# Technology-Assisted Weight Loss Interventions in Primary Care: A Systematic Review

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**BACKGROUND:** The US Preventive Services Task Force recommends screening for and treating obesity. However, there are many barriers to successfully treating obesity in primary care (PC). Technology-assisted weight loss interventions offer novel ways of improving treatment, but trials are overwhelmingly conducted outside of PC and may not translate well into this setting. We conducted a systematic review of technology-assisted weight loss interventions specifically tested in PC settings.

**METHODS:** We searched the literature from January 2000 to March 2014. Inclusion criteria: (1) Randomized controlled trial; (2) trials that utilized the Internet, personal computer, and/or mobile device; and (3) occurred in an ambulatory PC setting. We applied the Cochrane Effective Practice and Organization of Care (EPOC) and Delphi criteria to assess bias and the Pragmatic-Explanatory Continuum Indicator Summary (PRECIS) criteria to assess pragmatism (whether trials occurred in the real world versus under ideal circumstances). Given heterogeneity, results were not pooled quantitatively.

**RESULTS:** Sixteen trials met inclusion criteria. Twelve (75 %) interventions achieved weight loss (range: 0.08 kg -5.4 kg) compared to controls, while 5-45 % of patients lost at least 5 % of baseline weight. Trial duration and attrition ranged from 3-36 months and 6-80 %, respectively. Ten (63 %) studies reported results after at least 1 year of follow-up. Interventions used various forms of personnel, technology modalities, and behavior change elements; trials most frequently utilized medical doctors (MDs) (44 %), webbased applications (63 %), and self-monitoring (81 %), respectively. Interventions that included clinician-guiding software or feedback from personnel appeared to promote more weight loss than fully automated interventions. Only two (13%) studies used publically available technologies. Many studies had fair pragmatism scores (mean: 2.8/4), despite occurring in primary care.

**DISCUSSION:** Compared to usual care, technologyassisted interventions in the PC setting help patients achieve

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Received December 9, 2013 Revised June 2, 2014 Accepted July 23, 2014 Published online August 19, 2014 weight loss, offering evidence-based options to PC providers. However, best practices remain undetermined. Despite occurring in PC, studies often fall short in utilizing pragmatic methodology and rarely provide publically available technology. Longitudinal, pragmatic, interdisciplinary, and opensource interventions are needed.

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## INTRODUCTION

The obesity epidemic accounts for unprecedented rates of chronic disease<sup>1–3</sup> and influences numerous interactions in primary care (PC).<sup>4</sup> The prevalence of overweight and obese adults, while having plateaued over the past decade, remains high, such that among US adults today, 34.9 % are obese,<sup>5</sup> up from 10.4 % in 1960 and 19.9 % in 1994.<sup>6</sup>

Unfortunately, there is a paucity of available weight loss interventions suitable to the real-world PC setting, with most research and guideline formulation conducted inside academic silos,<sup>7–10</sup> not in practice settings such as PC offices.<sup>11</sup> With 577 million yearly PC visits, effective weight loss interventions within PC could potentially have a large health impact.<sup>12</sup>

Weight loss, even when modest ( $\geq$  5 %), is associated with significant improvement in cardiovascular disease (CVD) risk factors,<sup>13</sup> quality of life,<sup>14</sup> and all-cause mortality for those with comorbidities.<sup>15–18</sup> Intensive, tailored lifestyle modification programs result in significant weight loss, health benefits,<sup>19–21</sup> and perhaps cost savings.<sup>22</sup> Those who receive advice from their PC provider (PCP) to lose weight are more likely to do so,<sup>23</sup> yet existing programs are challenged by high attrition rates, weight regain, resource-intensive requirements, and poor scalability.<sup>24–27</sup> Concurrent provider barriers include limited visit time,<sup>28</sup> inadequate reimbursement,<sup>29</sup> lack of training,<sup>30</sup> and poor competency.<sup>31</sup> As a result, PCPs struggle to properly counsel their obese patients,<sup>32–34</sup> even as patients are genuinely interested in receiving counseling from them.<sup>35,36</sup>

Given these large hurdles, there is a strong need to identify novel weight loss interventions that are applicable to the PC office. Technology-assisted interventions have the potential to address barriers to providing care in the PC office through time and cost savings, improved feedback, enhanced selfmonitoring, and convenience of use.<sup>37</sup> Many studies<sup>38,39</sup> and multiple reviews<sup>40–42</sup> have assessed technology-assisted weight loss interventions, yet few have placed emphasis on interventions suitable to the real-world PC practice. Many interventions have yet to be integrated into PC, while others are publically unavailable, poorly studied, or expensive.<sup>43</sup> Other trials are not pragmatic,<sup>44</sup> where pragmatism refers to the extent to which studies operate in real-world circumstances.<sup>45</sup>

This systematic review examines technology-assisted weight loss interventions specifically provided in PC settings, and aims to highlight their innovation, impact, and pragmatism. To our knowledge, this is the first such systematic review to examine this topic from the lens of the pragmatic PC practice.

