Review

The Effectiveness of Wearable Electronic Device System– Supported Physical Activity Programs for Cancer Survivors: Meta-Analysis of Randomized Controlled Trials

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Abstract

Background: As cancer is increasingly regarded as a chronic disease, it is essential to support cancer survivors' self-management and enhance their quality of life (QoL). Although a physically active lifestyle can help alleviate symptom burden, improve QoL, and even benefit survival among cancer survivors, many remain physically inactive. Wearable electronic device systems (WEDSs) have become increasingly integrated into daily life and may offer a potential solution to promote physical activity (PA) and improve QoL in this population. However, existing findings remain modest and inconclusive.

Objective: This meta-analysis aims to evaluate (1) the effects of WEDS-supported PA programs on improving PA, sedentary behavior, BMI, and QoL in cancer survivors; and (2) the effects of various types of these interventions.

Methods: A comprehensive literature search was conducted across PubMed, Embase, Web of Science, CENTRAL, and MEDLINE from database inception through July 31, 2024. Two authors independently screened the articles, extracted the data, and evaluated the methodological quality of the included studies using the Cochrane Risk-of-Bias tool 2. Data synthesis was performed using R Studio. The effects of the interventions were determined by calculating standard mean differences (SMDs) and 95% CIs, while heterogeneity was assessed using *P* statistics and *P* values. Subgroup analysis was conducted to assess whether the effects differed by the formats of the partnering tools and the duration of the intervention. Sensitivity analysis was performed using the one-study-out method to evaluate the robustness of the results, and the Egger test was conducted to assess small study effects. Statistical significance for the overall effect was considered when the 2-tailed *P* value was less than .05.

Results: A total of 46 randomized controlled trials, involving 3727 patients, were included in this meta-analysis. The results indicated that WEDS-supported PA programs significantly improved objectively measured moderate-to-vigorous-intensity physical activity (MVPA; SMD 0.66, 95% CI 0.47-0.86, P<.001, I^2 =69%), subjectively reported PA (SMD 0.5, 95% CI 0.23-0.77, P<.001, I^2 =79%), steps per day (SMD 0.5, 95% CI 0.23-0.77, P=.009, I^2 =79%), and QoL (SMD 0.19, 95% CI 0.08-0.31, P<.001, I^2 =33%) among cancer survivors. Subgroup analysis revealed that interventions incorporating multipartnering tools (no fewer than 2 formats) were effective in improving subjectively reported PA, steps per day, and QoL. Long-term interventions (\geq 12 weeks) improved objectively measured MVPA, subjectively reported PA, steps per day, and QoL. Interventions tailored to specific cancer types significantly improved steps per day (SMD 0.59, 95% CI 0.1-1.08, P=.008, I^2 =83%) and QoL (SMD 0.14, 95% CI 0.04-0.23, I^2 =006, I^2 =0%).

Conclusions: We observed that WEDS-supported PA programs are effective in improving the level of PA (both objectively and subjectively), steps per day, and QoL among cancer survivors, but showed no significant effects on sedentary behavior or BMI. In the future, the use of multipartnering tools, appropriate intervention duration, and tailored PA programs should be

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carefully considered when developing WEDS-supported PA interventions. Further promotion and refinement of WEDS-supported PA programs are warranted.

Trial Registration: PROSPERO CRD42024582905; https://www.crd.york.ac.uk/PROSPERO/view/CRD42024582905

J Med Internet Res 2025;27:e74347; doi: 10.2196/74347

Keywords: oncology; digital health; physical activity; quality of life; PRISMA

Introduction

Owing to advancements in early detection and breakthroughs in cancer treatment, the number of long-term cancer survivors has significantly increased, transforming cancer into a chronic condition and making it a growing global public health concern [1]. When cancer is considered a chronic disease, it is essential to support cancer survivors' self-management and enhance their quality of life (QoL). However, cancer survivors often experience various symptom burdens, including reduced cardiorespiratory fitness and muscle strength, fatigue, sleep disturbances, and emotional distress, all of which affect their QoL and mortality [1-8]. These symptom burdens and decreased QoL are even associated with patient survival [9-12]. Maintaining physical activity (PA) has emerged as a promising lifestyle approach for cancer survivors. PA refers to any bodily movement produced by skeletal muscles that requires energy expenditure [13]. For cancer survivors, adequate PA, especially moderate-tovigorous-intensity PA (MVPA), can improve their cardiorespiratory and muscular fitness, alleviate symptom burden (such as cancer-related fatigue), and enhance QoL [14]. However, when cancer survivors are diagnosed with cancer, their engagement in even little or leisure PA significantly decreases, whereas their time spent sitting increases, which negatively impacts their survival [15,16]. Barriers that prevent cancer survivors from participating in PA may include their symptom burden (such as pain, fatigue, and lymphedema), social factors (such as lack of time, motivation, and support from health care professionals [HCPs]), and lack of information (such as recommendations on PA) [17]. Thus, it is essential to identify interventions that can encourage or remind cancer survivors to increase their level of PA and provide support to enhance their physical fitness, such as muscular fitness and cardiorespiratory fitness, ultimately improving their QoL and survival.

Digital health, defined as the use of "digital technologies for health" [18], including mobile health (mHealth) apps, electronic health records, electronic medical records, wearable electronic devices (WEDs), telehealth and telemedicine, and personalized medicine [19], is an influential force in the progression of global health care toward improved accessibility and quality [20]. WEDs refer to any kind of electronic device designed to be worn on the user's body, as either an accessory or an implant [21]. In the context of behavior change techniques [22], interventions supported by WEDs have become increasingly prevalent among cancer survivors to increase their PA by collecting physical and physiological information, enabling continuous real-time self-health surveillance, and providing stimuli for behavior

change [23-26]. Moreover, WEDs can be combined with partnering tools to form a wearable electronic device system (WEDS), where partnering tools may include telephone calls, SMS text messages, apps, or websites. Interventions supported by WEDS are effective intervention modalities and can offer an optional and novel approach to promoting PA among cancer survivors [25,27,28].

In WEDS-supported PA programs, HCPs usually set PA goals for participants, WEDs facilitate self-monitoring and data collection, and partnering tools typically enable patient contact with HCPs or provide timely feedback from HCPs (such as consultation, goal resetting, and guidance) [28]. Owing to these benefits and their portability, studies exploring WEDS-supported PA in oncology rehabilitation have surged dramatically over the past decade [23,29,30]. Positive effects have been observed in improving PA among older adults, adults, and patients with diabetes, cardiovascular-related diseases, and chronic obstructive pulmonary disease [31-35]. Although evidence suggests that WEDS-supported PA programs can benefit patients with cancer by increasing PA levels and improving health-related outcomes (such as fatigue, muscular fitness, aerobic fitness, and QoL), findings from existing studies on the effects of these programs remain inconsistent [8,23,29,30]. For example, Singh et al [23] reported that WEDS-supported PA programs led to a statistically significant increase in patients' daily step counts, whereas Teo et al [30] found no statistically significant difference in daily step counts between the experimental and control groups. Additionally, in other forms of eHealthsupported interventions for cancer survivors, effectiveness varies depending on the type of eHealth and the duration of the intervention. For example, both Li et al [36] and Su et al [37] reported that the effectiveness of internet-based digital health interventions differed across different subgroups based on the format or duration of the intervention. Moreover, previous studies have consistently overlooked the role of partnering tools, and no researcher has investigated which types of partnering tools may better integrate with WEDs to enhance their effectiveness. Additionally, although numerous studies on this topic have been published, most have focused only on specific types of cancer or examined a limited range of health-related outcomes [29,30]. Furthermore, prior research in this area has predominantly focused on examining the feasibility, acceptability, or overall effects of WED-supported PA programs on cancer survivors. To our knowledge, no meta-analysis has focused specifically on WEDS-supported PA programs, nor has any examined subgroup effects on diverse outcomes, such as different formats of partnering tools for improving PA levels, BMI, or QoL, and decreasing sedentary behavior.

Thus, the objectives of this meta-analysis are (1) to evaluate the effects of WEDS-supported PA programs on increasing PA-related outcomes and QoL, and (2) to explore which type of WEDS is most effective and the optimal duration of the intervention for cancer survivors.

Methods

Study Design

This meta-analysis was conducted in accordance with the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) 2020 statement [38] and has been duly registered with the International Prospective Register of Systematic Reviews (PROSPERO; registration number CRD42024582905). As the data utilized in this research were exclusively sourced from previously published studies, ethical approval and informed consent were not required.

Search Strategy

After consulting a professor of statistics, a comprehensive literature search was carried out, covering the period from the inception of the databases to July 31, 2024. The search was conducted across PubMed, Embase, Web of Science, CENTRAL, and MEDLINE, with access to the full text of the articles. The development of the search strategy was guided by the PICOS (Participants, Interventions, Comparisons, Outcomes, and Study Design) framework, along with the guidelines provided by the Cochrane Collaboration to ensure the integrity of the analysis. This strategy incorporated the use of MeSH (Medical Subject Headings) terms, textual keyword searches, and Boolean logic operations, supplemented by keywords from titles or abstracts, including terms such as neoplasms, carcinomas, tumors, cancer, caregivers, PA trackers, wearable, telemedicine, and telerehabilitation. All search strategies used are presented in Multimedia Appendix 1. The search was limited to studies involving humans and randomized controlled trials and was conducted in English. Moreover, we conducted a rigorous manual review of the bibliographies of the retrieved articles to identify and obtain supplementary relevant scholarly works, thereby enhancing the depth of our analytical inquiry.

Study Eligibility Criteria

This meta-analysis considered studies for inclusion based on the following criteria: (1) participants were survivors of any type of cancer, regardless of sex or cancer stage; (2) patients in the intervention group received WEDS-supported PA programs, which included reminders to change behavior, consultations with HCPs, or social support from other patients; (3) patients in the control group received usual care or were placed on a waitlist; (4) the outcomes included at least one of the following indices—objectively measured MVPA, subjectively reported PA, sedentary behavior, QoL, or BMI—without restrictions on the measures used; (5) the publications were written in English; and (6) the studies were designed as randomized controlled trials. Studies were excluded if they were only registered but not yet conducted or if relevant data were incomplete.

Study Selection and Data Extraction

The reference management software EndNote X9 (Clarivate Plc) was used to import and screen the titles and abstracts of the studies. Duplications were first removed automatically by EndNote and then meticulously screened by researchers. To ensure alignment with the inclusion criteria, 2 independent authors (ZW and YL) concurrently conducted a thorough screening of the titles and abstracts. Subsequently, they carefully evaluated the full texts of the papers based on the predetermined eligibility criteria. Any discrepancies in the screening process were resolved through discussion or by consulting a third author (QW). Data extraction from the included studies was performed independently by 2 authors (ZW and YL), who meticulously recorded the information using a predefined data extraction template. This template encompassed a range of details, including the first author's name, year of publication, country where the study was conducted, participants' ages (mean and SD), sample size, type of cancer diagnosed, types of WEDs and associated tools used, intervention content, duration of the interventions, outcome measures employed, and timing of assessments.

Quality Assessment

The methodological quality and risk of bias in the included studies were meticulously assessed by 2 independent reviewers, using the Risk of Bias Tool 2, version 5.1.0. A total of 7 domains were evaluated: random sequence generation (selection bias), allocation concealment (selection bias), blinding of participants and personnel (performance bias), blinding of outcome assessment (detection bias), incomplete outcome data (attrition bias), selective reporting (reporting bias), and other potential sources of bias. Each domain was graded as "low risk" of bias, "high risk" of bias, and "unclear risk" of bias. The official Cochrane Excel tool was used to automatically compute the overall risk. Disagreements were resolved through discussion or, when necessary, by consulting a third author.

Data Synthesis and Analysis

Utilizing R Studio (R Foundation), we conducted heterogeneity evaluations and performed the meta-analysis. To quantify the intervention effects, we computed the standard mean difference (SMD) along with its corresponding 95% CI, and presented the results using forest plots. To obtain more robust results, all data were pooled and analyzed using a randomeffects model, while a fixed-effects model was applied when the number of included studies was small (no more than 5) [39]. In cases where a multiarm trial was included, the shared group was divided into subgroups of approximately equal size, 1 for each experimental group [40,41]. In addition, we assessed statistical heterogeneity across all included studies using the I^2 statistic and P value. When there were 10 or more studies, the Egger test was conducted to assess small-study effects, with a P value below .05 indicating the possible presence of such effects [42]. To evaluate the robustness and reliability of the pooled results, a sensitivity analysis was performed, using the one-study-out method. Statistical significance for the overall effect was established when the 2-tailed *P* value was less than .05.

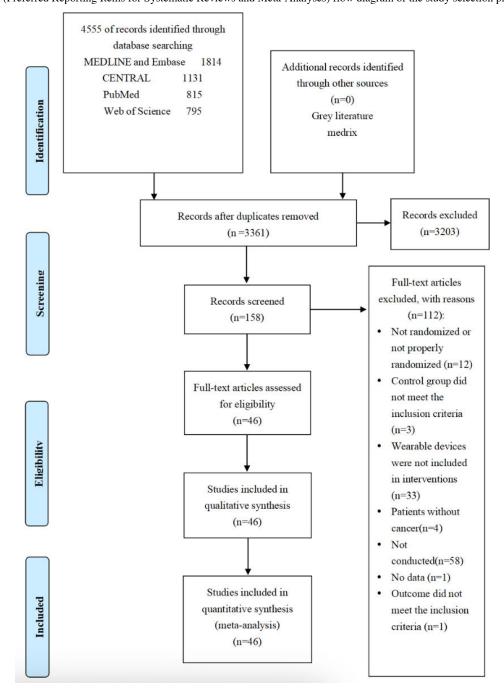
Results

Search Results and Selection

The initial search across 5 electronic databases identified 4555 articles. After 1194 duplicates were removed both

automatically and manually, 3361 articles were excluded based on their titles and abstracts. Following this initial screening, the full texts of the remaining 153 articles were retrieved, resulting in a final total of 46 studies included in the meta-analysis. The procedures for search and selection are delineated in Figure 1.

