**, *13*, 438.
- Reid, R.E.; Insogna, J.A.; Carver, T.E.; Comptour, A.M.; Bewski, N.A.; Sciortino, C.; Andersen, R.E. Validity and reliability of Fitbit activity monitors compared to ActiGraph GT3X+ with female adults in a free-living environment. *J. Sci. Med. Sport* 2017, 20, 578–582. [CrossRef]
- Fuller, D.; Colwell, E.; Low, J.; Orychock, K.; Tobin, M.A.; Simango, B.; Buote, R.; Van Heerden, D.; Luan, H.; Cullen, K.; et al. Reliability and validity of commercially available wearable devices for measuring steps, energy expenditure, and heart rate: Systematic review. *JMIR mHealth uHealth* 2020, *8*, e18694. [CrossRef]
- 50. Evenson, K.R.; Goto, M.M.; Furberg, R.D. Systematic review of the validity and reliability of consumer-wearable activity trackers. *Int. J. Behav. Nutr. Phys. Act.* 2015, 12, 159. [CrossRef]
- 51. Kooiman, T.J.M.; Dontje, M.L.; Sprenger, S.R.; Krijnen, W.P.; van der Schans, C.P.; de Groot, M. Reliability and validity of ten consumer activity trackers. *BMC Sports Sci. Med. Rehabil.* 2015, 7, 24. [CrossRef]

- 52. Bodine, K.; Gemperle, F. Effects of functionality on perceived comfort of wearables. In Proceedings of the Seventh IEEE International Symposium on Wearable Computers, White Plains, NY, USA, 21–23 October 2003; p. 57.
- 53. Wendel, N.; Macpherson, C.E.; Webber, K.; Hendron, K.; DeAngelis, T.; Colon-Semenza, C.; Ellis, T. Accuracy of activity trackers in Parkinson disease: Should we prescribe them? *Phys. Ther.* **2018**, *98*, 705–714. [CrossRef]
- Paul, S.S.; Ellis, T.D.; Dibble, L.E.; Earhart, G.M.; Ford, M.P.; Foreman, K.B.; Cavanaugh, J.T. Obtaining reliable estimates of ambulatory physical activity in people with Parkinson's disease. J. Parkinson's Dis. 2016, 6, 301–305. [CrossRef]
- 55. Treuth, M.S.; Catellier, D.J.; Schmitz, K.H.; Pate, R.R.; Elder, J.P.; McMurray, R.G.; Blew, R.M.; Yang, S.; Webber, L. Weekend and weekday patterns of physical activity in overweight and normal-weight adolescent girls. *Obesity* 2007, *15*, 1782–1788. [CrossRef]
- 56. Ramirez-Rico, E.; Hilland, T.A.; Foweather, L.; Fernandez-Garcia, E.; Fairclough, S.J. Weekday and weekend patterns of physical activity and sedentary time among Liverpool and Madrid youth. *Eur. J. Sport Sci.* **2014**, *14*, 287–293. [CrossRef]
- Fukushima, N.; Kikuchi, H.; Amagasa, S.; Machida, M.; Kitabayashi, M.; Hayashi, T.; Odagiri, Y.; Takamiya, T.; Inoue, S. Exposure to prolonged sedentary behavior on weekdays rather than weekends in white-collar workers in comparison with blue-collar workers. J. Occup. Health 2021, 63, e12246. [CrossRef]
- 58. Morales-Briceño, H.; Cervantes-Arriaga, A.; Rodríguez-Violante, M.; Calleja-Castillo, J.; Corona, T. Overweight is more prevalent in patients with Parkinson's disease. *Arq. Neuropsiquiatr.* **2012**, *70*, 843–846. [CrossRef]
- 59. Cai, G.; Huang, Y.; Luo, S.; Lin, Z.; Dai, H.; Ye, Q. Continuous quantitative monitoring of physical activity in Parkinson's disease patients by using wearable devices: A case-control study. *Neurol. Sci.* 2017, *38*, 1657–1663. [CrossRef]
- 60. Semplonius, T.; Willoughby, T. Long-term links between physical activity and sleep quality. *Med. Sci. Sports Exerc.* 2018, 50, 2418–2424. [CrossRef]
- 61. Martin, K.E.; Wood, L.; Christian, H.; Trapp, G.S.A. Not just "a walking the dog": Dog walking and pet play and their association with recommended physical activity among adolescents. *Am. J. Health Promot.* **2015**, *29*, 353–356. [CrossRef]
- 62. Ham, S.A.; Epping, J. Dog Walking and Physical Activity in the United States. Prev. Chronic Dis. 2006, 3, A47.
- 63. Klenk, J.; Büchele, G.; Rapp, K.; Franke, S.; Peter, R.; Group, A.S. Walking on sunshine: Effect of weather conditions on physical activity in older people. *J. Epidemiol. Community Health* **2012**, *66*, 474–476. [CrossRef]
- 64. Tudor-Locke, C.; Williams, J.E.; Reis, J.P.; Pluto, D. Utility of pedometers for assessing physical activity. *Sports Med.* 2002, 32, 795–808. [CrossRef]