#### **METHODS**

Our review follows standard PRISMA methodology,<sup>46</sup> and the detailed protocol is available on PROPSERO (CRD42013003998)<sup>47</sup> and in Appendix 1, available online. We briefly describe our methods below.

## **Eligibility Criteria**

Our aim was to identify and synthesize PC-based weight loss intervention studies that incorporated various technologies. We included peer-reviewed, randomized controlled trials (RCTs) that used the Internet, a personal or in-office computer, and/or mobile device, had weight loss as a primary outcome, and took place in an ambulatory PC (internal medicine, family medicine, OBGYN) setting. We excluded studies that utilized pedometers, postal mail, or telephone calls as the sole technology, non-RCTs, studies conducted in a specialty or nonambulatory setting, and studies involving pediatric populations. We chose to exclude pedometers, as they have previously been reviewed and demonstrated modest weight loss.<sup>48,49</sup> We only considered trials conducted after the year 2000, as few PC-based studies were published before that date and any prior technologyassisted interventions would likely be obsolete.<sup>50</sup>

# Data Sources and Search Strategy

In late 2012, we performed initial scoping searches with Google Scholar. We then finalized our search strategies with the help of a research librarian and searched the following databases without language restrictions to identify citations and trials from 1 January 2000 to 11 March 2014: PubMed/ MEDLINE, EMBASE, Cochrane Database of Systematic Reviews, and Cochrane CENTRAL. To limit to RCTs, we used validated search filters.<sup>51</sup> We also searched the reference lists of identified studies and reviews.

#### Study Selection and Data Extraction

To determine the selection of studies, two authors (DML and SS) independently investigated the titles of all 1.201 initial citations. We excluded 934 articles by title and 166 articles by abstract, leaving 101 for full-text review (Fig. 1). Any disagreements were resolved through consensus, and when needed, a third reviewer (MJ). Two reviewers (DML and SS) independently extracted data for the following variables: baseline demographics, recruitment procedures, setting, intensity,<sup>52</sup> mode of customization,<sup>53</sup> motivating theory, technology modality, personnel, attrition, weight loss (kg, percent, and percent achieving  $\geq$  5 % loss), bias, and pragmatism. Because recruitment (all PC offices) and customization (all tailored) did not vary, we do not discuss them further. We similarly do not discuss motivating theory, as studies infrequently reported this.

## **Bias Assessment**

The Delphi<sup>54</sup> and Cochrane Effective Practice and Organization of Care (EPOC)<sup>55</sup> criteria were used to assess bias. Delphi and EPOC contain assessment items that pertain to interventions occurring in real-world PC settings, and are commonly used tools for this type of review.<sup>56</sup> Two reviewers (DML and SS) independently evaluated the trials (Supplemental Table 1).

#### Pragmatism Assessment

We evaluated the degree of pragmatism of each intervention with the Pragmatic-Explanatory Continuum Indicator Summary (PRECIS).<sup>45</sup> This tool helps researchers evaluate the likelihood that a given intervention will work in a real-world setting versus a clinical trial in an ideal setting. An intervention is considered more pragmatic if it is conducted in real-world circumstances; it is considered more explanatory if it occurs in ideal circumstances. PRECIS was initially created by the CONSORT Work Group to assist in trial design and has been adapted to evaluate studies in systematic reviews.<sup>57</sup> We modeled our use of PRECIS based on a recent adaptation evaluating the Practice-Based Opportunities for Weight Reduction (POWER) trials (two of which are used in this review), using a 0 (explanatory) to 4 (pragmatic) scale in ten

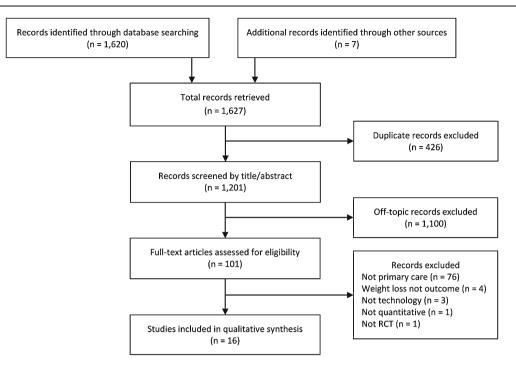


Figure 1. PRISMA flow diagram.

domains.<sup>44</sup> A detailed tabular explanation of each of these domains can be found in Fig. 4 and elsewhere.<sup>45</sup>

#### **Statistical Analysis**

Study heterogeneity did not allow us to conduct a metaanalysis. We therefore aggregated our results with ranges and narrative summation in most instances.