Figure 1. PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) flow diagram of the study selection process.



Description of Included Studies

Study Characteristics

The attributes of the 46 studies included in this analysis are presented in Multimedia Appendix 2. All studies, conducted

as randomized controlled trials, were published between 2005 and 2024 across 8 countries: the United States of America (31 studies) [43-73], Canada (1 study) [74], the United Kingdom (2 studies) [75,76], New Zealand (1 study) [77], Australia (6

studies) [78-83], the Netherlands (2 studies) [84,85], Korea (2 studies) [86,87] and China (1 study) [88].

Characteristics of Cancer Survivors

A total of 3727 cancer survivors were enrolled in the studies, with the number of participants ranging from 11 [65] to 412 [85]. The mean age of the included cancer survivors ranged from 12.7 (SD 7.87) years [53] to 73.79 (SD 7.74) years [65]. Regarding cancer types, 19 studies enrolled participants diagnosed with nonspecific types of cancer [45-47,53,54,56,59,67,72,73,75-80,82,85,88], while 27 focused solely on a single cancer type, including breast cancer (15 studies) [43,44,51,52,57,58,60,61,63,64,66,71,74,81,83], colorectal cancer (6 studies) [48,49,62,68,69,86], prostate cancer (4 studies) [55,65,70,87], leukemia or lymphoma (1 study) [50], and glioma (1 study) [84].

Characteristics of WEDS-Supported PA Programs

In the included studies, the intervention duration ranged from 4 weeks [45] to 48 weeks [51], with an average duration of 13.4 weeks. The WEDS-supported PA programs consisted of 2 components: WEDs and partnering tools.

WEDs play a role in step counting, reminders, and data storage. The WEDs used in these studies included pedometers (n=13) [43,56,57,61-63,65,66,75,77,79,85,86], smartwatches (n=4) [64,74,83,84], breath monitors (n=1) [45], smart bands (n=25) [44,46-52,54,55,58,59,67-73,76,78,80-82,87], intelligent sports bracelets (n=1) [88], headbands (n=1) [60], and activity monitors, with no mention of the specific type (n=1) [53]. There are some similarities between smart bands, intelligent sports bracelets, and smartwatches; however, intelligent sports bracelets are considered more fashionable due to their appearance resembling a traditional bracelet, while smart bands are slimmer and simpler in design, focusing primarily on fitness tracking and health monitoring [89]. In comparison, smartwatches have a watch-like form and offer more versatile functionalities, including apps and notifications [90].

Partnering tools in WEDS differ in their functions, including reminders, consultation, education, and data transmission for researchers. The types of partnering tools used included websites/web pages (n=5) [51,53,79,80,85], apps (n=7) [45,50,58,60,64,66,87], telephone calls (n=13) [43,56,57,61-63,65,71,74,75,77,83,88], SMS text messages (n=2) [48,76], or their combinations (n=19) [44,46,47,49,52,54,55,59,67-70,72,73,78,81,82,84,86].

In the 46 included articles, the behavior change techniques used in the interventions included goal setting, self-monitoring, feedback and monitoring, and social support. All interventions used goal setting, self-monitoring, and feedback and monitoring, while 8 studies [44,48,51,55,60,65,68,73] incorporated social support.

Characteristics of the Controls

Most of the patients in the control groups received usual care (n=35), which included education from HCPs (n=33)

or only access to websites or an app without reminders (n=2) [43,44,47-50,53-56,58-70,72,75,77,78,80-82,85-88]. Others were placed on a waiting list (n=11) to receive the respective interventions after the trials [45,46,51,52,57,71,73,74,79,83,84].

Outcome Measures

Outcome measures encompassed a diverse array, with assessments conducted at varying intervals and across different follow-up periods for participants.

Objectively Measured Moderate-to-Vigorous-Intensity Physical Activity

Researchers in 20 studies assessed objectively measured MVPA using an ActiGraph accelerometer [44,46,47,50,52-55,58,59,64,67-69,71,72,78,80-82,85]. Anderson et al [75] used a SenseWear PA monitor to assess patients' objectively measured PA, while Ferrante et al [51] used a Fitbit to evaluate patients' objectively measured PA.

Steps Per Day

Researchers in 11 studies assessed steps per day using an ActiGraph accelerometer [44,46,47,50,55,57,64,67,68,72,81]. Anderson et al [75] used a SenseWear PA monitor to assess steps per day, while Ferrante et al [51] and Walsh et al [76] used Fitbit devices. In addition, a pedometer was used to evaluate steps per day by Sajid et al [65] and Frensham et al [79].

Sedentary Behavior

Overview

Researchers in 13 studies used ActiGraph accelerometers to assess sedentary behavior [46,50,54,58,59,64,67,71,81,82].

Subjectively Measured Physical Activity

Six scales were used in 17 studies to assess cancer survivors' subjectively measured PA: the International Physical Activity Questionnaire Short Form [75], the Community Healthy Activities Model Program for Seniors [49,57,62,77], the International Physical Activity Questionnaire [66,84,87], the Short Questionnaire to Assess Health-Enhancing Physical Activity [85], the Godin Leisure-Time Exercise Questionnaire [67,76,86], and the Seven-Day Physical Activity Recall [43,56,61-63].

Quality of Life

Nine scales were used to assess the QoL of cancer survivors in 22 studies: the Patient-Reported Outcome Measurement Information System [50], the 36-item Short Form Health Survey—Physical Component [46,61,73,77,79], the Quality of Life in Adult Cancer Survivors [51], the EORTC QLG Core Questionnaire-30 [56,60,66,85,87], the Functional Assessment of Cancer Therapy—General [48,88], the Functional Assessment of Cancer Therapy—Breast [44,58,74], the Functional Assessment of Cancer Therapy—Colorectal [62,86], the RAND-36 Measure of Health-Related

Quality of Life [76], and the Pediatric Quality of Life Inventory [53,59].

Feasibility

Researchers in 12 studies reported feasibility, which was assessed by retention rate (n=8), wearing time (n=2), whether steps per day improved or not (n=1), and adherence to interventions (n=1) [39-41,44,45,48,49,59,62,63,67,69,77,78].

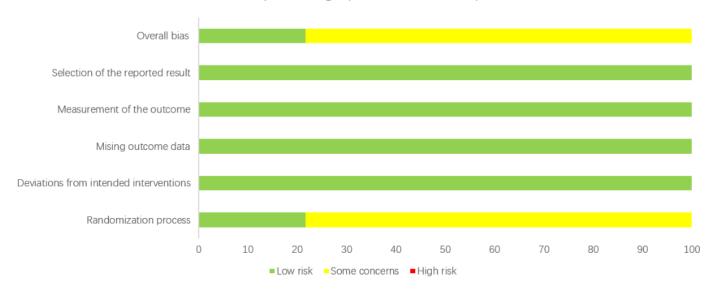
Risk of Bias

Utilizing the revised Cochrane risk-of-bias tool, the 24 studies that utilized intention-to-treat analysis within the inclusion criteria were classified as follows: 6 (25%) studies [44,51,55,77,78,84] were deemed to have a low risk of bias, while 18 (75%) studies [43,46,48,54,57,59,61,62,67,70,71,74,76,80,82,85,86,88] were identified as having some concerns regarding bias.

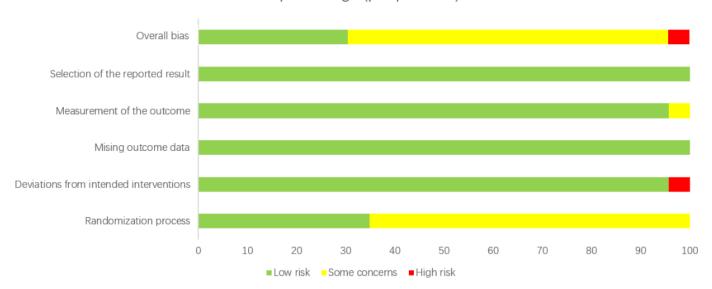
Furthermore, among the studies that used per-protocol analysis (N=22), 6 (27%) studies [47,49,64,75,79,81] were assessed as having a low risk of bias, whereas 15 (68%) studies [50,52,53,56,58,60,63,65,66,68,69,72,73,83,87] were classified as having some concerns, and 1 (5%) study [45] was classified as having a high risk of bias. Concerns regarding risk of bias emerged due to the randomization process (33/47 studies) [43,46,48,50,52-54,56-63,65-74,76,80,82,83,85-88] and the measurement of the outcome (1 of 47 studies) [52]. A high risk of bias was associated with deviation from the intended interventions (1 of 47 studies) [45]. The assessments of risk of bias are comprehensively presented in Figure 2. In addition, the results of the Egger test revealed no evidence of small study effects (objectively measured MVPA: P=.26; subjectively reported PA: P=.09; steps per day: P=.12; sedentary behavior: P=.15; BMI: P=.13; QoL: P=.24; Multimedia Appendix 3).

Figure 2. Results of the assessments of the risk of bias.





As precentage (per protocol)



Meta-Analysis Results

The summary of all outcomes included in this meta-analysis is detailed in Multimedia Appendix 3.

Primary Outcome: Objectively Measured MVPA Total Effects of WEDS-Supported PA Programs

Investigators from 23 studies, encompassing a total of 1853 participants, quantified the influence of WEDS-supported PA

programs on the objectively reported MVPA among cancer survivors. The random-effects model used for pooling the data yielded a significant improvement in the intervention groups (SMD 0.66, 95% CI 0.47-0.86, P<.001, I²=69%; Figure 3). Additionally, the meta-analysis results remained stable after the omission of individual studies (Multimedia Appendix 4).

Figure 3. Total effects on objectively measured moderate-to-vigorous-intensity physical activity [43,46,48,49,51-54,57,58,62,63,66-68,70,71,78-81,84]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
Dt	0.4	00.00	04.0000	00	20.00	74.0000	1 : -	4.40	10.40.4.701	4.40/
Bertram et al., 2019	24	69.00	84.0000	23	-20.00	74.0000		1.10	[0.49; 1.72]	4.1%
Chan J et al., 2020	38	30.00	60.0000	36	0.00	56.2500		0.51	[0.05; 0.97]	4.9%
Chow et al., 2021	24	24.50	16.4589	17	1.20	15.8824		1.41	[0.71; 2.11]	3.7%
Golstejin et al., 2018	226	60.00	223.3900	204	-82.00	272.2400	-	0.57	[0.38; 0.77]	6.4%
Hardcastle et al., 2024	33	68.70	76.7095	41	18.90	75.8778		0.65	[0.18; 1.12]	4.9%
Hardcastle et al., 2021	33	66.50	186.9230	29	-14.40	134.2080		0.49	[-0.02; 0.99]	4.7%
Hartman et al., 2018	43	14.20	13.9800	44	-0.70	9.7400	1 1 1	1.23	[0.77; 1.69]	5.0%
Howell et al., 2018	52	4.70	119.9000	24	-24.30	89.7000	 	0.26	[-0.23; 0.74]	4.8%
Johnson et al., 2022	26	0.00	0.2000	23	-0.20	0.2000		0.98	[0.39; 1.58]	4.2%
Kenfield et al., 2019	32	1.60	3.8500	32	-5.50	4.5750	-	1.66	[1.09; 2.23]	4.3%
Lynch et al., 2019	40	66.50	103.9662	40	-2.10	103.9662	- 8	0.65	[0.20; 1.10]	5.0%
Maxwell-Smith et al., 2018	34	45.00	123.2400	34	-21.00	108.9000	-	0.56	[0.08; 1.05]	4.8%
McNeil et al., 2019-Lower PA	15	42.00	32.5038	7	6.00	34.7513		1.04	[0.08; 2.00]	2.6%
McNeil et al., 2019-Higher PA	14	24.00	31.1752	6	6.00	34.7513		0.54	[-0.44; 1.51]	2.6%
Mendoza et al., 2017	29	4.40	25.6323	30	5.00	23.1651	—— i	-0.02	[-0.53; 0.49]	4.7%
Phillips et al., 2024	25	2.90	13.5000	24	-2.10	14.2100	- 1 1 1	0.36	[-0.21; 0.92]	4.4%
Pinto et al., 2021	12	71.02	144.1210	8	-34.00	64.9463	-	0.84	[-0.10; 1.78]	2.7%
Pinto et al., 2015	36	56.90	57.6221	32	2.20	28.6658		1.17	[0.65; 1.68]	4.6%
Pope et al., 2018	12	3.50	16.6460	8	7.60	18.6580	- 1	-0.22	[-1.12; 0.67]	2.8%
Valle et al., 2023	140	24.67	59.7645	140	11.41	60.1871	- !	0.22	[-0.01; 0.46]	6.2%
Van et al., 2022	17	-14.70	52.8000	20	-8.30	55.6000	i	-0.12	[-0.76; 0.53]	3.9%
Van et al., 2019	20	13.70	42.3864	19	3.70	23.0883		0.28	[-0.35; 0.92]	4.0%
Weiner et al., 2019	43	14.20	14.0000	44	-0.70	9.8000	- 10	1.22	[0.76; 1.68]	5.0%
Random effects model	968			885			🗼	0.66	[0.47; 0.86]	100.0%
Heterogeneity: $I^2 = 69\%$, $\tau^2 = 0.148$		1					-2 -1 0 1 2			

Subgroup Analysis

Studies grouped by the use of multipartnering tools suggested that WEDS-supported PA programs, whether with both multipartnering tools (SMD 0.68, 95% CI 0.44-0.92, P<.001, I²=70%) or without (SMD 0.63, 95% CI 0.26-1.01, P<.001, I²=71%), showed significant improvements in objectively measured MVPA (Figure 4).

