

Results from the dissemination of an evidence-based telephone-delivered intervention for healthy lifestyle and weight loss: the Optimal Health Program

A Goode, PhD,¹ M Reeves, PhD,¹ N Owen, PhD,^{1,2} E Eakin, PhD

¹Cancer Prevention Research Centre, School of Population Health, The University of Queensland, Herston Rd., Herston, Queensland 4006, Australia

²Baker IDI Heart and Diabetes Institute, Melbourne, Victoria 3004, Australia

Correspondence to: A Goode
a.goode@sph.uq.edu.au;

Cite this as: *TBM* 2013;3:340–350
doi: 10.1007/s13142-013-0210-7

ABSTRACT

Despite proven efficacy, there are few published evaluations of telephone-delivered interventions targeting physical activity, healthy eating, and weight loss in community dissemination contexts. This study aims to evaluate participant and program outcomes from the Optimal Health Program, a telephone-delivered healthy lifestyle and weight loss program provided by a primary health care organization. Dissemination study used a single-group, repeated measures design; outcomes were assessed at 6-month (mid-program; $n=166$) and 12-month (end of program; $n=88$) using paired analyses. The program reached a representative sample of at-risk, primary care patients, with 56 % withdrawing before program completion. Among completers, a statistically significant improvement between baseline and end of program was observed for weight [mean change (SE) -5.4 (7.0) kg] and waist circumference [-4.8 (9.7) cm], underpinned by significant physical activity and dietary change. Findings suggest that telephone-delivered weight loss and healthy lifestyle programs can provide an effective model for use in primary care settings, but participant retention remains a challenge.

KEYWORDS

Dissemination, Telephone, Weight loss, Physical activity, Diet, Intervention

INTRODUCTION

Despite well-known health benefits of physical activity and healthy eating [1–3], majority of adults in economically developed countries fail to meet minimal public health recommendations [4, 5]. Related to this are increasing rates of overweight and obesity, and lifestyle-related diseases [6–8]. Thus, effective and broad reaching interventions to promote regular physical activity and healthy eating as well as modest weight loss are required. Mediated (non-face-to-face) intervention delivery in the form of print, Internet, and telephone [9–11] offers a potentially flexible, convenient, and cost-effective means of providing the repeated contacts necessary to achieve and maintain behavior change [12]. Telephone delivery is the most widely researched

Implications

Research: To enhance impact of evidence-based telephone-delivered interventions for physical activity, healthy eating, and weight loss, research that elucidates ways to improve retention in “real-world” settings is needed.

Practice: Healthy lifestyle interventions delivered via telephone by community primary care organizations can have a significant impact on patient health behaviors (i.e., physical activity and diet) and health outcomes (i.e., weight, waist circumference, and cholesterol).

Policy: Programs like the OHP offer potential to deliver a more accessible and cost-effective intervention to promote longer term behavior change and weight management, but funding is needed to help determine ways to improve program retention.

of these modalities to date and remains the most accessible [9, 12]. Such interventions are distinctly promising given their potential to be adopted by health organizations that operate telephone information and support centers [12, 13].

A strong randomized trial evidence base supports efficacy of telephone-delivered interventions targeting physical activity and/or dietary change and weight loss in a range of settings and target populations [9, 10, 14–19]. In order to achieve their potential public health impact, such interventions need to be implemented and evaluated in diverse community practice settings [20–23]. To date, only three large-scale dissemination studies have reported on the outcomes of telephone-delivered lifestyle interventions, two targeting physical activity only [24–26] and one targeting modest weight loss via physical activity and diet [27]. Findings from these telephone-delivered dissemination studies suggest that evidence-based interventions can be delivered successfully to achieve results comparable to those observed in controlled research settings and, in so doing, may even reach more diverse samples [24–26, 28].

The Logan Healthy Living Program is an evidence-based telephone-delivered 12-month inter-

vention targeting physical activity and healthy eating [29, 30]. Its recent uptake by a primary health care organization provided opportunity to evaluate participant and program outcomes within an applied practice setting. Detailed study methods and outcomes from the randomized controlled trial of the Logan Healthy Living Program have been described [29–31]. In brief, the trial, which targeted adults with type 2 diabetes and/or hypertension, demonstrated significant between-group improvement, favoring the intervention group, for all dietary outcomes, including total and saturated fat, and vegetable, fruit, and fiber intake. Significant within-group improvement was observed for physical activity for both intervention and usual care groups [30]. Diet and physical activity improvements were largely maintained at a 6-month postintervention follow-up [32] and the intervention was shown to be cost effective [33].