#### RESULTS

Sixteen trials<sup>58–73</sup> met inclusion criteria and are summarized in Table 1. For readers interested in additional tabular detail and narrative summation of each trial, see Supplemental Table 2 and Appendix 2, available online. Participants were more often female (62 %), white (71 %), and middle-aged. Fifteen (94 %)<sup>58–66,68–73</sup> trials were of high intensity and employed tailored interventions. Interventions ranged from 3 – 36 months in duration, with the majority of studies (10/16, 63 %)<sup>58,60–64,68,70,71,73</sup> reporting results after at least 1 year of follow-up. Participant attrition rates ranged from 6 to 80 %.

Compared to the control group, most  $(12/16, 75 \%)^{58-63,66-70,73}$  technology-assisted interventions achieved weight loss at the end of the study period. Weight loss in active treatment arms ranged from 0.08 kg to 5.4 kg (0.8 % – 5.8 % of initial body weight). The percentage of patients losing at least 5 % of baseline weight ranged from 5 % to 45 %.

## Personnel

Figure 2 illustrates the association between weight loss and mix of personnel delivering the intervention. Seven  $(44 \%)^{58,60,63,66,70,71,73}$  trials used more than one type of personnel. Trials most frequently utilized medical doctors (MDs)  $(7/16, 44 \%)^{58,60-62,66,70,71}$ . When MDs participated, they were solely in a supportive role (no active counseling) in 3/7 (43 %)<sup>58,60,71</sup> studies, whereas they actively counseled patients in 3/7 (43 %)<sup>61,62,66</sup> studies. In 1/7 (14 %)<sup>70</sup> studies, they co-taught group lifestyle sessions with registered dieticians (RDs) and psychologists. Four studies utilized RDs who most often counseled participants one-on-one (3/4, 75 %),<sup>59,63,73</sup> using a combination of in-person, telephone, and email-based exchanges. In one study, RDs co-taught group sessions with MDs and psychologists.<sup>70</sup> Several studies used "other" personnel (5/16, 31 %).58,60,63,68,70 These ranged from "fitness instructors,"63 to "weight loss coaches,"58 to "community health educators."60 These personnel had varied levels of training and were most often involved in lifestyle coaching delivered via various modalities, including in-person, telephone, and email-based exchanges similar to the RD studies.

Only one study included nurse practitioners (NPs), where their counseling was guided by software.<sup>71</sup> Although this did not demonstrate significant weight loss compared with usual PC by MDs at the study's 3 year conclusion, significant weight loss was demonstrated at 1 year.<sup>74</sup>

Three  $(19 \%)^{64,69,72}$  studies utilized no personnel, instead relying entirely on automated advice based on stage of change<sup>72</sup> or based on dietary and physical activity

	Parti	cipants	5			Intervention		Outcomes		
Study	n	Age $(\overline{x})$	% F	Ethnicity (%)	<b>BMI</b> $(\overline{x})$	Intervention (n)	Duration (mos)	Attrition (%)	Weight $\Delta$ (kg)	≥ 5 % weight loss (%)
Appel <sup>58</sup>	415	54	64	56W, 41B	37	<ul> <li>All: PCP visit at 6/12/24 mos for encouragement</li> <li>IG1 (138): in-person support, website: self-monitoring, auto feedback, &amp; email reminder to login</li> <li>IG2 (139): remote support via telephone, same website &amp; emails as IG1</li> <li>CG (138): 1 weight loss coach meeting, brochures, weight loss websites</li> </ul>	24	16	IG1: -5.1* IG2: -4.6* CG: -0.8	IG1: 41* IG2: 38* CG: 19
Bennett (2010) <sup>59</sup>	101	54	47	50W, 31B, 5H	35	IG (51): self-monitoring website, 4 RD visits via phone and in- person CG (50): usual care & basic	3	16	IG: -2.3* CG: +0.3	IG: 26 <sup>NR</sup> CG: 0
Bennett (2012) <sup>60</sup>	365	55	69	71B, 13H	37	materials IG (180): web-based tailored behavior change goals, skills training via website or interac- tive voice response, telephone counseling w trained commu- nity health educator, primary care provider endorsement, 12 optional in-person group sup- port sessions, walking kit w pedometer CG (185): usual care & self-help booklet	24	14	IG: -1.5* CG: -0.5	IG: 20 CG: 20
Christian (2008) <sup>61</sup>	310	53	66	100H	35	IG (155): goal-setting computer program, then regular MD clinic visits w 3/6/9 mos goals reassessment	12	12	IG: -0.08 CG: +0.6	IG: 21* CG: 11
Christian (2011) <sup>62</sup>	279	50	68	51W, 44H	34	CG (155): usual care & info packet IG (140): goal-setting computer program, then MD clinic visits to reinforce goals, 6 mos goals reassessment w computer CG (139): usual care & info packet	12	6	IG: -1.5* CG: +0.1	IG: 26* CG: 8
Ma <sup>63</sup> / Xiao† <sup>79</sup>	171	53	46	79W, 4H	32	All: Heart360 website, standardized monthly emails, ability to submit questions online IG1 (79): 3-mo intensive in- person weekly DPP w physical activity & food tastings, per- sonalized monthly emails on Heart360 progress IG2 (81): 3-mo intensive at- home DPP DVD, standardized weekly emails	24	29	IG1: -5.4* IG2: -4.5* CG: -2.4	IG1: 45* IG2: 30 CG: 17
McConnon <sup>64</sup>	221	46	77	95W	34	CG (81): usual care IG (111): website: tailored advice, tools & information to support behavior change in terms of dictary & physical activity pat- terns, reminder emails CG (110): usual care, small info	12	41	IG: -1.3 CG: -1.9	IG: 22 <sup>NR</sup> CG: 18
McDoniel <sup>65</sup>	111	46	61	78W, 20H	37	booklet All: MI counseling at wk 4 & wk 8; core topic email newsletters weekly IG (55): MedGem Analyzer for nutrition program & Balance- Log for SM CG (56): Usual care: 3-day food	3	28	IG: -3.5 CG: -3.7	IG: 31 CG: 42
Mehring <sup>66</sup>	186	48	69	NR	34	menu, paper journal) IG (109): HausMed eHealth web- based coaching program w MD input and 3 phone calls from MD or MA at wks 1,5,12 for motivation CG (77): usual care	3	20	IG: -2.9* CG: -1.6	IG: 26 CG: 16