Upon categorizing the studies based on the duration of the intervention, the pooled results indicated that WEDS-supported PA programs with long-term durations

(\geq 12 weeks) were effective in increasing objectively measured MVPA (SMD 0.72, 95% CI 0.53-0.92, P<.001, I^2 =67%; Figure 5). When grouped by whether the intervention was designed for a specific cancer type, both the "yes" group (SMD 0.74, 95% CI 0.52-0.96, P<.001, I^2 =64%) and the "no" group (SMD 0.37, 95% CI 0.04-0.7, P=.02, I^2 =52%) showed significant differences (Figure 6). Heterogeneity in these 2 subgroups showed a modest to notable decrease (Multimedia Appendix 2). Duration and whether the intervention was designed for patients with a specific cancer type may be sources of heterogeneity.

Figure 4. Subgroup analysis on objectively measured moderate-to-vigorous-intensity physical activity, grouped by the use of multipartnering tools [43,46,48,49,51-54,57,58,62,63,66-68,70,71,78-81,84]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
							1 .			
Yes			04.0000		00.00	74.0000	i			4.40/
Bertram et al., 2019	24	69.00	84.0000	23	-20.00	74.0000		1.10	[0.49; 1.72]	4.1%
Chan J et al., 2020	38	30.00	60.0000	36	0.00	56.2500		0.51	[0.05; 0.97]	4.9%
Chow et al., 2021	24	24.50	16.4589	17	1.20	15.8824	<u> </u>	1.41	[0.71; 2.11]	3.7%
Hardcastle et al., 2024	33	68.70	76.7095	41	18.90	75.8778		0.65	[0.18; 1.12]	4.9%
Hartman et al., 2018	43	14.20	13.9800	44	-0.70	9.7400	i	1.23	[0.77; 1.69]	5.0%
Johnson et al., 2022	26	0.00	0.2000	23	-0.20	0.2000		0.98	[0.39; 1.58]	4.2%
Kenfield et al., 2019	32	1.60	3.8500	32	-5.50	4.5750	<u>i</u>	1.66	[1.09; 2.23]	4.3%
Lynch et al., 2019	40	66.50	103.9662	40	-2.10	103.9662		0.65	[0.20; 1.10]	5.0%
Maxwell-Smith et al., 2018	34	45.00	123.2400	34	-21.00	108.9000	-	0.56	[0.08; 1.05]	4.8%
McNeil et al., 2019-Lower PA	15	42.00	32.5038	7	6.00	34.7513		1.04	[0.08; 2.00]	2.6%
McNeil et al., 2019-Higher PA	14	24.00	31.1752	6	6.00	34.7513		0.54	[-0.44; 1.51]	2.6%
Mendoza et al., 2017	29	4.40	25.6323	30	5.00	23.1651	 !	-0.02	[-0.53; 0.49]	4.7%
Phillips et al., 2024	25	2.90	13.5000	24	-2.10	14.2100		0.36	[-0.21; 0.92]	4.4%
Pinto et al., 2021	12	71.02	144.1210	8	-34.00	64.9463		0.84	[-0.10; 1.78]	2.7%
Valle et al., 2023	140	24.67	59.7645	140	11.41	60.1871	-	0.22	[-0.01; 0.46]	6.2%
Van et al., 2022	17	-14.70	52.8000	20	-8.30	55.6000	-	-0.12	[-0.76; 0.53]	3.9%
Van et al., 2019	20	13.70	42.3864	19	3.70	23.0883		0.28	[-0.35; 0.92]	4.0%
Random effects model	566			544			•	0.68	[0.44; 0.92]	71.7%
Heterogeneity: $I^2 = 70\%$, $\tau^2 = 0.16$	25, p < 0.0	1					!			
							!			
No Colotella et al. 2010	000	60.00	200 2000	20.4	00.00	070.0400		0.57	10.00.0.771	0.40/
Golstejin et al., 2018	226	60.00	223.3900	204	-82.00	272.2400		0.57	[0.38; 0.77]	6.4%
Hardcastle et al., 2021	33	66.50	186.9230	29	-14.40	134.2080		0.49	[-0.02; 0.99]	4.7%
Howell et al., 2018	52	4.70	119.9000	24	-24.30	89.7000		0.26	[-0.23; 0.74]	4.8%
Pinto et al., 2015	36	56.90	57.6221	32	2.20	28.6658	_	1.17	[0.65; 1.68]	4.6%
Pope et al., 2018	12	3.50	16.6460	8	7.60	18.6580		-0.22	[-1.12; 0.67]	2.8%
Weiner et al., 2019	43	14.20	14.0000	44	-0.70	9.8000		1.22	[0.76; 1.68]	5.0%
Random effects model	402			341				0.64	[0.26; 1.01]	28.3%
Heterogeneity: $I^2 = 71\%$, $\tau^2 = 0.15$	82, p < 0.0	1					!			
Random effects model	968			885			→	0.66	[0.47; 0.86]	100.0%
Heterogeneity: $I^2 = 69\%$, $\tau^2 = 0.14$	85, p < 0.0	1								
Test for subgroup differences: χ ²	0.03, df =	1 (p = 0.86	3)				-2 -1 0 1 2			

Figure 5. Subgroup analysis on objectively measured moderate-to-vigorous-intensity physical activity, grouped by intervention duration [43,46,48,49,51-54,57,58,62,63,66-68,70,71,78-81,84]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
≥12 weeks	202000	0.000.000.00	76.0.7000330	2027.209	1920200000	0.0000000000000000000000000000000000000	l i –	70027099938		
Bertram et al., 2019	24	69.00	84.0000	23	-20.00	74.0000		1.10	[0.49; 1.72]	4.1%
Chan J et al., 2020	38	30.00	60.0000	36	0.00	56.2500		0.51	[0.05; 0.97]	4.9%
Chow et al., 2021	24	24.50	16.4589	17	1.20	15.8824		1.41	[0.71; 2.11]	3.7%
Golstejin et al., 2018	226	60.00	223.3900	204	-82.00	272.2400	-	0.57	[0.38; 0.77]	6.4%
Hardcastle et al., 2024	33	68.70	76.7095	41	18.90	75.8778		0.65	[0.18; 1.12]	4.9%
Hardcastle et al., 2021	33	66.50	186.9230	29	-14.40	134.2080	i _	0.49	[-0.02; 0.99]	4.7%
Hartman et al., 2018	43	14.20	13.9800	44	-0.70	9.7400	_ i	1.23	[0.77; 1.69]	5.0%
Howell et al., 2018	52	4.70	119.9000	24	-24.30	89.7000		0.26	[-0.23; 0.74]	4.8%
Johnson et al., 2022	26	0.00	0.2000	23	-0.20	0.2000		0.98	[0.39; 1.58]	4.2%
Kenfield et al., 2019	32	1.60	3.8500	32	-5.50	4.5750		1.66	[1.09; 2.23]	4.3%
Lynch et al., 2019	40	66.50	103.9662	40	-2.10	103.9662	- 	0.65	[0.20; 1.10]	5.0%
Maxwell-Smith et al., 2018	34	45.00	123.2400	34	-21.00	108.9000	-	0.56	[0.08; 1.05]	4.8%
McNeil et al., 2019-Lower PA	15	42.00	32.5038	7	6.00	34.7513	- 10	1.04	[0.08; 2.00]	2.6%
McNeil et al., 2019-Higher PA	14	24.00	31.1752	6	6.00	34.7513		0.54	[-0.44; 1.51]	2.6%
Phillips et al., 2024	25	2.90	13.5000	24	-2.10	14.2100		0.36	[-0.21; 0.92]	4.4%
Pinto et al., 2021	12	71.02	144.1210	8	-34.00	64.9463	100	0.84	[-0.10; 1.78]	2.7%
Pinto et al., 2015	36	56.90	57.6221	32	2.20	28.6658	- B	1.17	[0.65; 1.68]	4.6%
Valle et al., 2023	140	24.67	59.7645	140	11.41	60.1871	-	0.22	[-0.01; 0.46]	6.2%
Van et al., 2022	17	-14.70	52.8000	20	-8.30	55.6000		-0.12	[-0.76; 0.53]	3.9%
Van et al., 2019	20	13.70	42.3864	19	3.70	23.0883		0.28	[-0.35; 0.92]	4.0%
Weiner et al., 2019	43	14.20	14.0000	44	-0.70	9.8000	i — B	1.22	[0.76; 1.68]	5.0%
Random effects model	927			847			-	0.72	[0.53; 0.92]	92.5%
Heterogeneity: $I^2 = 67\%$, $\tau^2 = 0.124$	48, p < 0.0	1								
<12 weeks							l i			
Mendoza et al., 2017	29	4.40	25.6323	30	5.00	23.1651		-0.02	[-0.53; 0.49]	4.7%
Pope et al., 2018	12	3.50	16.6460	8	7.60	18.6580		-0.22	[-1.12; 0.67]	2.8%
Random effects model	41			38			- i	-0.07	[-0.52; 0.37]	7.5%
Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, $p =$	0.70						!			
Random effects model	968			885			—	0.66	[0.47; 0.86]	100.0%
Heterogeneity: $I^2 = 69\%$, $\tau^2 = 0.148$	85, p < 0.0	1								
Test for subgroup differences: χ_1^2 =	10.41, df	= 1 (p < 0.6)	01)				-2 -1 0 1 2			

Figure 6. Subgroup analysis on objectively measured moderate-to-vigorous-intensity physical activity, grouped by whether the intervention was designed for a specific cancer type [43,46,48,49,51-54,57,58,62,63,66-68,70,71,78-81,84]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
							1 1			
Yes							l i -			
Bertram et al., 2019	24	69.00	84.0000	23	-20.00	74.0000		1.10	[0.49; 1.72]	4.1%
Chan J et al., 2020	38	30.00	60.0000	36	0.00	56.2500		0.51	[0.05; 0.97]	4.9%
Chow et al., 2021	24	24.50	16.4589	17	1.20	15.8824		1.41	[0.71; 2.11]	3.7%
Golstejin et al., 2018	226	60.00	223.3900	204	-82.00	272.2400		0.57	[0.38; 0.77]	6.4%
Hardcastle et al., 2024	33	68.70	76.7095	41	18.90	75.8778		0.65	[0.18; 1.12]	4.9%
Hardcastle et al., 2021	33	66.50	186.9230	29	-14.40	134.2080		0.49	[-0.02; 0.99]	4.7%
Hartman et al., 2018	43	14.20	13.9800	44	-0.70	9.7400	i	1.23	[0.77; 1.69]	5.0%
Kenfield et al., 2019	32	1.60	3.8500	32	-5.50	4.5750		1.66	[1.09; 2.23]	4.3%
Lynch et al., 2019	40	66.50	103.9662	40	-2.10	103.9662	- - 11	0.65	[0.20; 1.10]	5.0%
Maxwell-Smith et al., 2018	34	45.00	123.2400	34	-21.00	108.9000	- E	0.56	[0.08; 1.05]	4.8%
McNeil et al., 2019-Lower PA	15	42.00	32.5038	7	6.00	34.7513		1.04	[0.08; 2.00]	2.6%
McNeil et al., 2019-Higher PA	14	24.00	31.1752	6	6.00	34.7513	- 1	0.54	[-0.44; 1.51]	2.6%
Phillips et al., 2024	25	2.90	13.5000	24	-2.10	14.2100		0.36	[-0.21; 0.92]	4.4%
Pinto et al., 2015	36	56.90	57.6221	32	2.20	28.6658	-	1.17	[0.65; 1.68]	4.6%
Pope et al., 2018	12	3.50	16.6460	8	7.60	18.6580		-0.22	[-1.12; 0.67]	2.8%
Van et al., 2022	17	-14.70	52.8000	20	-8.30	55.6000		-0.12	[-0.76; 0.53]	3.9%
Van et al., 2019	20	13.70	42.3864	19	3.70	23.0883		0.28	[-0.35; 0.92]	4.0%
Weiner et al., 2019	43	14.20	14.0000	44	-0.70	9.8000	!	1.22	[0.76; 1.68]	5.0%
Random effects model	709			660			-	0.74	[0.53; 0.96]	77.5%
Heterogeneity: $I^2 = 64\%$, $\tau^2 = 0.135$	6, p < 0.0	1					l i			
							l i			
No							l i			
Howell et al., 2018	52	4.70	119.9000	24	-24.30	89.7000		0.26	[-0.23; 0.74]	4.8%
Johnson et al., 2022	26	0.00	0.2000	23	-0.20	0.2000		0.98	[0.39; 1.58]	4.2%
Mendoza et al., 2017	29	4.40	25.6323	30	5.00	23.1651		-0.02	[-0.53; 0.49]	4.7%
Pinto et al., 2021	12	71.02	144.1210	8	-34.00	64.9463	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	0.84	[-0.10; 1.78]	2.7%
Valle et al., 2023	140	24.67	59.7645	140	11.41	60.1871	i i	0.22	[-0.01; 0.46]	6.2%
Random effects model	259			225				0.37	[0.04; 0.70]	22.5%
Heterogeneity: $I^2 = 52\%$, $\tau^2 = 0.074$	$17, \rho = 0.0$	8					l i			
Random effects model	968			885			.	0.66	[0.47; 0.86]	100.0%
Heterogeneity: $I^2 = 69\%$, $\tau^2 = 0.148$	35, p < 0.0)1								
Test for subgroup differences: χ_1^2 =	3.44, df =	1 (p = 0.0	5)				-2 -1 0 1 2			

Secondary Outcomes: Subjectively Reported

Total Effects of WEDS-Supported PA Programs

Data gathered from 15 studies, involving a total of 2016 participants, were used to assess the efficacy of WEDS-supported PA programs in increasing subjectively

reported PA. The results of the random-effects model suggested a significant improvement in the experimental groups (SMD 0.5, 95% CI 0.23-0.77, P<.001, I^2 =79%; Figure 7). Furthermore, utilizing the one-study-out approach for sensitivity analysis, the findings remained stable (Multimedia Appendix 4).