Adaptations and supports that were necessary to facilitate adoption and implementation of the program, now known as the Optimal Health Program (OHP), in a community setting have also been described; the adaptations include broadening program focus to include modest weight loss (i.e., –5 to 10 % of initial body weight), in addition to promotion of physical activity and healthy eating [34], for overweight patients without chronic illness. This was done to avoid duplication of services as the adopting organization offers a range of program and support initiatives around self-management of chronic illness including diabetes and heart disease. This paper describes the evaluation of the OHP. Given the dissemination context, indicators of both internal and external validity are addressed [35–37] via reporting on both program (i.e., adoption, reach, characteristics of participants vs. nonparticipants and completers vs. dropouts, and implementation) and participant outcomes (i.e., weight; waist circumference; HDL-, LDL-, and total cholesterol; systolic and diastolic blood pressure; fruit and vegetable intake; total time for moderate-to-vigorous physical activity (MVPA); and total screen time).

METHODS

Study design

Given that effectiveness of the telephone-delivered intervention had previously been established in a randomized trial and that the primary research question in this study was about outcomes that could be achieved in an applied practice setting (i.e., dissemination context), a single group, pre-postdesign was used, as is common for dissemination research [24, 25, 38–40]. As the OHP program is ongoing, data presented here come from a “snap shot” of participant and program outcomes after approximately 2.5 years of the program becoming fully operational, with a census date of April 15, 2012. OHP participants were assessed at baseline and at 6 months (mid-program and end of the more

intensive phase of telephone contacts) and 12 months (end of program). The study protocol was approved by the School of Population Health Research and Ethics Committee, The University of Queensland, Australia.

Setting

The OHP was taken up for delivery by the Greater Metro South Medicare Local. The Greater Metro South Medicare Local is a state- and federal-funded organization that provides administrative, technical, professional development, and educational support to primary medical care practices within the Logan area south of Brisbane, Queensland, Australia. The Logan area (population 277,000) is a large, ethnically diverse community characterized by higher levels of social disadvantage compared to Brisbane (the state capital) and Queensland, including a greater percentage of single-parent families, unemployment, and residents born overseas [41]. At the time the program was initiated, the area was supported by 80 primary care practices with 304 general practitioners (GPs).

Practice and patient recruitment

The OHP began recruitment of general practices within the Logan area primarily through notices within general practice newsletters. Practices were also invited to participate through expression of interest at committee meetings, promotional events, and conferences. Once an expression of interest was received, OHP staff completed a practice visit with GPs and/or practice nurses. During these visits, practices were provided with information kits detailing the program including eligibility criteria for participants, program brochures, referral forms, and participant outcome reporting forms. GPs screened potentially eligible patients for OHP referral. To be eligible for participation, patients needed to be at least 18 years of age, have a body mass index (BMI) equal or greater than 25 kg/m², and have no chronic disease (other than hypertension, arthritis, osteoporosis, dyslipidemia, depression, or anxiety). Patients were excluded if they were unable to participate in telephone counseling (e.g., no telephone and unreachable by phone for extended periods) and if the doctor determined that participation in unsupervised moderate-intensity physical activity or strength training was contraindicated. Once referrals were received by the Medicare Local, an additional screening call (to double check eligibility) was conducted by the OHP counselors before recruitment into the program. During this call, verbal informed consent was obtained for the collection of data for evaluation purposes.

Intervention: the Optimal Health Program

The OHP intervention protocol closely followed the original Logan Healthy Living Program [29]; given the change in target group, it also included evi-

dence-based weight loss protocols [42]. It involved delivery of a total of 18 intervention calls, delivered weekly for the first 4 weeks, then fortnightly until 4 months and then monthly for the remaining 8 months. Although the ideal frequency of calls was specified based on the Logan Healthy Living Program protocol [29, 30], flexibility in the timing of calls was allowed, consistent with the norms of the clinical practice-based approach being used. (For example, participants may have received an extra call during the monthly phase, if required). Program calls were intended to last approximately 20–30 min. Program protocol allotted up to five call attempts before a participant was withdrawn from the program.