### Table 1. Technology-Assisted Weight Loss Interventions in Primary Care

	Participants					Intervention	Outcomes			
Study	n	Age $(\overline{x})$	% F	Ethnicity (%)	<b>BMI</b> $(\overline{x})$	Intervention (n)	Duration (mos)	Attrition (%)	Weight $\Delta$ (kg)	≥ 5 % weight loss (%)
Nanchahal (2009) <sup>67</sup>	123	47	80	96W	36	IG (61): ProHealthClinical structured lifestyle support w tailored diet, self-monitoring w diary, coping skills, & RN feedback	3	15	IG: -4.0* CG: +1.2	IG: 17 <sup>NR</sup> CG: 10
Nanchahal (2012) <sup>68</sup>	381	49	72	73W	33	CG (62): usual care IG (191): structured one-on-one in-person sessions (6 in 1st 12 wks, 4 in 2nd 12 wks, 3 in 3rd 12 wks), perfect-diet- tracker.com for SM; adamsportionpot.com, pedometers CG (190): usual care & asked to	12	43	IG: -2.4 CG: -1.3	IG: 34* CG: 19
Rothert <sup>+69</sup>	2862	45	83	56W, 36B	32	seek weight loss from PCP All: 20min computer assessment IG (1475): Web-tailored weight management program x 6 wks w 1/3/6 wk email asking par- ticipants to enter weight CG (1387): Web-info-only ma- terials on Kaiser's website	6	80	IG: -2.8* CG: -1.1	NR
Spring <sup>70</sup>	70	58	14	75W, 25B, 6H	36	All: 2 wk run-in baseline IG (35): Weight Loss Phase (mos 0–6): twice weekly MOVE sessions, PDA for self- monitoring w automated feed- back, coach-derived feedback; Maintenance Phase (mos 7– 12): monthly MOVE sessions, telephone coach conversation if no data transmitted CG (34): All MOVE sessions as	12	23	IG: -2.9* CG: +0.02	IG: 30* CG: 15
ter Bogt <sup>71</sup>	457	56	52	NR	30	IG, but no PDA, no coach calls All: baseline online or paper questionnaire IG (225): 4 visits w NP using software, 1 telephone f/u in yr 1, then 1 visit & 2 telephone f/u's in yr 2 & yr 3 CG (232): MD usual care	36	22	IG: -1.1 CG: -0.5	IG: 5 CG: 5
Verheijden <sup>72</sup>	146	63	45	NR	29	IG (73): Heartweb online pro- gram: monthly stage-of- change questionnaire w subse- quent tailored nutrition sug- gestions, bulletin board, dietary fat tracker, low fat recipes	8	9	NR	NR
Wylie-Rosett <sup>73</sup>	588	52	82	83W	36	CG (73): usual care All: computer assessment IG1 (236): 6 group workshops, RD/MSW consult (telephone or face-to-face) up to 18times + IG2 + CG IG2 (236): kiosks weekly (20–30 mins) x 3 mos, then monthly + CG CG (116): workbook	12	21	IG1: -3.4* IG2: -2.1* CG: -1.0	IG1: 31* IG2: 23* CG: 15