Figure 7. Total effects on subjectively reported physical activity [42,48,55,56,60-62,65,66,75,76,83-86]. SMD: standardized mean difference.

		E	xperimental			Control	Standardised Mean		
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI
Bennett et al., 2006	27	1555.08	2117.7497	28	396.74	2174.1427	L==-	0.53	[-0.01; 1.07]
	1000						T		
Chan J et al., 2020	37	1.00	0.5000	36	0.00	0.5000		1.98	[1.41; 2.54]
Gehring et al., 2018	19	1200.00	4420.7847	10	811.00	5833.9410	-	0.08	[-0.69; 0.84]
Golstejin et al., 2017	249	280.00	747.2556	229	89.00	800.7328	 :	0.25	[0.07; 0.43]
Kim et al., 2019	37	235.60	280.8000	34	16.30	197.1000	+ -	0.89	[0.40; 1.38]
Ligible et al., 2011	48	54.50	142.0000	51	14.60	117.2000	 	0.30	[-0.09; 0.70]
Matthews et al., 2006	22	21.50	30.6000	14	5.00	13.8000		0.63	[-0.06; 1.32]
Park et al., 2020	75	1454.80	2595.0394	73	1002.80	2568.6301	-	0.17	[-0.15; 0.50]
Pinto et al., 2005	43	15.97	24.8110	43	-0.89	19.3472		0.75	[0.31; 1.19]
Pinto et al., 2015	36	97.70	63.6279	32	7.90	59.2719	-	1.44	[0.90; 1.98]
Pinto(B) et al., 2012	106	59.70	126.6205	86	30.82	119.8225		0.23	[-0.05; 0.52]
Pinto(C) et al., 2012	19	59.40	127.5774	24	68.30	135.0460		-0.07	[-0.67; 0.54]
Uhm et al., 2016	167	976.30	2350.9615	172	468.90	2135.6040		0.23	[0.01; 0.44]
Valle et al., 2023	140	123.28	171.0500	140	83.40	205.8600	i i	0.21	[-0.02; 0.45]
Walsh et al., 2021	11	3.61	19.5215	8	3.65	23.0054		-0.00	[-0.91; 0.91]
Random effects model	1036			980			.	0.50	[0.23; 0.77]
Heterogeneity: $I^2 = 79\%$, $\tau^2 =$	= 0.2245, µ	< 0.01							
							-2 -1 0 1 2		

Weight

6.3% 6.2%

5.0%

8.2% 6.6%

7.2% 5.5% 7.6%

6.4%

7.8%

6.0% 8.1%

8.0% 4.3% 100.0%

Subgroup Analysis

In the pooled analysis based on the use of multipartnering tools, the results in the subgroup without multipartnering tools (SMD 0.39, 95% CI 0.17-0.61, P<.001, I²=62%) showed a significant improvement in subjectively reported PA (Figure 8).

Upon categorizing the studies based on the duration of intervention, subjectively reported PA significantly increased in the long-term intervention groups (no less than 12 weeks;

SMD 0.52, 95% CI 0.24-0.81, P<.001, I^2 =80%; Figure 9). When grouped by whether the intervention was designed for a specific cancer type, patients' subjectively reported PA improved in both the "yes" group (SMD 0.56, 95% CI 0.23-0.89, P<.001, I^2 =82%) and the "no" group (SMD 0.25, 95% CI 0.04-0.06, P=.02, I^2 =0%; Figure 10). The use of multipartnering tools and whether interventions were designed for patients with specific cancer types may be sources of heterogeneity (Multimedia Appendix 2).

Figure 8. Subgroup analysis on subjectively reported physical activity, grouped by the use of multipartnering tools [42,48,55,56,60-62,65,66,75,76,83-86]. SMD: standardized mean difference.

		E	xperimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
No							1 2			
Bennett et al., 2006	27	1555.08	2117.7497	28	396.74	2174.1427		0.53	[-0.01; 1.07]	6.3%
Golstejin et al., 2017	249	280.00	747.2556	229	89.00	800.7328	!	0.25	[0.07; 0.43]	8.2%
Ligible et al., 2011	48	54.50	142.0000	51	14.60	117.2000	 	0.30	[-0.09; 0.70]	7.2%
Matthews et al., 2006	22	21.50	30.6000	14	5.00	13.8000		0.63	[-0.06; 1.32]	5.5%
Park et al., 2020	75	1454.80	2595.0394	73	1002.80	2568.6301		0.17	[-0.15; 0.50]	7.6%
Pinto et al., 2005	43	15.97	24.8110	43	-0.89	19.3472	- 1	0.75	[0.31; 1.19]	6.9%
Pinto et al., 2015	36	97.70	63.6279	32	7.90	59.2719		1.44	[0.90; 1.98]	6.4%
Pinto(B) et al., 2012	106	59.70	126.6205	86	30.82	119.8225	1	0.23	[-0.05; 0.52]	7.8%
Pinto(C) et al., 2012	19	59.40	127.5774	24	68.30	135.0460		-0.07	[-0.67; 0.54]	6.0%
Uhm et al., 2016	167	976.30	2350.9615	172	468.90	2135.6040	 	0.23	[0.01; 0.44]	8.1%
Walsh et al., 2021	11	3.61	19.5215	8	3.65	23.0054		-0.00	[-0.91; 0.91]	4.3%
Random effects model	803			760			•	0.39	[0.17; 0.61]	74.1%
Heterogeneity: $I^2 = 62\%$, $\tau^2 = 60\%$	= 0.0832, j	2 < 0.01					l j			
Yes										
Chan J et al., 2020	37	1.00	0.5000	36	0.00	0.5000		1.98	[1.41; 2.54]	6.2%
Gehring et al., 2018	19	1200.00	4420.7847	10	811.00	5833.9410		0.08	[-0.69; 0.84]	5.0%
Kim et al., 2019	37	235.60	280.8000	34	16.30	197.1000	1	0.89	[0.40; 1.38]	6.6%
Valle et al., 2023	140	123.28	171.0500	140	83.40	205.8600	} → -!	0.21	[-0.02; 0.45]	8.0%
Random effects model	233			220				0.79	[-0.05; 1.63]	25.9%
Heterogeneity: $I^2 = 92\%$, $\tau^2 = 10$	= 0.6620, j	> < 0.01								
Random effects model	1036			980			-	0.50	[0.23; 0.77]	100.0%
Heterogeneity: $I^2 = 79\%$, $\tau^2 = 79\%$	= 0.2245, /	< 0.01								
Test for subgroup difference	s: $\chi_1^2 = 0.8$	1. df = 1 (p =	0.37)				-2 -1 0 1 2			

Figure 9. Subgroup analysis on subjectively reported physical activity, grouped by intervention duration [42,48,55,56,60-62,65,66,75,76,83-86]. SMD: standardized mean difference.

		E	xperimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
≥12 weeks							1 !			
Bennett et al., 2006	27	1555.08	2117.7497	28	396.74	2174.1427		0.53	[-0.01; 1.07]	6.3%
Chan J et al., 2020	37	1.00	0.5000	36	0.00	0.5000	i 	1.98	[1.41; 2.54]	6.2%
Gehring et al., 2018	19	1200.00	4420.7847	10	811.00	5833.9410	- i	0.08	[-0.69; 0.84]	5.0%
Golstejin et al., 2017	249	280.00	747.2556	229	89.00	800.7328		0.25	[0.07; 0.43]	8.2%
Kim et al., 2019	37	235.60	280.8000	34	16.30	197.1000	÷ = -	0.89	[0.40; 1.38]	6.6%
Ligible et al., 2011	48	54.50	142.0000	51	14.60	117.2000	 	0.30	[-0.09; 0.70]	7.2%
Matthews et al., 2006	22	21.50	30.6000	14	5.00	13.8000	 	0.63	[-0.06; 1.32]	5.5%
Park et al., 2020	75	1454.80	2595.0394	73	1002.80	2568.6301		0.17	[-0.15; 0.50]	7.6%
Pinto et al., 2005	43	15.97	24.8110	43	-0.89	19.3472	+ -	0.75	[0.31; 1.19]	6.9%
Pinto et al., 2015	36	97.70	63.6279	32	7.90	59.2719	i 	1.44	[0.90; 1.98]	6.4%
Pinto(B) et al., 2012	106	59.70	126.6205	86	30.82	119.8225	l i	0.23	[-0.05; 0.52]	7.8%
Pinto(C) et al., 2012	19	59.40	127.5774	24	68.30	135.0460	- 1	-0.07	[-0.67; 0.54]	6.0%
Uhm et al., 2016	167	976.30	2350.9615	172	468.90	2135.6040	 -	0.23	[0.01; 0.44]	8.1%
Valle et al., 2023	140	123.28	171.0500	140	83.40	205.8600	·	0.21	[-0.02; 0.45]	8.0%
Random effects model	1025			972			-	0.52	[0.24; 0.81]	95.7%
Heterogeneity: $I^2 = 80\%$, $\tau^2 =$	= 0.2322, /	0 < 0.01								
<12 weeks										
Walsh et al., 2021	11	3.61	19.5215	8	3.65	23.0054	- •	-0.00	[-0.91; 0.91]	4.3%
Random effects model	1036			980			—	0.50	[0.23; 0.77]	100.0%
Heterogeneity: $I^2 = 79\%$, $\tau^2 =$	0.2245,	o < 0.01								
Test for subgroup differences	s: $\chi_1^2 = 1.1$	7, df = 1 (p =	0.28)				-2 -1 0 1 2			

Figure 10. Subgroup analysis on subjectively reported physical activity, grouped by whether the intervention was designed for a specific cancer type [42,48,55,56,60-62,65,66,75,76,83-86]. SMD: standardized mean difference.

		_	xperimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
No							1 !			
Bennett et al., 2006	27	1555.08	2117.7497	28	396.74	2174.1427	-	0.53	[-0.01; 1.07]	6.3%
Valle et al., 2023	140	123.28	171.0500	140	83.40	205.8600	 !	0.21	[-0.02; 0.45]	8.0%
Walsh et al., 2021	11	3.61	19.5215	8	3.65	23.0054		-0.00	[-0.91; 0.91]	4.3%
Random effects model	178			176			◆ i	0.25	[0.04; 0.46]	18.6%
Heterogeneity: $t^2 = 0\%$, $\tau^2 =$	0. p = 0.48	3					l i			
Yes										
Chan J et al., 2020	37	1.00	0.5000	36	0.00	0.5000	!	1.98	[1.41; 2.54]	6.2%
Gehring et al., 2018	19	1200.00	4420.7847	10	811.00	5833.9410		0.08	[-0.69; 0.84]	5.0%
Golstejin et al., 2017	249	280.00	747.2556	229	89.00	800.7328	· · · !	0.25	[0.07; 0.43]	8.2%
Kim et al., 2019	37	235.60	280.8000	34	16.30	197.1000	1 1	0.89	[0.40; 1.38]	6.6%
Ligible et al., 2011	48	54.50	142.0000	51	14.60	117.2000	+	0.30	[-0.09; 0.70]	7.2%
Matthews et al., 2006	22	21.50	30.6000	14	5.00	13.8000	4	0.63	[-0.06; 1.32]	5.5%
Park et al., 2020	75	1454.80	2595.0394	73	1002.80	2568.6301	-	0.17	[-0.15; 0.50]	7.6%
Pinto et al., 2005	43	15.97	24.8110	43	-0.89	19.3472		0.75	[0.31; 1.19]	6.9%
Pinto et al., 2015	36	97.70	63.6279	32	7.90	59.2719		- 1.44	[0.90; 1.98]	6.4%
Pinto(B) et al., 2012	106	59.70	126.6205	86	30.82	119.8225	 	0.23	[-0.05; 0.52]	7.8%
Pinto(C) et al., 2012	19	59.40	127.5774	24	68.30	135.0460		-0.07	[-0.67; 0.54]	6.0%
Uhm et al., 2016	167	976.30	2350.9615	172	468.90	2135.6040	- i	0.23	[0.01; 0.44]	8.1%
Random effects model	858			804			-	0.56	[0.23; 0.89]	81.4%
Heterogeneity: $I^2 = 82\%$, $\tau^2 =$	0.2807, p	0.01					l į			
Random effects model	1036			980			.	0.50	[0.23; 0.77]	100.0%

Secondary Outcomes: Steps Per Day

Total Effects of WEDS-Supported PA Programs

Fifteen studies assessed the impact of WEDS-supported PA programs on the steps per day of patients with cancer [44,46,47,50,55,57,64,65,67,68,72,75,76,79,81]. The

random-effects model revealed a significant difference between the intervention and control groups (SMD 0.54, 95% CI 0.14-0.94, P=.002, I²=81%; Figure 11). Utilizing the one-study-out approach for sensitivity analysis, the pooled findings remained robust upon the sequential exclusion of individual studies (Multimedia Appendix 4).