Participants were mailed a workbook, pedometer, stretch band, tape measure, calorie (fat and fiber) counter, community lifestyle directory with details of subsidized physical activity programs within the local area, and off-the-shelf brochures on diet and physical activity guidelines. In addition to sections on physical activity, diet, and weight loss, the workbook addressed behavior change strategies consistent with social cognitive theory [43], including goal setting, problem solving, self-rewards, social support, positive self-talk, and relapse prevention [29]. Telephone counselors regularly referred to the workbook during the 12-month program, emphasizing the development and ongoing review of achievable physical activity, diet, and weight loss goals. A patient-centered motivational interviewing approach [44] to the telephone counseling was used.

Targets for diet and physical activity were consistent with national guidelines [45–49]. Drawing on newer evidence on the importance of reducing sitting time [50], participants were encouraged to limit nonwork-related screen time to no more than 2 h/day. Consistent with the evidence on weight loss for chronic disease prevention, participants were encouraged to lose 5–10 % of their body weight over the 12-month program and weight loss protocols followed evidence-based guidelines [51, 52].

Staff training

Telephone counselors were accredited practicing dietitians, all with bachelor's level training in nutrition and dietetics. Counselors initially received an intensive 5-day in-house training program conducted by research staff on intervention procedures, recruitment, screening and assessment methods, follow-up protocols, data entry, and motivational interviewing strategies [44]. Additionally, a half-day training workshop with an exercise physiologist was provided to ensure adequate skills related to physical activity promotion (specifically around strength training). Regular phone and email contact, and monthly to bimonthly face-to-face meetings with research staff supported implementation and addressed quality control of program delivery (via case conferences) and data collection

(via regular checks for accuracy of entry); however, only call delivery and duration were systematically tracked and recorded. A total of 2.2 full-time equivalents were devoted to OHP program delivery by three counselors.

Outcomes

Program outcomes

Participant baseline sociodemographic variables (i.e., age, sex, marital status, highest education attainment, employment status, and income) were collected via telephone by OHP counselors in order to be able to describe the characteristics of participants vs. nonparticipants and OHP completers vs. dropouts. Data related to program delivery (i.e., number and duration of calls completed) were tracked by OHP counselors. At both the mid- and end-of-program assessments, participants were asked by counselors to rate how helpful they found the program overall on a 10-point Likert scale, from 1 “not helpful at all” to 10 “extremely helpful”.

Participant outcomes

Participant outcomes included objectively measured clinical (weight; waist circumference; HDL-, LDL-, and total cholesterol; and systolic and diastolic blood pressure) and self-reported behavioral outcomes [fruit and vegetable intake, total time for moderate-to-vigorous physical activity (MVPA), and total screen time]. For all participant outcomes, mid- and end-of-program assessments were scheduled at approximately 4–6 months and 12 months, allowing more flexibility with the scheduling than in a controlled research study in view of the constraints of conducting program evaluation in a community setting with rolling recruitment.

Clinical outcomes were collected via GP or practice nurse at baseline, and each follow-up time point. Behavioral outcomes were collected via telephone by OHP counselors and included: the same validated measures used in the original randomized controlled trial [29, 30] as well as demographic data at baseline. Servings of fruit and vegetables were assessed using two items from the validated Australian National Nutrition Survey [53, 54]. Self-reported physical activity was measured using the Active Australia Survey [4]. Total weekly minutes of MVPA was calculated from the sum of walking, moderate, and 2× vigorous minutes, first truncating each activity at 840 min/week and truncating total MVPA at 1,680 min/week [4]. Four items were used to assess total time spent sitting in the last week across two domains: (1) watching television, videos, or playing electronic games and (2) leisure-time computer use [55]. Adverse outcomes were assessed by asking participants if they had any new health problems since the previous assessment.

Data analysis

The “snap shot” evaluation utilized data from participants enrolled in the program from its inception until mid-April 2012. Analyses that required data from mid- or end-of-program assessments excluded those participants who had not been enrolled in the program long enough to have reached those assessment time points.

Data analyses were conducted in SPSS for Windows (version 18). Statistical significance was set at $p < 0.05$, two-tailed. Baseline characteristics of participants vs. non participants as well as those who completed the mid- (6 months) and those who completed end-of-program (12 months) assessments (completers) vs. those who withdrew before each assessment point (dropouts) were compared using independent-sample *t* tests, Wilcoxon signed-rank tests (for variables not normally distributed), and chi-square test statistics. Similarly, changes in program outcomes from baseline to 6 months were also compared between those who completed the 12-month assessment and those who withdrew between mid- and end-of-program assessments. Statistically significant and meaningful differences are noted (the latter defined as ≥ 10 % absolute difference for categorical variables or ≥ 10 % difference in means for continuous variables).