B Black; BMI Body Mass Index ( $kg/m^2$ ); CG Control Group;  $\Delta$  Change; DPP Diabetes Prevention Program; H Hispanic; IG Intervention Group; NR Not Reported; PCP Primary Care Provider; PDA Personal Digital Assistant; W White; TMean

Technology Modality: App Mobile App; CS Clinician Software; HC Home Computer (no internet); Int Internet; K Kiosk; EP Exercise Physiologist; MSW Master of Social Work; MD Medical Doctor; None No Personnel; NP Nurse Practitioner; O Other Personnel; RD Registered Dietician; RN Registered Nurse

Elements: AF Automated Feedback; Con Contests; IPF In-Person Feedback; LC Lifestyle Coaching; MDF MD Feedback; O Other; P2P Peer-to-Peer Support; Rem Reminders; SM Self Monitoring

\* p < 0.05

<sup>†</sup> Xiao et al. re-consented the patients from Ma et al. and followed them for nine additional months; we report their data with this extension period. Also, their data is for  $\geq 7\%$  weight loss, not  $\geq 5\%$ . ‡ Completers-only analysis (i.e., not intention to treat)

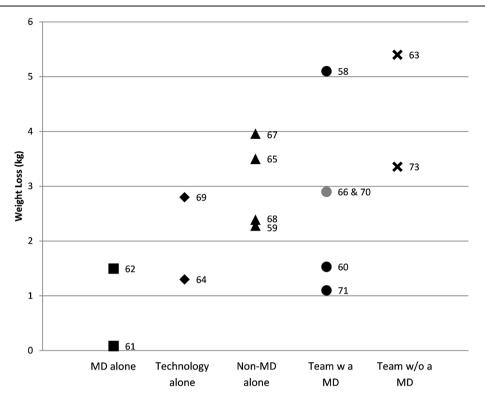


Figure 2. Weight loss by personnel type. *Square*: MD alone (e.g., no additional personnel assisted MD). *Diamond*: Technology alone (e.g., no personnel, intervention automated). *Triangle*: Non-MD alone (e.g., RN, weight loss coach, etc.). *Circle*: Team with an MD. X: Team without an MD. *Gray*: Two overlapping data points. Numbers beside each shape indicate citation.

patterns.<sup>64,69</sup> All employed web-based self-monitoring. Importantly, 2/3 (67 %)<sup>64,72</sup> of these studies did not demonstrate significant weight loss, while 1/3 (33 %)<sup>69</sup> showed 2.8 kg weight loss, but had 80 % attrition, did not use intention to treat analysis, and used self-reported data.

## **Technology Modality**

Technology modalities included web-based applications (63 %),  ${}^{58-60,63-66,68,69,72}$  clinician-guiding software (44 %),  ${}^{58,60-62,66,67,71}$  kiosks (19 %),  ${}^{61,62,73}$  home PCs (13 %),  ${}^{65,68}$  mobile applications (6 %),  ${}^{70}$  and short message services (SMS, "texting") (6 %).  ${}^{66}$  Half of the studies  ${}^{59,64,67,69-73}$  employed a single technology modality (weight loss: 1.1 – 3.96 kg), while others  ${}^{58,60-63,65,66,68}$  used multiple modalities (weight loss: 0.082 – 5.4 kg). Two out of 16 (13 %)  ${}^{63,73}$  used publically available open-source technology.

Ten  $(63 \%)^{58-60,63-66,68,69,72}$  trials employed web-based applications. All utilized proprietary technology for self-monitoring with the exception of Heart360.org.<sup>63</sup> Spring et al.<sup>70</sup> used a proprietary mobile app, while Mehring et al. employed texting through a commercial website.<sup>66</sup>

Clinician-guiding software, used in  $7/16 (44 \%)^{58,60-62,66,67,71}$  studies, assisted various practitioners (MDs, RNs, and NPs) in

guiding participants toward achieving their weight loss goals. In Christian et al.,<sup>61,62</sup> for example, participants took a kiosk-based survey that provided the MD with a one-page assessment of the patient and assisted in goal-setting and goal-resetting over 9 months. Participants in all seven studies had significant weight loss at 3 months,<sup>66,67</sup> 1 year,<sup>61,62,74</sup> and 2 years,<sup>58,60</sup> although in ter Bogt et al., weight loss in the intervention group was no longer significantly different from that in the control group at 3 years.<sup>71</sup>

Three  $(19 \%)^{61,62,73}$  studies used kiosks to obtain baseline health and behavior-related patient information<sup>61,62</sup> or to facilitate a computer-based weight loss program while in the waiting room.<sup>73</sup> In Wylie-Rosett et al.,<sup>73</sup> only about 50 % of participants enjoyed working on the kiosks, 75 % found them a poor substitute for human contact, yet 75 % found the kiosk easy to use. In contrast, about 75 % and 67 % of participants in Nanchahal (2009) et al.<sup>67</sup> found automated and RN web-based feedback and motivation very/extremely helpful, respectively.