Figure 11. Total effects on steps per day [45,46,49,50,54,56,63,64,66-68,71,74,75,78,80]. SMD: standardized mean difference.

Co	ol Standardised Mean	
Mean	D Difference SMD 95%-CI	Weight
699.00 5167.	0.23 [-0.27; 0.74]	6.8%
-398.00 1751.	00 1.01 [0.40; 1.62]	6.4%
233.00 2402.	2 0.59 [-0.09; 1.27]	6.2%
260.00 2020.	0.20 [-0.42; 0.83]	6.4%
-205.47 2147.	0.04 [-0.62; 0.71]	6.3%
-978.00 560.0	3.43 [2.64; 4.21]	5.9%
308.00 2482.0	0.40 [-0.04; 0.85]	6.9%
-559.10 1326.	0.81 [0.11; 1.51]	6.2%
-689.10 2446.	0.42 [-0.15; 0.98]	6.6%
-1209.18 1736.	77 0.96 [0.01; 1.91]	5.3%
334.40 1870.	0.00 [-0.89; 0.90]	5.5%
-383.40 1583.	0.93 [-0.35; 2.22]	4.3%
299.92 2052.	0.19 [-0.04; 0.43]	7.4%
-431.00 2993.0	-0.23 [-0.88; 0.42]	6.3%
712.00 4962.	0.07 [-0.56; 0.70]	6.4%
257.67 3916.	0.00 [-0.35; 0.36]	7.2%
	0.54 [0.14; 0.94]	100.0%
		0.54 [0.14; 0.94] -4 -2 0 2 4

Subgroup Analysis

In the pooled results of the partnering tools used in WEDS-supported PA programs, compared with the subgroup without multipartnering tools, the subgroup using multipartnering showed a significant difference in the number of steps per day (SMD 0.59, 95% CI 0.07-1.1, P=.006, I²=85%; Figure 12). Heterogeneity decreased in the subgroup without multipartnering tools (I²=46%).

The pooled findings from the subgroup analysis indicated a significant increase in steps per day when the duration of WEDS-supported PA programs was no less than 12 weeks (SMD 0.55, 95% CI 0.11-0.99, P=.003 I²=83%; Figure 13). Heterogeneity decreased in the group with a duration of less than 12 weeks (I²=27%).

When grouped by whether the intervention was designed for a specific cancer type, patients' steps per day improved in the "yes" group (SMD 0.59, 95% CI 0.1-1.08, P=.008, I²=83%; Figure 14). Heterogeneity decreased in the group without specific cancer types (I²=43%).

The use of multipartnering tools, intervention duration, and whether patients were allocated based on specific cancer types may be sources of heterogeneity.

Figure 12. Subgroup analysis on steps per day, grouped by the use of multipartnering tools [45,46,49,50,54,56,63,64,66-68,71,74,75,78,80]. SMD: standardized mean difference.

		E	xperimental			Control		Standard	lised Mean			
Study	Total	Mean	SD	Total	Mean	SD		Diffe	erence	SMD	95%-CI	Weight
Yes									H			
Anderson et al., 2018	30	1760.00	3619.3835	30	699.00	5167.2959		-		0.23	[-0.27; 0.74]	6.8%
Bertram et al., 2019	24	1470.00	1881.0000	23	-398.00	1751.0000			1	1.01	[0.40; 1.62]	6.4%
Blair et al., 2021	17	1675.00	2394.2310	18	233.00	2402.0272			-	0.59	[-0.09; 1.27]	6.2%
Chow et al., 2021	24	711.00	2278.2012	17	260.00	2020.8010		-		0.20	[-0.42; 0.83]	6.4%
Ferrante et al., 2020	18	-107.07	2184.9400	17	-205.47	2147.7900		-	i i	0.04	[-0.62; 0.71]	6.3%
Kenfield et al., 2019	32	849.00	490.5000	32	-978.00	560.0000			Ti -	3.43	[2.64; 4.21]	5.9%
Lynch et al., 2019	40	1241.00	2064.6300	40	308.00	2482.6800			-	0.40	[-0.04; 0.85]	6.9%
Phillips et al., 2019	25	335.80	2387.0000	24	-689.10	2446.0000			+	0.42	[-0.15; 0.98]	6.6%
Pinto et al., 2021	12	1487.79	3156.9956	8	-1209.18	1736.7177				0.96	[0.01; 1.91]	5.3%
Valle et al., 2023	140	670.69	1796.6000	140	299.92	2052.7000				0.19	[-0.04; 0.43]	7.4%
Van et al., 2022	17	-1139.00	3058.0000	20	-431.00	2993.0000			-!	-0.23	[-0.88; 0.42]	6.3%
Van et al., 2019	20	1040.00	4111.7450	19	712.00	4962.5700		_	-	0.07	[-0.56; 0.70]	6.4%
Random effects model	399			388						0.59	[0.07; 1.10]	76.9%
Heterogeneity: I^2 = 85%, τ^2 =	= 0.7235, /	o < 0.01							i			
No												
Matthews et al., 2007	22	1152.50	2408.8000	14	-559.10	1326.5000				0.81	[0.11; 1.51]	6.2%
Pope et al., 2018	12	342.70	2105.8209	8	334.40	1870.5392		_	 	0.00	[-0.89; 0.90]	5.5%
Sajid et al., 2016	6	1950.40	2713.2009	5	-383.40	1583.7820		_	1 -	0.93	[-0.35; 2.22]	4.3%
Walsh et al., 2021	61	275.65	3916.7265	62	257.67	3916.7300		4	1	0.00	[-0.35; 0.36]	7.2%
Random effects model	101			89						0.32	[-0.17; 0.81]	23.1%
Heterogeneity; $I^2 = 46\%$, $\tau^2 =$	= 0.1136. /	0 = 0.14							l i			
Random effects model	500			477					•	0.54	[0.14; 0.94]	100.0%
Heterogeneity: $I^2 = 81\%$, $\tau^2 =$	= 0.5493,	o < 0.01							1 1	1		
Test for subgroup differences	s: $\chi_1^2 = 0.5$	3, df = 1 (p =	0.47)				-4	-2	0 2	4		

Figure 13. Subgroup analysis on steps per day, grouped by intervention duration [45,46,49,50,54,56,63,64,66-68,71,74,75,78,80]. SMD: standardized mean difference.

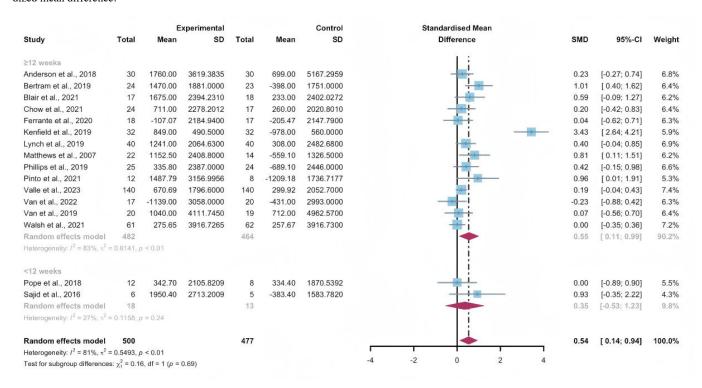


Figure 14. Subgroup analysis on steps per day, grouped by whether the intervention was designed for a specific cancer type [45,46,49,50,54,56,63,64,66-68,71,74,75,78,80]. SMD: standardized mean difference.

		E	xperimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
/es							Li			
Anderson et al., 2018	30	1760.00	3619.3835	30	699.00	5167.2959	-	0.23	[-0.27; 0.74]	6.8%
Bertram et al., 2019	24	1470.00	1881.0000	23	-398.00	1751.0000	1	1.01	[0.40; 1.62]	6.4%
Blair et al., 2021	17	1675.00	2394.2310	18	233.00	2402.0272	-	0.59	[-0.09; 1.27]	6.2%
Chow et al., 2021	24	711.00	2278.2012	17	260.00	2020.8010	— 	0.20	[-0.42; 0.83]	6.4%
errante et al., 2020	18	-107.07	2184.9400	17	-205.47	2147.7900	- - 1	0.04	[-0.62; 0.71]	6.3%
Cenfield et al., 2019	32	849.00	490.5000	32	-978.00	560.0000	l i —	3.43	[2.64; 4.21]	5.9%
ynch et al., 2019	40	1241.00	2064.6300	40	308.00	2482.6800	H 1	0.40	[-0.04; 0.85]	6.9%
Natthews et al., 2007	22	1152.50	2408.8000	14	-559.10	1326.5000		0.81	[0.11; 1.51]	6.2%
hillips et al., 2019	25	335.80	2387.0000	24	-689.10	2446.0000	+	0.42	[-0.15; 0.98]	6.6%
ope et al., 2018	12	342.70	2105.8209	8	334.40	1870.5392	I	0.00	[-0.89; 0.90]	5.5%
ajid et al., 2016	6	1950.40	2713.2009	5	-383.40	1583.7820		0.93	[-0.35; 2.22]	4.3%
an et al., 2022	17	-1139.00	3058.0000	20	-431.00	2993.0000	i	-0.23	[-0.88; 0.42]	6.3%
an et al., 2019	20	1040.00	4111.7450	19	712.00	4962.5700	- i	0.07	[-0.56; 0.70]	6.4%
tandom effects model	287			267			-	0.59	[0.10; 1.08]	80.1%
leterogeneity: $I^2 = 83\%$, $\tau^2 =$	0.6827, /	0.01								
No										
Pinto et al., 2021	12	1487.79	3156.9956	8	-1209.18	1736.7177	1	0.96	[0.01; 1.91]	5.3%
/alle et al., 2023	140	670.69	1796.6000	140	299.92	2052.7000	-	0.19	[-0.04; 0.43]	7.4%
Valsh et al., 2021	61	275.65	3916.7265	62	257.67	3916.7300	- i	0.00	[-0.35; 0.36]	7.2%
Random effects model	213			210			∳ i	0.17	[-0.02; 0.36]	19.9%
Heterogeneity: $I^2 = 43\%$, $\tau^2 =$	< 0.0001	$\rho = 0.17$					l i			
Random effects model	500			477			→	0.54	[0.14; 0.94]	100.0%
Heterogeneity: $I^2 = 81\%$, $\tau^2 =$	0.5493, /	0.01				Г	-2 0 2			

Secondary Outcomes: Sedentary Behavior Total Effects of WEDS-Supported PA Programs

Data gathered from 13 studies, involving a total of 912 participants, were used to assess the efficacy of WEDS-supported PA programs in decreasing sedentary

behavior. The results of the random-effects model demonstrated that WEDS-supported PA programs did not significantly decrease cancer survivors' sedentary behavior (SMD -0.63, 95% CI -1.34 to 0.07, P=.08, I²=92%; Figure 15). Utilizing the one-study-out approach for sensitivity analysis, the findings remained robust (Multimedia Appendix 4).

Figure 15. Total effects on sedentary behavior [43,45,49,53,57,58,63,66,70,71,79-81]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
Blair et al., 2021	17	7.90	77.8221	18	-2.50	75.4090	:	0.13	[-0.53; 0.80]	7.2%
Chow et al., 2021	24	-4.10	77.4399	17	-0.50	68.6567	1	-0.05	[-0.67; 0.57]	7.3%
Hardcastle et al., 2024	33	-194.50	434.7341	41	-129.40	430.8720	<u> </u>	-0.15	[-0.61; 0.31]	7.4%
Hardcastle et al., 2021	33	-2.00	5.3584	29	-0.70	5.3894	4	-0.24	[-0.74; 0.26]	7.4%
Johnson et al., 2022	26	-52.40	12.1000	23	2.50	12.8000	i]	-4.34	[-5.40; -3.29]	6.6%
Lynch et al., 2019	37	-23.50	70.4824	40	13.10	75.6686	-	-0.49	[-0.95; -0.04]	7.4%
Maxwell-Smith et al., 2018	34	-3.00	5.7320	33	-3.00	7.0505	-	0.00	[-0.48; 0.48]	7.4%
McNeil et al., 2019-Lower PA	15	-54.00	70.4249	7	12.00	74.4671	 	-0.89	[-1.83; 0.06]	6.8%
McNeil et al., 2019-Higher PA	14	-6.00	67.5462	6	12.00	74.4671		-0.25	[-1.21; 0.71]	6.7%
Mendoza et al., 2017	29	-4.50	82.6806	30	1.00	56.6407	L in the second	-0.08	[-0.59; 0.43]	7.4%
Pinto et al., 2021	11	-290.67	828.3160	8	-781.12	1091.8827	!	0.50	[-0.43; 1.42]	6.8%
Pope et al., 2018	12	2.40	44.8920	8	0.40	54.0700	+	0.04	[-0.86; 0.93]	6.8%
Valle et al., 2023	140	-7.57	68.9706	140	-7.13	79.0219	i 😅	-0.01	[-0.24; 0.23]	7.6%
Weiner et al., 2019	43	-24.90	5.9000	44	-4.80	5.9000		-3.38	[-4.04; -2.71]	7.2%
Random effects model	468			444			-	-0.63	[-1.34; 0.07]	100.0%
Heterogeneity: $I^2 = 92\%$, $\tau^2 = 1.687$	72, ρ < 0.0)1					-4 -2 0 2 4			

Subgroup Analysis

In the pooled results for the subgroups of WEDS-supported PA programs, usage of multipartnering tools, durations of interventions, and whether interventions were designed

for specific cancer types, no significant differences were observed (Figures 16-18 and Multimedia Appendix 2).