Effectiveness of the program was assessed by examining whether participants who completed the program assessments changed significantly from baseline to mid- or end-of-program in their clinical and behavioral outcomes using paired *t* tests and Wilcoxon signed-rank tests. Program outcomes are presented as means (standard deviations) for normally distributed outcomes and medians (minimum and maximum values) for outcomes that did not follow a normal distribution.

RESULTS

Program outcomes

Adoption, reach, and characteristics of participants vs. nonparticipants

After approximately two and a half years of being fully operational, the OHP had been adopted by 23/80 general practices (29 %) and had received 377 referrals, with 317 participants consenting to participate and completing the baseline assessment. Recruitment and retention of participants are shown in Fig. 1.

Participant characteristics at baseline are presented in Table 1. At time of entry into the program, the age range of participants was 18 to 77 years [mean (SD) = 46.4 (11.8) years] and BMI ranged from 25.3 to 76.8 kg/m² [mean (SD) = 37.0 (7.7) kg/m²], with 48.5 % having a BMI of greater than or equal 35 kg/m². Participants were predominantly female, Caucasian, and married. However, the sample also included a notable percentage of ethnic minorities, including Aboriginal/Torres Strait Islander and Pacific

Islander populations (7 %), those unemployed (11 %), and with low educational attainment—junior high school or less (41 %). According to referral data (gender, age, BMI, weight, and waist circumference), there was no statistically significant or meaningful difference between those who consented vs. those who declined to participate (data not shown).

Attrition and characteristics of completers vs. dropouts

As of the census date, of the 279 participants enrolled in the program long enough to complete the mid-program assessment, 166 completed it, 107 withdrew from the program, and six had not withdrawn but had assessments outstanding. Approximately one-third of those who dropped out before the mid-program assessment did so after completing only one counseling call. Of those who completed the mid-program assessments, 136 had also been in the program for long enough to complete their end-of-program assessment; 88 completed it, 39 withdrew, and nine remained outstanding. The withdrawal rates were 38 % (107/279) up to the mid-program assessment and 29 % (39/136) between mid- and end-of-program assessments. In total, approximately 44 % of participants who commenced the OHP completed the program and 12-month assessment.

Table 2 shows the demographic and baseline variables of those who completed the mid-program assessment (completers; $n=166$) and those who withdrew before completion (dropouts; $n=107$). A comparison of the two groups revealed that completers were significantly more likely to be older than those who withdrew. There was a nonsignificant but meaningful difference in physical activity, with completers reporting higher levels of MVPA per week at baseline than dropouts.

Table 2 also shows the demographic and baseline variables for those who completed the end-of-program assessment (completers; $n=88$) and those who withdrew between the mid- and end-of-program assessments (dropouts; $n=39$). A comparison of the two groups indicates that completers and dropouts at the end-of-program time point varied on similar indicators as was observed for the mid-program assessment time point, except that non-Caucasians were more likely to complete end-of-program assessments, as were those with lower incomes, and lower fruit intake at baseline. Compared to those who completed the end-of-program assessment, those who withdrew between mid- and end-of-program assessments achieved smaller adiposity changes from baseline to the mid-program assessment [weight: mean change (SD) = -3.7 (5.5) kg vs. -1.3 (5.9) kg, $p=0.1$; waist circumference: -4.2 (6.9) vs. -1.9 (5.5) cm, $p=0.3$, respectively] but reported larger behavioral changes [including MVPA: 63.7 (215.8) mins/week vs. 153.5 (201.8) mins/week, $p=0.03$; and fruit intake: 0.4 (1.2) vs. 0.2 (1.2), $p=0.3$].