## **Behavior Change Elements**

All studies used evidence-based elements to promote behavior change (Fig. 3). The most common was self-monitoring (13/16, 81 %),  $^{58-60,63-72}$  where participants recorded daily dietary and physical activity behaviors, usually on web-based software. This technology-supported self-monitoring occurred in 11/13 (85 %)

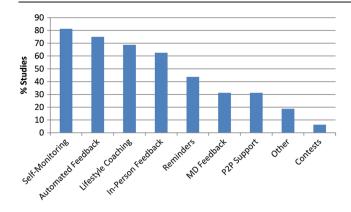


Figure 3. Behavior change elements: Elements used to promote behavior change. P2P: Person-To-Person.

studies (weight loss: 1.3 - 5.4 kg), compared to paper and pencil methods in 2/13 (15 %) (weight loss: 1.1 - 3.96 kg).<sup>67,71</sup> Of the automated response feedback studies,<sup>64,69,72</sup> only one (33 %) was a positive trial, while of the feedback delivered by personnel (phone, in-person, or email),<sup>58–63,65–68,70,71,73</sup> 11/13 (85 %) were positive trials.

## **Bias**

Trials scored between 4 and 8 of nine points for both the Delphi and EPOC bias criteria (Supplemental Table 1). Of studies meeting seven or more EPOC criteria, 58,60,62,63,65,67,68,72 weight loss ranged from 1.5 kg to 5.4 kg, whereas those meeting less than seven EPOC criteria, 59,61,64,66,69-71,73 reported 0.08 to 3.36 kg weight loss. Blinding of the provider, patient, and outcome assessor were the most common sources of bias.

#### Pragmatism

PRECIS scores generally, leaned more toward pragmatic (mean: 2.8 [SD 0.46], range: 2 - 3.6) (Fig. 4 and Supplemental Table 3). On average, studies scored lower (more explanatory) on follow-up intensity (1.8/4 [SD 1.2]) (i.e., subjects had more frequent visits and additional data collection than in routine practice), experimental intervention flexibility (2.0/4 [SD 1.1]) (i.e., strict intervention protocols), participant compliance (2.2/4 [SD 0.66]) (i.e., compliance was closely monitored and included measures to maintain and regain high compliance), and practitioner expertise in the experimental groups (2.4/4 [SD 1.3]) (i.e., used only highly trained practitioners). Studies scored highest (more pragmatic) in analysis of the primary outcome (3.4/4 [SD 1.2]) (e.g., intention-to-treat analysis), flexibility in the control group (3.3/4 [SD 1.4]) (i.e., usual practice control), and practitioner expertise in the control group (3.6/4 [SD 0.81))(i.e., additional expertise and training were not required). Five (31 %) trials were highly pragmatic, scoring on average > 3/4 on the PRECIS scale.<sup>62-64,67,72</sup> Of note.

our scores remained within one standard deviation of previously published data.<sup>44</sup>

#### DISCUSSION

Results of this review demonstrate that compared to usual PC, most  $(12/16, 75 \%)^{58-63,66-70,73}$  technology-assisted interventions in the PC setting help patients to achieve significant weight loss, indicating that technology can supplement and enhance the work of the PCP for weight loss outcomes. These technology-assisted interventions employ elements already demonstrated to be effective in facilitating weight loss: selfmonitoring, in-person feedback, and targeted, structured lifestyle coaching.<sup>75</sup>

The degree of weight loss in this review compares favorably to other PC-based weight loss interventions without technology. Weight loss results in Tsai and Wadden's review<sup>50</sup> of PC obesity treatment via counseling and pharmacotherapy (0.1 kg–7.7 kg) and in McCombie and colleagues' appraisal<sup>76</sup> of PC options in the UK (1.1 kg–6.6 kg) were overall similar to our findings (0.08 kg–5.4 kg). Attrition rates were also comparable to those in our review. At the very least, this portrays technology-assisted weight loss in the PC setting as having similar outcomes to traditional methods and offers further options for PCPs to consider.

Moreover, technology may give PCPs and patients the option to undergo weight loss intervention semi-remotely. Appel et al.<sup>58</sup> demonstrated that remote treatment produced weight loss comparable to an in-person intervention. Ma et al.<sup>63</sup> showed that in-person and self-directed Diabetes Prevention Program (DPP) arms produced comparable weight loss, while Spring et al.<sup>70</sup> similarly confirmed the additive benefit of remote support. This is promising, given that many practices are moving toward the patient-centered medical home model with office visits less central to PC practice.