Heterogeneity decreased in the group with a duration of less than 12 weeks ($I^2=0\%$).

Figure 16. Subgroup analysis on sedentary behaviors, grouped by the use of multipartnering tools [43,45,49,53,57,58,63,66,70,71,79-81]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
Yes							!]			
Blair et al., 2021	17	7.90	77.8221	18	-2.50	75.4090	<u> </u>	0.13	[-0.53; 0.80]	7.2%
Chow et al., 2021	24	-4.10	77.4399	17	-0.50	68.6567	8	-0.05	[-0.67; 0.57]	7.3%
Hardcastle et al., 2024	33	-194.50	434.7341	41	-129.40	430.8720	i i	-0.15	[-0.61; 0.31]	7.4%
Johnson et al., 2022	26	-52.40	12.1000	23	2.50	12.8000	- i	-4.34	[-5.40; -3.29]	6.6%
Lynch et al., 2019	37	-23.50	70.4824	40	13.10	75.6686	-	-0.49	[-0.95; -0.04]	7.4%
Maxwell-Smith et al., 2018	34	-3.00	5.7320	33	-3.00	7.0505	-	0.00	[-0.48; 0.48]	7.4%
McNeil et al., 2019-Lower PA	15	-54.00	70.4249	7	12.00	74.4671		-0.89	[-1.83; 0.06]	6.8%
McNeil et al., 2019-Higher PA	14	-6.00	67.5462	6	12.00	74.4671	1	-0.25	[-1.21; 0.71]	6.7%
Mendoza et al., 2017	29	-4.50	82.6806	30	1.00	56.6407		-0.08	[-0.59; 0.43]	7.4%
Pinto et al., 2021	11	-290.67	828.3160	8	-781.12	1091.8827	1 2	0.50	[-0.43; 1.42]	6.8%
Valle et al., 2023	140	-7.57	68.9706	140	-7.13	79.0219	i 🛗	-0.01	[-0.24; 0.23]	7.6%
Random effects model	380			363			*	-0.47	[-1.18; 0.24]	78.6%
Heterogeneity: $I^2 = 86\%$, $\tau^2 = 1.319$	94. p < 0.0	01					j			
No							!			
Hardcastle et al., 2021	33	-2.00	5.3584	29	-0.70	5.3894		-0.24	[-0.74; 0.26]	7.4%
Pope et al., 2018	12	2.40	44.8920	8	0.40	54.0700	1 1	0.04	[-0.86; 0.93]	6.8%
Weiner et al., 2019	43	-24.90	5.9000	44	-4.80	5.9000	- i	-3.38	[-4.04; -2.71]	7.2%
Random effects model	88			81				-1.20	[-3.35; 0.95]	21.4%
Heterogeneity: $I^2 = 97\%$, $\tau^2 = 3.472$	28, p < 0.0)1					į			
Random effects model	468			444				-0.63	[-1.34; 0.07]	100.0%
Heterogeneity: I^2 = 92%, τ^2 = 1.687	72, p < 0.0	01								
Test for subgroup differences: χ_1^2 =	0.40, df =	1 (p = 0.53))				-4 -2 0 2 4			

Figure 17. Subgroup analysis on sedentary behaviors, grouped by intervention duration [43,45,49,53,57,58,63,66,70,71,79-81]. SMD: standardized mean difference.

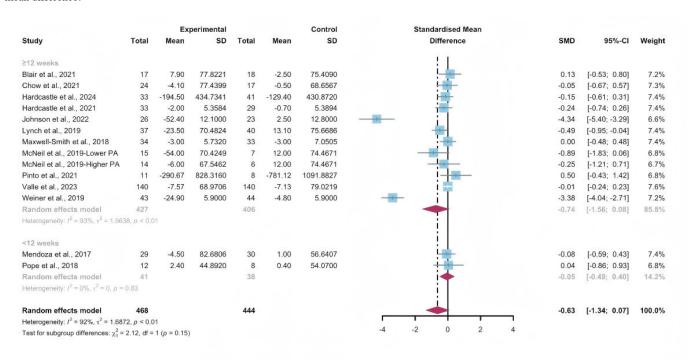


Figure 18. Subgroup analysis on sedentary behaviors, grouped by whether the intervention was designed for a specific cancer type [43,45,49,53,57,58,63,66,70,71,79-81]. SMD: standardized mean difference.

		Ex	perimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weight
Yes							! 1			
Blair et al., 2021	17	7.90	77.8221	18	-2.50	75.4090	!_ 	0.13	[-0.53; 0.80]	7.2%
Chow et al., 2021	24	-4.10	77.4399	17	-0.50	68.6567	1-1	-0.05	[-0.67; 0.57]	7.3%
Hardcastle et al., 2024	33	-194.50	434.7341	41	-129.40	430.8720	<u> </u>	-0.15	[-0.61; 0.31]	7.4%
Hardcastle et al., 2021	33	-2.00	5.3584	29	-0.70	5.3894	1	-0.24	[-0.74; 0.26]	7.4%
Lynch et al., 2019	37	-23.50	70.4824	40	13.10	75.6686	-	-0.49	[-0.95; -0.04]	7.4%
Maxwell-Smith et al., 2018	34	-3.00	5.7320	33	-3.00	7.0505	-	0.00	[-0.48; 0.48]	7.4%
McNeil et al., 2019-Lower PA	15	-54.00	70.4249	7	12.00	74.4671		-0.89	[-1.83; 0.06]	6.8%
McNeil et al., 2019-Higher PA	14	-6.00	67.5462	6	12.00	74.4671	- 1 - 1	-0.25	[-1.21; 0.71]	6.7%
Pope et al., 2018	12	2.40	44.8920	8	0.40	54.0700	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 	0.04	[-0.86; 0.93]	6.8%
Weiner et al., 2019	43	-24.90	5.9000	44	-4.80	5.9000	!	-3.38	[-4.04; -2.71]	7.2%
Random effects model	262			243			•	-0.52	[-1.17; 0.13]	71.7%
Heterogeneity: $I^Z = 90\%$, $\tau^Z = 0.979$	92, p < 0.0)1					i			
No										
Johnson et al., 2022	26	-52.40	12.1000	23	2.50	12.8000		-4.34	[-5.40; -3.29]	6.6%
Mendoza et al., 2017	29	-4.50	82.6806	30	1.00	56.6407	! 	-0.08	[-0.59; 0.43]	7.4%
Pinto et al., 2021	11	-290.67	828.3160	8	-781.12	1091.8827	!+	0.50	[-0.43; 1.42]	6.8%
Valle et al., 2023	140	-7.57	68.9706	140	-7.13	79.0219	1 22	-0.01	[-0.24; 0.23]	7.6%
Random effects model	206			201				-0.95	[-3.12; 1.21]	28.3%
Heterogeneity: $l^2 = 95\%$, $\tau^2 = 4.730$	0.04, p < 0.0)1					i l			
Random effects model	468			444				-0.63	[-1.34; 0.07]	100.0%
Heterogeneity: $I^2 = 92\%$, $\tau^2 = 1.687$	72, p < 0.0)1								
Test for subgroup differences: χ_1^2 =	0.14, df =	1(p = 0.71))				-4 -2 0 2 4			

Secondary Outcomes: BMI

Total Effects of WEDS-Supported PA Programs on BMI

Data gathered from 12 studies, involving a total of 1134 participants, were used to assess the efficacy of WEDS-supported PA programs in decreasing BMI. The

results of the random-effects model demonstrated no significant difference between the experimental and control groups (SMD -0.07, 95% CI -0.18 to 0.05, P=.27, $I^2=0\%$; Figure 19). In addition, the pooled findings remained robust upon the sequential exclusion of individual studies (Multimedia Appendix 4).

Figure 19. Total effects on BMI [42,50,51,57,65,74,75,77,79,81,83,86]. SMD: standardized mean difference.

		Expe	rimental			Control		Stan	dardised	Mean				
Study	Total	Mean	SD	Total	Mean	SD			Differenc	е		SMD	95%-CI	Weight
Anderson et al., 2018	30	-1.10	1.6100	30	-0.10	0.5400			!			-0.82	[-1.35; -0.29]	4.8%
Ferrante et al., 2020	18	-0.74	0.9900	17	-0.91	1.3900					-	0.14	[-0.53; 0.80]	3.0%
Frensham et al., 2018	46	-0.20	4.9000	45	0.00	4.4508			-			-0.04	[-0.45; 0.37]	7.9%
Gehring et al., 2018	19	-0.40	0.6224	9	0.00	0.9107		-	-11-			-0.54	[-1.34; 0.27]	2.0%
Hardcastle et al., 2021	33	-0.30	0.5640	29	-0.30	0.5258		_	- 11			0.00	[-0.50; 0.50]	5.4%
Hartman et al., 2018	43	0.10	0.5600	44	0.10	0.6000		_	- 10			0.00	[-0.42; 0.42]	7.5%
Maxwell-Smith et al., 2018	34	-0.44	0.6000	34	-0.47	0.6300		_	- 0			0.05	[-0.43; 0.52]	5.9%
McNeil et al., 2019-Lower PA	15	-0.10	1.0835	7	0.30	1.0756		-				-0.36	[-1.26; 0.55]	1.6%
McNeil et al., 2019-Higher PA	14	-0.02	1.0392	6	0.30	1.0756	(t)	-		-		-0.29	[-1.25; 0.67]	1.4%
Park et al., 2021	75	0.20	2.6058	73	0.10	2.9052		-	1 2	_		0.04	[-0.29; 0.36]	12.8%
Pinto et al., 2005	43	0.65	4.8401	43	0.75	5.4808			-			-0.02	[-0.44; 0.40]	7.5%
Uhm et al., 2017	167	0.00	3.1512	172	0.00	3.3511			-			0.00	[-0.21; 0.21]	29.4%
Walsh et al., 2021	61	-0.69	3.8987	62	-0.18	5.0452			-	-		-0.11	[-0.47; 0.24]	10.7%
Random effects model	598			571					4			-0.07	[-0.18; 0.05]	100.0%
Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, $\rho =$	0.49							1	-1-					
							-1	-0.5	0	0.5	1			

Subgroup Analysis (BMI)

In the pooled analysis of subgroups within WEDS-supported PA programs, considering factors such as the use of multipartnering tools, the duration of interventions, and

whether the interventions were tailored to specific cancer types, no significant differences were found (Figures 20 and 21 and Multimedia Appendix 2).

Figure 20. Subgroup analysis on BMI, grouped by the use of multipartnering tools [42,50,51,57,65,74,75,77,79,81,83,86]. SMD: standardized mean difference.

D Difference SMD 95%-0 -0.82 [-1.35; -0.28 0.0	4.8% 3.0% 2.0% 7.5%
0.0 0.14 [-0.53; 0.8] 0.07 -0.54 [-1.34; 0.2] 0.00 [-0.42; 0.4]	3.0% 2.0% 7.5%
0.0 0.14 [-0.53; 0.8] 0.07 -0.54 [-1.34; 0.2] 0.00 [-0.42; 0.4]	3.0% 2.0% 7.5%
-0.54 [-1.34; 0.2] 00	2.0% 7.5%
0.00 [-0.42; 0.43]	7.5%
· · ·	
0.05 [-0.43: 0.5]	1 E 09/
	0.970
-0.36 [-1.26; 0.56	1.6%
-0.29 [-1.25; 0.6]] 1.4%
-0.23 [-0.53; 0.08] 26.4%
-0.04 [-0.45; 0.3	7.9%
0.00 [-0.50; 0.50	5.4%
0.04 [-0.29; 0.30]	12.8%
0.02 [-0.44; 0.44]	7.5%
1 0.00 [-0.21; 0.2] 29.4%
-0.11 [-0.47; 0.24] 10.7%
	73.6%
-0.02 [-0.15; 0.13	
] 100.0%

Figure 21. Subgroup analysis on BMI, grouped by whether the intervention was designed for a specific cancer type [42,50,51,57,65,74,75,77,79,81,83,86]. SMD: standardized mean difference.

		Expe	rimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weigh
No							9			
Anderson et al., 2018	30	-1.10	1.6100	30	-0.10	0.5400		-0.82	[-1.35; -0.29]	4.89
Frensham et al., 2018	46	-0.20	4.9000	45	0.00	4.4508		-0.04	[-0.45; 0.37]	7.99
Walsh et al., 2021	61	-0.69	3.8987	62	-0.18	5.0452		-0.11	[-0.47; 0.24]	10.79
Random effects model	137			137				-0.29	[-0.74; 0.16]	23.39
Heterogeneity: $I^2 = 67\%$, $\tau^2 = 0.110$	$08, \rho = 0.0$)5					i			
Yes							il i			
Ferrante et al., 2020	18	-0.74	0.9900	17	-0.91	1.3900		0.14	[-0.53; 0.80]	3.09
Gehring et al., 2018	19	-0.40	0.6224	9	0.00	0.9107 —		-0.54	[-1.34; 0.27]	2.0
Hardcastle et al., 2021	33	-0.30	0.5640	29	-0.30	0.5258		0.00	[-0.50; 0.50]	5.4
Hartman et al., 2018	43	0.10	0.5600	44	0.10	0.6000		0.00	[-0.42; 0.42]	7.5
Maxwell-Smith et al., 2018	34	-0.44	0.6000	34	-0.47	0.6300		0.05	[-0.43; 0.52]	5.99
McNeil et al., 2019-Lower PA	15	-0.10	1.0835	7	0.30	1.0756 —		-0.36	[-1.26; 0.55]	1.6
McNeil et al., 2019-Higher PA	14	-0.02	1.0392	6	0.30	1.0756 —		-0.29	[-1.25; 0.67]	1.49
Park et al., 2021	75	0.20	2.6058	73	0.10	2.9052		0.04	[-0.29; 0.36]	12.89
Pinto et al., 2005	43	0.65	4.8401	43	0.75	5.4808		-0.02	[-0.44; 0.40]	7.5
Uhm et al., 2017	167	0.00	3.1512	172	0.00	3.3511	_	0.00	[-0.21; 0.21]	29.49
Random effects model	461			434			*	-0.01	[-0.15; 0.12]	76.79
Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, $\rho =$	0.97						i i			
Random effects model	598			571			4	-0.07	[-0.18; 0.05]	100.09
Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, $\rho =$	0.49									
Test for subgroup differences: χ_1^2 =	1.36, df =	1(p = 0.2)	24)				-1 -0.5 0 0.5 1			

Secondary Outcomes: Quality of Life Total Effects of WEDS-Supported PA Programs

Researchers from 21 studies assessed the effectiveness of WEDS-supported PA programs on the QoL of cancer survivors. The random-effects model revealed a significant

difference between the experimental and control groups in the pooled results (SMD 0.19, 95% CI 0.08-0.31, P<.001, I^2 =33%; Figure 22). A sensitivity analysis was conducted using the one-study-out method, and the results remained robust.