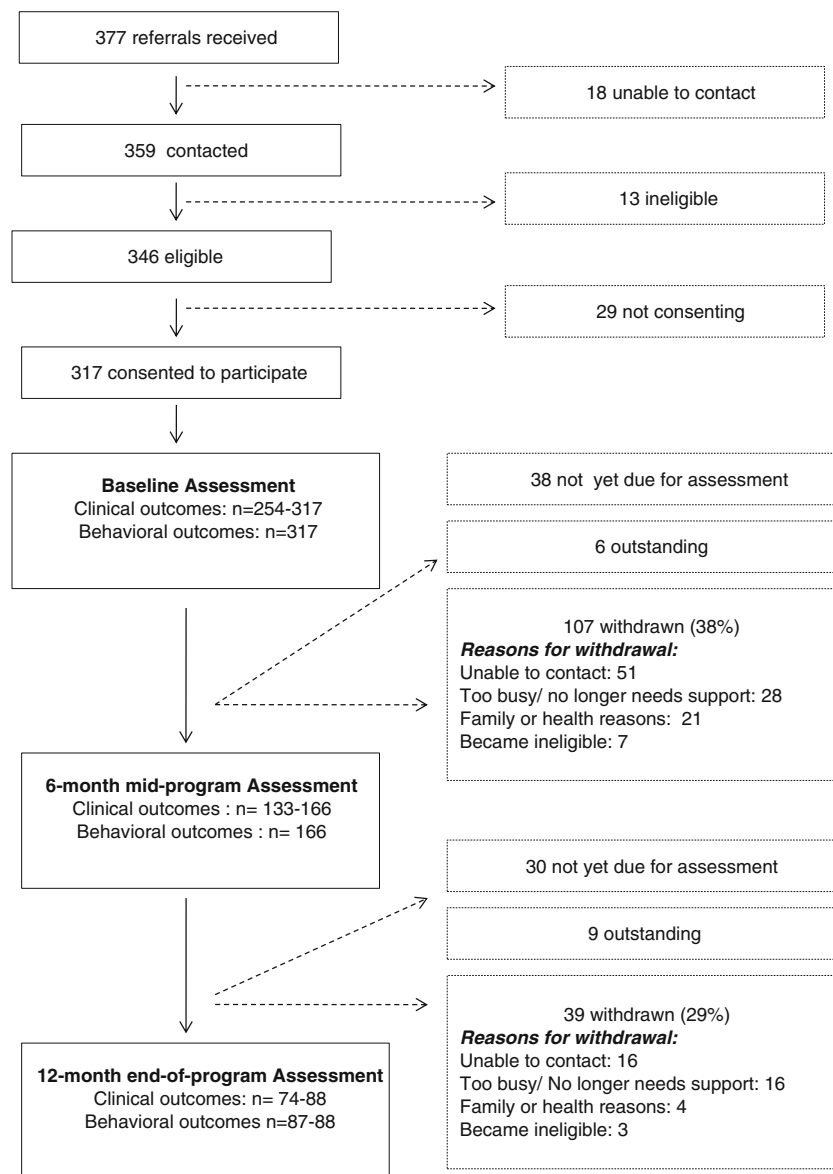


Fig 1 | Flow chart of participant recruitment

Reasons for attrition are listed in Fig. 1. Most commonly, for the mid-program assessment, participants were unable to be contacted ($n=51$). Participants who actively withdrew from the program before completing the mid-program assessment most commonly cited family or health reasons ($n=21$) or that they had become too busy or no longer needed support ($n=28$). Similarly, participants who did not complete the end-of-program assessment were most commonly unable to be contacted ($n=16$).

Implementation

For those remaining in the program at 6 months ($n=166$), the median number of calls received was 10 out of approximately 12 recommended calls (range=1–15). For those who had completed the entire 12-month program ($n=88$), the median num-

ber of calls received was 16 of approximately 18 recommended calls (range=7–23). Call duration ranged from 6 to 55 min, with an average duration of 29 min. On a scale of 1 to 10, with 10 being “extremely helpful”, 69/78 (86 %) rated the program an “8” or above at end-of-program.

Participant outcomes

Change from baseline to mid-program

As shown in Table 3, for those completing the mid-program assessment, there was a statistically significant improvement between baseline and 6 months for all clinical outcomes (i.e., BMI, weight, waist circumference, total cholesterol, and systolic and diastolic blood pressure), except HDL-cholesterol. Mean weight loss of 3.3 % (± 5.6) of initial body weight [mean change (SD): -3.3 (5.7) kg] was observed at 6 months.