Given the variation among the 12 studies with significant weight loss, surmising the most effective practices and/or their amalgamation is difficult. Our results suggest that interventions employing clinician-guiding software and feedback from personnel may be more likely to promote weight loss, as 86 % and 85 % of studies using these tools showed significant weight loss, respectively. In contrast, interventions without personnel (fully automated) were less likely to do so, with only 33 % demonstrating weight loss, suggesting that technology cannot fully replace human interactions with the healthcare team. Furthermore, whether employing multiple interventions (e.g., Bennett [2012] et al.<sup>60</sup>) is more effective than utilizing fewer technology modalities (e.g., Christian et al.<sup>61</sup>) remains unclear, as in this example both studies led to 1.5 kg weight loss. Similarly, we cannot determine the ideal combination of practitioners. While teams without MDs showed the highest weight loss (Fig. 2), the *n* is too small to justify any robust conclusions. Further trials could study the effect of specific team groupings on weight outcomes.

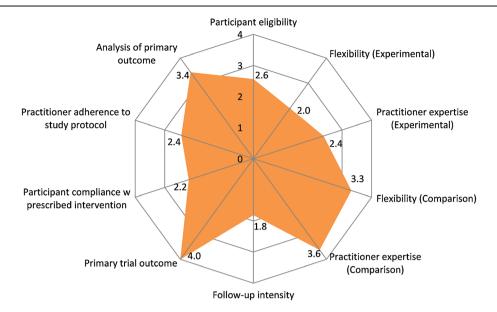


Figure 4. Mean PRagmatic Explanatory Continuum Indicator Summary (PRECIS) Scores. 0-Completely Explanatory (ideal circumstances)  $\rightarrow$  4-Completely Pragmatic (real-world circumstances). The following descriptions indicate entirely pragmatic trials (see Thorpe et al.<sup>45</sup> for detail): *Participant eligibility* – those with the condition of interest are enrolled without exclusions; *Flexibility* (experimental arm) – practitioners have considerable leeway in applying the intervention; *Practitioner expertise* (experimental arm) – no special training or clinical setting is required; *Flexibility* (comparison arm) – usual care is allowed; *Practitioner expertise* (comparison arm) – same as experimental arm; *Follow-up intensity* – no formal follow up; *Primary trial outcome* – clinically meaningful and testable under usual conditions; *Participant compliance* – unobtrusive or no measurement of compliance with no rescue strategies; *Practitioner compliance* – same as participant compliance; *Analysis of primary outcome* – employs intention to treat analysis.

Of the four studies that did not achieve significant weight loss, each had specific deficiencies. ter Bogt et al.<sup>71</sup> randomized at the level of the patient, not practice, potentially introducing contamination bias as the study progressed. They also transitioned to three yearly NP visits after the first year, an intensity less likely to promote weight loss.<sup>52</sup> Verheijden et al.<sup>72</sup> suffered from poor website uptake (33 %) and no change in online social support. McConnon et al.<sup>64</sup> had similar problems of poor website utilization (29 %) and high attrition (41 %). Finally, McDoniel et al.<sup>65</sup> demonstrated an age discrepancy between those who completed the study (older) and those who did not (younger).

For web-based interventions, poor web utilization was common. For example, in Bennett (2012) et al.,<sup>60</sup> only 25 % of participants used a self-monitoring platform for at least 75 % of trial weeks for unclear reasons. We speculate this was due to outdated design unable to keep pace with rapidly changing web standards<sup>77,78</sup> or an age-related "digital divide" as seen in Verheijden et al.,<sup>72</sup> where users were significantly younger than nonusers.

Only 4/16 (25 %)<sup>58,60,63,71</sup> interventions lasted more than 1 year, with most lasting for 12 months (6/16, 38 %)<sup>61,62,64,68,70,73</sup> or less (6/16, 38 %).<sup>59,65–67,69,72</sup> All reported outcomes at the end of the intervention, except for Spring et al.,<sup>70</sup> Nanchahal (2012) et al.,<sup>68</sup> and Ma et al.<sup>63</sup> (extension Xiao et al.<sup>79</sup>), who had maintenance phases of 6, 4, and 9 months, respectively. Whether after 1 year shorter trials would have continued to show significant weight loss is unclear. Judging from other weight loss literature and ter Bogt et al., where encouraging 1 year data<sup>74</sup> unfortunately resulted in insignificant 3-year weight loss,<sup>71</sup> the need for long-term interventions and/or follow-up post intervention greater than 1 year is imperative to provide patients with long-term outcomes.

Overall, the studies were of moderate to excellent quality. Our findings on Delphi and EPOC are similar to previously published results.<sup>80</sup> Due to the nature of this research, blinding was difficult. Studies of higher quality ( $\geq$  7 EPOC criteria)<sup>58,60,62,63,65,67,68,72</sup> reported greater weight loss, an encouraging finding. Interestingly, and not part of our study's initial intent, Delphi and EPOC provide comparable quality rankings.