Figure 22. Total effects on quality of life. SMD: standardized mean difference. [36,44,47,49,50,52,55,58-61,65,72,73,75-78,84-86]. SMD: standardized mean difference.

		Exp	erimental			Control		Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD		Difference	SMD	95%-CI	Weight
Alberts et al., 2020	31	6.45	28.8400	32	-3.13	24.3900		1 30	0.35	[-0.14; 0.85]	3.9%
Bennett et al., 2007	28	2.63	8.3646	28	2.49	8.9366			0.02	[-0.51; 0.54]	3.6%
Chan H et al., 2022	20	3.50	3.5000	21	0.75	4.2500			0.69	[0.06; 1.32]	2.7%
Chow et al., 2021	24	2.70	4.6180	17	1.80	3.9871		-	0.20	[-0.42; 0.82]	2.7%
Ferrante et al., 2020	18	9.44	16.9700	17	4.65	24.2100			0.23	[-0.44; 0.89]	2.5%
Frensham et al., 2018	46	16.30	14.7482	45	16.50	13.7400			-0.01	[-0.42; 0.40]	5.1%
Golstejin et al., 2018	249	-0.20	16.5557	229	-1.40	14.5093		-	0.08	[-0.10; 0.26]	11.0%
Howell et al., 2018	52	3.83	10.5000	24	1.71	11.9700			0.19	[-0.29; 0.68]	4.0%
Kim et al., 2019	37	2.90	9.2000	34	1.60	11.3000			0.13	[-0.34; 0.59]	4.3%
Li et al., 2022	47	13.95	10.0300	48	1.74	12.0200			1.09	[0.66; 1.53]	4.8%
Ligible et al., 2012	48	4.30	16.0000	51	-1.50	18.8000		1	0.33	[-0.07; 0.73]	5.3%
Mendoza et al., 2017	29	0.60	19.0789	30	-1.90	21.6838			0.12	[-0.39; 0.63]	3.7%
Millstine et al., 2019	15	-4.40	13.2714	13	-4.20	16.4320		- + i	-0.01	[-0.76; 0.73]	2.0%
Park et al., 2021	75	4.30	26.2532	73	-0.40	20.4546		-	0.20	[-0.12; 0.52]	6.8%
Phillips et al., 2024	25	-0.30	3.0000	24	-0.80	2.9394			0.17	[-0.40; 0.73]	3.2%
Pinto(B) et al., 2013	106	3.73	21.6681	86	-2.74	20.4990		100	0.30	[0.02; 0.59]	7.7%
Pinto(C) et al., 2013	19	3.50	13.5503	24	7.50	14.8584	_	-	-0.27	[-0.88; 0.33]	2.9%
Rastogi et al., 2020	24	4.30	9.5000	23	1.20	5.9000			0.38	[-0.19; 0.96]	3.1%
Uhm et al., 2017	167	5.20	20.5117	172	5.30	22.8026		-	-0.00	[-0.22; 0.21]	9.9%
Vallance et al., 2020	40	3.20	15.7904	40	1.80	10.4748			0.10	[-0.34; 0.54]	4.7%
Walsh et al., 2021	61	7.48	15.0661	62	5.87	18.4008		-i	0.10	[-0.26; 0.45]	6.1%
Random effects model	1161			1093				.	0.19	[0.08; 0.31]	100.0%

Subgroup Analysis (QoL)

In the pooled results based on the usage of multipartnering tools, both the subgroup using multipartnering tools (SMD 0.35, 95% CI 0.05-0.65, P<.001, I^2 =56%) and the subgroup not using them (SMD 0.12, 95% CI 0.03-0.21, P=.02, I^2 =0%) showed significant improvement in QoL (Figure 23). Notably, heterogeneity sharply decreased in the noncombination group (I^2 =0%).

The pooled findings from the subgroup analysis indicated a significant increase in QoL when the duration of WEDS-supported PA programs was no less than 12 weeks,

accompanied by a sharp decrease in heterogeneity (SMD 0.12, 95% CI $0.04-0.21, P<.001, I^2=0\%$; Figure 24).

When grouped by whether the intervention was designed for a specific cancer type, patients' QoL improved in the specific cancer type group (SMD 0.14, 95% CI 0.04-0.23, P=.006, I²=0%; Figure 25), with a sharp decrease in heterogeneity. Whether interventions were designed for a specific cancer type, whether multipartnering tools were used, and the duration of interventions might be potential sources of heterogeneity.

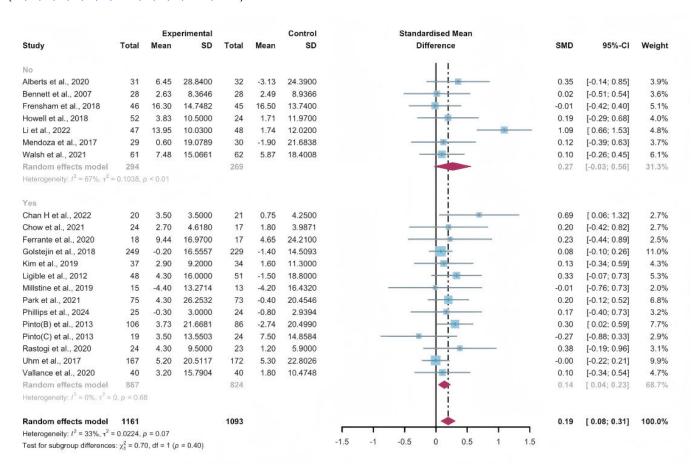
Figure 23. Subgroup analysis on quality of life, grouped by the use of multipartnering tools [36,44,47,49,50,52,55,58-61,65,72,73,75-78,84-86]. SMD: standardized mean difference.

		2000	erimental			Control	Standardised Mean			
Study	Total	Mean	SD	Total	Mean	SD	Difference	SMD	95%-CI	Weigh
No							1.5			
Alberts et al., 2020	31	6.45	28.8400	32	-3.13	24.3900		0.35	[-0.14; 0.85]	3.9%
Bennett et al., 2007	28	2.63	8.3646	28	2.49	8.9366		0.02	[-0.51; 0.54]	3.6%
Chan H et al., 2022	20	3.50	3.5000	21	0.75	4.2500	 	0.69	[0.06; 1.32]	2.79
Frensham et al., 2018	46	16.30	14.7482	45	16.50	13.7400	- · · · · · · · · · · · · · · · · · · ·	-0.01	[-0.42; 0.40]	5.19
Golstejin et al., 2018	249	-0.20	16.5557	229	-1.40	14.5093	-	80.0	[-0.10; 0.26]	11.09
Howell et al., 2018	52	3.83	10.5000	24	1.71	11.9700		0.19	[-0.29; 0.68]	4.09
igible et al., 2012	48	4.30	16.0000	51	-1.50	18.8000	1	0.33	[-0.07; 0.73]	5.3%
Millstine et al., 2019	15	-4.40	13.2714	13	-4.20	16.4320	+ + -	-0.01	[-0.76; 0.73]	2.09
Park et al., 2021	75	4.30	26.2532	73	-0.40	20.4546		0.20	[-0.12; 0.52]	6.89
Pinto(B) et al., 2013	106	3.73	21.6681	86	-2.74	20.4990	100	0.30	[0.02; 0.59]	7.79
Pinto(C) et al., 2013	19	3.50	13.5503	24	7.50	14.8584		-0.27	[-0.88; 0.33]	2.99
Jhm et al., 2017	167	5.20	20.5117	172	5.30	22.8026	- i i	-0.00	[-0.22; 0.21]	9.99
/allance et al., 2020	40	3.20	15.7904	40	1.80	10.4748		0.10	[-0.34; 0.54]	4.79
Walsh et al., 2021	61	7.48	15.0661	62	5.87	18.4008		0.10	[-0.26; 0.45]	6.19
Random effects model	957			900			 ◆	0.12	[0.03; 0.21]	75.79
Heterogeneity: $I^2 = 0\%$, $\tau^2 =$	0, p = 0.6	2								
fes										
Chow et al., 2021	24	2.70	4.6180	17	1.80	3.9871		0.20	[-0.42; 0.82]	2.79
errante et al., 2020	18	9.44	16.9700	17	4.65	24.2100		0.23	[-0.44; 0.89]	2.59
Kim et al., 2019	37	2.90	9.2000	34	1.60	11.3000		0.13	[-0.34; 0.59]	4.39
i et al., 2022	47	13.95	10.0300	48	1.74	12.0200		1.09	[0.66; 1.53]	4.89
Mendoza et al., 2017	29	0.60	19.0789	30	-1.90	21.6838		0.12	[-0.39; 0.63]	3.79
Phillips et al., 2024	25	-0.30	3.0000	24	-0.80	2.9394		0.17	[-0.40; 0.73]	3.29
Rastogi et al., 2020	24	4.30	9.5000	23	1.20	5.9000		0.38	[-0.19; 0.96]	3.19
Random effects model	204			193				0.35	[0.05; 0.65]	24.39
Heterogeneity: $I^2 = 56\%$, τ^2	= 0.0881, /	0.03					į			
Random effects model	1161			1093		30	•	0.19	[0.08; 0.31]	100.0%
Heterogeneity: $I^2 = 33\%$, τ^2	= 0.0224.	0 = 0.07						1		

Figure 24. Subgroup analysis on quality of life, grouped by intervention duration [36,44,47,49,50,52,55,58-61,65,72,73,75-78,84-86]. SMD: standardized mean difference.

Study	Total		erimental SD	Total	Maan	Control SD	Standardised Mean Difference	SMD	95%-CI	Weight
Study	Total	Mean	30	Total	Mean	30	Difference	SINID	95%-01	weight
<12 weeks							1.!			
Alberts et al., 2020	31	6.45	28.8400	32	-3.13	24.3900		0.35	[-0.14; 0.85]	3.9%
Li et al., 2022	47	13.95	10.0300	48	1.74	12.0200	i	1.09	[0.66; 1.53]	4.8%
Mendoza et al., 2017	29	0.60	19.0789	30	-1.90	21.6838		0.12	[-0.39; 0.63]	3.7%
Random effects model	107			110				0.54	[-0.05; 1.12]	12.4%
Heterogeneity: $I^2 = 78\%$, τ^2	= 0.2067,	0.01					l i			
							1 :			
≥12 weeks										
Bennett et al., 2007	28	2.63	8.3646	28	2.49	8.9366	* -	0.02	[-0.51; 0.54]	3.6%
Chan H et al., 2022	20	3.50	3.5000	21	0.75	4.2500	-	0.69	[0.06; 1.32]	2.7%
Chow et al., 2021	24	2.70	4.6180	17	1.80	3.9871		0.20	[-0.42; 0.82]	2.7%
Ferrante et al., 2020	18	9.44	16.9700	17	4.65	24.2100	- -	0.23	[-0.44; 0.89]	2.5%
Frensham et al., 2018	46	16.30	14.7482	45	16.50	13.7400	- † i	-0.01	[-0.42; 0.40]	5.1%
Golstejin et al., 2018	249	-0.20	16.5557	229	-1.40	14.5093	-	0.08	[-0.10; 0.26]	11.0%
Howell et al., 2018	52	3.83	10.5000	24	1.71	11.9700	- • -	0.19	[-0.29; 0.68]	4.0%
Kim et al., 2019	37	2.90	9.2000	34	1.60	11.3000		0.13	[-0.34; 0.59]	4.3%
Ligible et al., 2012	48	4.30	16.0000	51	-1.50	18.8000	 	0.33	[-0.07; 0.73]	5.3%
Millstine et al., 2019	15	-4.40	13.2714	13	-4.20	16.4320		-0.01	[-0.76; 0.73]	2.0%
Park et al., 2021	75	4.30	26.2532	73	-0.40	20.4546		0.20	[-0.12; 0.52]	6.8%
Phillips et al., 2024	25	-0.30	3.0000	24	-0.80	2.9394	- 4	0.17	[-0.40; 0.73]	3.2%
Pinto(B) et al., 2013	106	3.73	21.6681	86	-2.74	20.4990	i	0.30	[0.02; 0.59]	7.7%
Pinto(C) et al., 2013	19	3.50	13.5503	24	7.50	14.8584	* i i	-0.27	[-0.88; 0.33]	2.9%
Rastogi et al., 2020	24	4.30	9.5000	23	1.20	5.9000		0.38	[-0.19; 0.96]	3.1%
Uhm et al., 2017	167	5.20	20.5117	172	5.30	22.8026		-0.00	[-0.22; 0.21]	9.9%
Vallance et al., 2020	40	3.20	15.7904	40	1.80	10.4748		0.10	[-0.34; 0.54]	4.7%
Walsh et al., 2021	61	7.48	15.0661	62	5.87	18.4008		0.10	[-0.26; 0.45]	6.1%
Random effects model	1054			983			•	0.12	[0.04; 0.21]	87.6%
Heterogeneity: $I^2 = 0\%$, $\tau^2 =$	0, p = 0.8	6					1 !			
	440-			4005			l i			400.00/
Random effects model				1093				0.19	[0.08; 0.31]	100.0%
Heterogeneity: $I^2 = 33\%$, τ^2			0.47				-1.5 -1 -0.5 0 0.5 1 1.5			
Test for subgroup difference	98: $\chi_1^2 = 1.8$	ь, df = 1 (<i>j</i>	p = 0.1/)							

Figure 25. Subgroup analysis on quality of life, grouped by whether the intervention was designed for a specific cancer type [36,44,47,49,50,52,55,58-61,65,72,73,75-78,84-86]. SMD: standardized mean difference.