PRECIS scores suggest that existing PC-based technologyassisted weight loss studies have many pragmatic elements. The PRECIS tool is helpful in pinpointing areas for improving pragmatism. For example, Ma et al.<sup>63</sup> had uniformly excellent scores with the exception of participant compliance (2/4) to intervention, suggesting that measurement of participant compliance could be less obtrusive. In general, interventions frequently scored poorly with respect to follow-up intensity, participant compliance, flexibility, and practitioner expertise, all key areas for successful real-world translation and implementation. This is concerning, as trials with low pragmatism are likely difficult to replicate and disseminate. Thus, future studies should strive for pragmatic design, particularly with attention to the aforementioned areas. On the other hand, the five trials that averaged > 3/4 on the PRECIS scale can serve as models for future protocol design.

## **Areas for Future Research**

Unfortunately, most studies are still performed outside of the PC setting—we excluded 76/101 studies during full text review for this reason. Given the paucity of PC-based trials, we suggest adapting technology-assisted interventions from non-PC settings into the office. Multiple innovative technologies tested only outside of PC have been summarized elsewhere and are awaiting possible implementation.<sup>40,42,81,82</sup> For instance, Ma et al.<sup>63</sup> adapted the DPP to the PC office, while Appel et al.<sup>58</sup> and Bennett (2012) et al.<sup>60</sup> similarly demonstrated an important step toward tailoring interventions for PC. Also encouraging is Morgan et al.'s Self-Help, Exercise, and Diet using Information Technology (SHED-IT) series. Translation of SHED-IT from a university<sup>83</sup> to a community<sup>84</sup> sample is encouraging, as further adaptation into PC appears a straightforward step.

Some technologies are currently under-represented in the literature, despite their widespread use. Interestingly, only one intervention in this review utilized texting, a consistently popular tool despite continued technology advances. A focus on texting could prove fruitful, as seen in smoking cessation<sup>85,86</sup> and treatment adherence research.<sup>87,88</sup> Of note, Heart360.org added texting capability after initiation of Ma et al.'s<sup>63</sup> study. Similarly, only one study employed a mobile application, even though internet traffic is increasingly occurring on mobile platforms.<sup>89</sup> Another untapped area similarly lies in social media, for its peer support outlets.<sup>90</sup>

We found it remarkable that few (2/16, 13 %) studies presented open-source or non-proprietary interventions.<sup>63,73</sup> Most websites and tools studied in trials are entirely unavailable to the practicing PCP, even though many patients could conceivably benefit from access. Instead, a poorly or unstudied commercial alternative is often the only remaining choice.<sup>43</sup> Notable exceptions to this include the AHA's Heart360, the MOVE!23 questionnaire and its mobile app, and the Healthy Highways website.

Currently missing from the literature is a cost-effectiveness analysis of technology-assisted weight loss interventions in PC. Only four trials<sup>64,73,91,92</sup> provide a critical appraisal of their costs. Also presently unclear is the usability of many of the trials' technology. Formal usability studies may help to improve reach and uptake of these weight loss interventions. Furthermore, the role of different clinicians in providing weight loss interventions in PC remains undetermined. Non-MD providers as a group were utilized most often, but large differences in training and role existed among them. As we continue to transition into the era of accountable care organizations and PC medical homes, multiple providers other than MDs will be expected to execute technology-assisted interventions.

## Limitations

The search terms may not have identified all pertinent studies. We find this unlikely, given the large number of articles extracted and the use of a research librarian to conduct the search. The participants in the included trials also may not represent the ethnic, gender, or age breakdown of typical PC practices. Some studies focused solely on disadvantaged populations,<sup>60–62</sup> while others occurred in ethnically homogenous countries,<sup>72</sup> perhaps precluding generalization. Furthermore, a "digital divide" may exist both with respect to socioeconomic status (SES) and age, whereby those with lower SES and older age may have difficulty interacting with technology-assisted interventions.<sup>93</sup> Moreover, while there are only a small number of RCTs on this topic, there are many prospective cohort studies, yet these may not provide us with sufficient rigor to guide clinical decision-making.

#### CONCLUSIONS

At the present time, there is no "first-line" therapy for technology-assisted weight loss in PC, but several meaningful choices exist that compare favorably to traditional weight loss interventions. Results of this review reveal that little is known about the optimal use of technology for weight loss in the PC setting. From this review, we can conclude the following: (1) technology-assisted weight loss is a valid tool for PCPs as they counsel patients, although a best iteration has yet to be determined; (2) technology-assisted weight loss interventions compare favorably to other modalities; and (3) longer, pragmatic, interdisciplinary, open-source interventions are needed.

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