Discussion

Principal Findings

A total of 46 studies met the eligibility criteria for this meta-analysis. Compared with usual care or waitlists, WEDS-supported PA programs significantly improved cancer survivors' objectively measured MVPA, subjectively measured PA, steps per day, and QoL, but showed no significant effect on reducing sedentary behavior or improving BMI. The SMD of total effects ranged from -0.07 to 0.66, which is consistent with previous studies [23,29], thereby confirming the effectiveness of WEDS-supported PA programs.

Quality of Evidence and Methodology

Overall, the evidence quality and methodology were rated as moderate; 24 studies used the intention-to-treat analysis, while 22 utilized per-protocol analysis. Using the revised Cochrane Risk-of-Bias tool, it was determined that 12 of 46 (26%) studies were assessed as having a low risk of bias, 33 of 46 (72%) studies raised some concerns, and only 1 of 46 (2%) studies was deemed to have a high risk of bias. Specifically, of these 46 studies, 33 (72%) were flagged for potential bias related to the randomization process, and 1 (2%) raised concerns regarding bias in the measurement of outcomes. In detail, all studies were assessed as having either a low or unclear risk associated with the

randomization process, primarily because some investigators failed to provide comprehensive details on the randomization techniques or adequately describe the allocation concealment methods. Consequently, the overall methodological quality was deemed moderate. These findings underscore the need for additional randomized controlled trials in the future, with a focus on more transparent reporting to enhance the robustness of research findings.

PA-Related Outcomes

The results of this meta-analysis demonstrated that WEDS-supported PA programs significantly improved objectively measured MVPA, subjectively reported PA, steps per day, and QoL in cancer survivors, but had no significant effect on sedentary behavior or BMI. The findings related to objectively measured MVPA, subjectively reported PA, and steps per day are consistent with those of previous studies [23].

For cancer survivors, these interventions serve as tracking devices (continuously collecting current activity), feedback tools (providing immediate information on activity levels), and environmental cues (reminders to be active). Through these continuous influences, patients' levels of PA increase [91]. In addition, the partnering tools used in these interventions enable cancer survivors to record, report, and contact HCPs anytime and anywhere they need, ensuring timely revision of exercise prescriptions and providing knowledge related to their disease and symptoms [78,80]. Moreover,

the similar results between subjectively reported PA and objectively measured MVPA highlight the feasibility of WEDS-supported PA programs in improving PA among cancer survivors. On the contrary, we found that sedentary behavior did not improve significantly. This may be because patients may choose to ignore activity alarms while remaining sedentary [46]. More cognitive behavioral therapy is needed to enhance patients' awareness and motivation to reduce sedentary behavior [92]. Regarding BMI, the lack of a significant difference might be due to patients gaining muscle mass, which can offset weight loss, resulting in no apparent change in BMI despite positive effects on fitness [93]. Furthermore, diet management has been shown to play a more significant role in weight loss programs than PA alone in many studies [94,95].

The results of the subgroup analysis on the usage of multipartnering tools revealed that objectively measured MVPA significantly improved regardless of whether multipartnering tools were used. This outcome may be because, at the beginning of the interventions, researchers set PA goals for participants and adjusted those goals based on their performance through the partnering tools. Whether or not multipartnering tools were used, the partnering tools could still serve as reminders for patients to complete more MVPA. Thus, the use of multipartnering tools may not have influenced objectively measured MVPA. By contrast, for subjectively reported PA, a significant difference was observed in the nonusage of multipartnering tools. When patients are assisted by multipartnering tools to remind them to improve PA, they may lack initiative, and their self-efficacy regarding PA may decrease, along with their perception of subjectively reported PA [96]. For steps per day, the multipartnering tools groups showed significant differences. This may be because various partnering tools play a more comprehensive role, such as providing real-time conversations through telephone calls and offering relevant knowledge via apps or websites. Through this type of multimedia stimulation, participants can receive more comprehensive reminders and encouragement to remain physically active.

Moreover, we observed significant improvements in objectively measured MVPA, subjectively reported PA, steps per day, and QoL among participants who received long-term (no less than 12 weeks) interventions, which is consistent with previous similar findings [36]. Research has shown that longer-term interventions are conducive to forming healthier lifestyles and developing lasting habits. Therefore, the patients can derive enjoyment from PA and are more willing to complete additional PA programs [97,98]. Similar to the overall effects, the results showed no significant improvements in sedentary behavior or BMI in the subgroups categorized by duration.

When studies were grouped by whether the intervention was designed for a specific cancer type, the results showed that both steps per day and QoL significantly improved in patients who received interventions tailored to their specific cancer type. This may be because tailored programs could better meet patients' specific needs, such as providing information related to their cancer type and offering

PA programs suited to their condition. By contrast, regardless of whether the interventions were tailored or not, the effectiveness of WEDS-supported PA programs in improving objectively measured MVPA, subjectively reported PA, sedentary behavior, and BMI did not change.

In brief, WEDS is promising for improving MVPA, subjectively reported PA, steps per day, and QoL. Long-term interventions (≥12 weeks) are effective in improving PA-related outcomes, except for sedentary behavior, and the use of multipartnering tools should depend on the patients' preferences and habits. Attention should also be given to the proper use of multipartnering tools, the optimal duration, and whether the intervention is tailored to specific cancer types when developing new WEDS-supported interventions. Further studies are needed to explore the most effective intervention characteristics for improving patients' sedentary behavior and BMI. This approach will facilitate the development of more effective WEDS-supported interventions.

Quality of Life

This meta-analysis demonstrated that, compared with usual care or waitlists, WEDS-supported PA programs have a significant effect on the QoL of cancer survivors, which is consistent with the findings of previous studies [23]. The QoL assessed in the included studies was health-related QoL, which encompasses not only basic physical functioning but also patient participation in activities such as work and entertainment [99]. WEDS-supported PA programs significantly improved cancer survivors' inactive lifestyles, enhanced their self-efficacy and feelings of self-worth, increased their satisfaction with life, and indirectly influenced their QoL [100]. Moreover, appropriate social relationships, cancer and self-care education, and psychological support provided through partnering tools could further help improve cancer survivors' QoL [101,102].

In the subgroup analysis, regarding the use of multipartnering tools, QoL was significantly improved in both the usage and nonusage groups. This may be because interventions in both groups provided reminders to patients, which significantly enhanced participants' PA levels and indirectly reduced their symptom burden, thereby improving QoL. When grouped by intervention duration, QoL was significantly improved in the long-term subgroups. Longer intervention durations enable patients to develop sustained habits of positive PA. Additionally, patients may have more opportunities to access diverse forms of support over the long term, which provides greater encouragement for engaging in PA and fosters the adoption of self-management strategies, thereby improving their QoL [103]. Furthermore, we observed a significant improvement in QoL among patients who received interventions designed for a specific cancer type. Researchers could tailor specific programs, such as PA regimens and psychological support from HCPs, according to the characteristics of each patient's cancer.

In essence, WEDS-supported PA programs enable cancer survivors to engage positively with WEDs and partnering tools, with the potential to reduce negative affective states and consequently enhance their QoL. Researchers should carefully consider the duration of intervention when designing WEDS-supported strategies. Additionally, further investigation is warranted to evaluate the effectiveness of partnering tools in addressing the specific needs of cancer survivors.

Limitations

This study has several limitations. First, heterogeneity existed due to variations in the format of partnering tools, durations of intervention, and types of cancer. Second, the reporting of study results may have been influenced by commercial interests associated with PA improvements, posing a potential risk of publication bias. Additionally, a significant proportion of the research was conducted in Western countries, and responses to WEDS-supported PA programs may vary among participants from different regions [104]. Finally, despite conducting an exhaustive literature search, publication bias could not be completely eliminated. Therefore, the outcomes of this meta-analysis should be interpreted with caution, and more high-quality randomized controlled trials are needed in the future.

Implications

In this study, we quantitatively integrated existing findings and found that WEDS-supported PA programs were effective in improving PA levels (both objectively and subjectively), daily steps, and QoL. The mechanisms through which WEDS-supported PA programs bring clinical benefits may include providing persistent reminders to encourage PA, offering convenient access to consultations with HCPs, collecting health-related data, recording electronic health records, and facilitating social groups for patients to communicate with others facing similar conditions [47,50]. Thus, HCPs can use WEDS as a supplementary tool to monitor patients' physiological data, manage care, adjust exercise prescriptions, and provide timely feedback and disease-related information.

With increasing research focusing on WEDs and other forms of eHealth as interventions to promote PA among cancer survivors, WEDS has the potential to become a valuable tool for HCPs and a novel reminder and management resource for cancer survivors. It can automatically sync data, thereby reducing the self-monitoring burden associated with traditional web-based interventions [105]. Additionally, previous studies often failed to adequately consider the role of partnering tools, resulting in their underutilization and a missed opportunity to maximize the benefits for patients' PA engagement. Furthermore, we observed that certain aspects of the intervention, such as the use of multipartnering tools, the duration of the intervention, and whether the intervention was tailored for specific cancer types, influenced its overall efficacy. This underscores the need for further standardization and more rigorous quantitative studies to refine the WEDS-supported intervention framework and to fully explore the potential benefits of WEDS-supported PA programs. Moreover, efforts should be made to enable data intercommunication between different commercial WEDs, thereby improving the feasibility and accessibility of these interventions

Conclusions

WEDS-supported PA programs offer a convenient and affordable method for assisting cancer survivors by serving as reminders and records of their PA. This meta-analysis of randomized controlled trials revealed that WEDS-supported PA programs significantly improved cancer survivors' level of PA (both objectively and subjectively), steps per day, and QoL, but had no significant effect on reducing sedentary behavior or BMI. These results varied based on the use of multipartnering tools, intervention duration, and patients' cancer type. Further standardization and promotion of WEDS-supported PA programs are warranted in the future.

Acknowledgments

This work was supported by the National Natural Science Foundation of China (grant 82172842); the China Medical Board (grant 22-482); the Ministry of Education University-Industry Collaborative Education Program (grant 230720523707281); the Sichuan University Graduate Students Education and Teaching Reform Research Program (grants GSSCU2023090 and GSSCU2023095); and the Chengdu Eastern New Area Municipal Administration Committee Program (grants 200304 and 00402053A29YN).

Data Availability

The datasets generated or analyzed during this study are available from the corresponding author (YS) on reasonable request.

Authors' Contributions

Conceptualization: YS, ZW, YL

Data curation: ZW, YL
Formal analysis: ZW, YL
Funding acquisition: YS
Investigation: ZW, YL

Methodology: YS, ZW, YL, QW Project administration: YS

Resources: YS Supervision: YS, QW Validation: ZW, YL Visualization: ZW, YL Writing—original draft: ZW

Writing-review & editing: YS, ZW, YL, QW

Conflicts of Interest

None declared.

Multimedia Appendix 1

Search strategies.

[DOCX File (Microsoft Word File), 17 KB-Multimedia Appendix 1]

Multimedia Appendix 2

Characteristics of included studies.

[DOCX File (Microsoft Word File), 39 KB-Multimedia Appendix 2]

Multimedia Appendix 3

Summary of all outcomes included in the meta-analysis.

[DOCX File (Microsoft Word File), 33 KB-Multimedia Appendix 3]

Multimedia Appendix 4

Results of the sensitivity analysis.

[DOCX File (Microsoft Word File), 851 KB-Multimedia Appendix 4]

Checklist 1

PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) checklist. [PDF File (Adobe File), 176 KB-Checklist 1]

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Abbreviations

HCP: health care professional **MeSH:** Medical Subject Headings

mHealth: mobile health

MVPA: moderate-to-vigorous-intensity physical activity

PA: physical activity

PICOS: Participants, Interventions, Comparisons, Outcomes, and Study Design **PRISMA:** Preferred Reporting Items for Systematic Reviews and Meta-Analyses

QoL: quality of life

SMD: standard mean difference **WED:** wearable electronic device

WEDS: wearable electronic device system

Edited by Javad Sarvestan; peer-reviewed by Antonio Martinko, Frans Van der Ouderaa; submitted 23.03.2025; final revised version received 09.06.2025; accepted 10.06.2025; published 14.08.2025

Please cite as:

Wang Z, Li Y, Wang Q, Su Y

The Effectiveness of Wearable Electronic Device System–Supported Physical Activity Programs for Cancer Survivors: Meta-Analysis of Randomized Controlled Trials

J Med Internet Res 2025;27:e74347 URL: <u>https://www.jmir.org/2025/1/e74347</u>

doi: <u>10.2196/74347</u>

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