Table 1 | Baseline characteristics of participants

Baseline characteristic	Mean (SD) or %N=317 ^a
Female	74 %
Age (years)	46 (12)
BMI (kg/m ²)	37.0 (7.7)
Obesity class I	34 %
Obesity class II	27 %
Obesity class III	25 %
Weight (kg)	103.1 (23)
Waist circumference (cm)	112.1 (16.8)
Total cholesterol (mmol/L)	5.3 (1.0)
HDL-cholesterol (mmol/L)	3.3 (0.9)
LDL-cholesterol (mmol/L)	1.3 (0.4)
Systolic BP (mmHg)	128.1 (17.4)
Diastolic BP (mmHg)	80.8 (10.7)
Ethnicity	
Caucasian	88 %
Marital status	
Married/living together	73 %
Education	
≤High school	41 %
Completed high school	43 %
Tertiary qualification (trade/diploma/university degree)	16 %
Employment	61 %
Employed (FT, PT, casual)	21 %
Retired/home duties	18 %
Unemployed/student/other	
Household income	30 %
≤\$999/week	59 %
≥\$1,000/week	11 %
Declined to answer/don't know	

Obesity class I=BMI 30.0–34.9 kg/m²; class II=35.0–39.9 kg/m²; class III≥40.0 kg/m²

^a Weight and BMI, n=315; waist circumference, n=297; cholesterol, n=306; HDL- and LDL-cholesterol, n=254; diastolic 7 systolic BP, n=304; for income, n=312

There were statistically significant improvements for all self-reported behavioral outcomes, including physical activity [+105 (231) total MVPA min/week] and dietary behaviour [fruit: +0.4 (1.1) serves/day; vegetables: +0.9 (1.5) serves/day] between baseline and mid-program for those completing the 6-month assessment. No adverse outcomes as a result of participating in the program were reported at the mid-program assessment.

Change from baseline to end of program

As shown in Table 4, for those completing the end-of-program assessments, there were improvements between baseline and 12 months for all clinical outcomes, with these reaching statistical significance for BMI, weight, waist circumference, and diastolic blood pressure. Mean weight loss of 5.5 % (±6.8) initial body weight [mean change (SD): −5.4 (7.0) kg] was observed for those completing the 12-month assessment, with 48 % (28/59) of participants having

met or exceeded the 5 % weight loss goal of the program.

As shown in Table 4, participants who completed the end-of-program assessment reported statistically significant improvements in self-reported behavioral outcomes including physical activity [+83 (249.7) min/week] and vegetable intake [+1.0 (1.7) serves/day], but not fruit, with median intake remaining at the recommended two serves per day. No adverse outcomes were reported at the end of program.

DISCUSSION

There have been numerous calls for increased efforts to disseminate effective chronic disease prevention and management interventions [22, 25, 35, 37, 56], with more recent attention to their translation into “real-world” settings [57, 58]. The OHP is unique to our knowledge, as it represents the first effort to translate and evaluate a telephone-delivered lifestyle intervention targeting weight loss within an applied primary health care setting. Overall, participant outcomes from the “snap shot” evaluation indicate promising effectiveness for weight loss and other clinical outcomes, underpinned by dietary and physical activity change. Those who completed the end-of-program assessment showed clinically meaningful improvement, losing on average 5.5 % of their body weight from baseline. Almost half of participants who completed the program achieved at least 5 % weight loss, although it is important to note this was based on a small number of participants, given 44 % retention at 12 months. This magnitude of weight loss has been associated with beneficial health outcomes and is meaningful in terms of both individual and population health [59–61].

An Australian telephone-delivered lifestyle program [offered as a statewide government health department-funded service—the Get Healthy Information and Coaching Service (GHS)] provides the most comparable source of data for OHP outcomes [62]. Similar to the OHP, the GHS targeted physical activity and diet as well as modest weight loss, but with all outcomes collected via self-report. In contrast to the OHP, the GHS involved 6 months of telephone coaching and broadly targeted the general adult population, mainly based on self-referrals following ongoing media campaigns [27, 28], with a smaller number of participants coming from secondary referral sources that included health practitioner referrals [28]. From a snap shot of 1,440 participants, the GHS reported statistically significant improvements in weight [−3.9 kg (5.1)] and waist circumference [−5.0 cm (6.0)], remarkably similar to the corresponding objectively measured anthropometric outcomes seen in the mid-program assessment of the OHP.

Overall, weight loss achieved within the OHP compares favorably to evidence from the broader array of studies that have attempted to translate the intensive Diabetes Prevention Program into delivery

Table 2 | Baseline characteristics of those who completed vs. those who withdrew before completion of 6-month assessment and those who completed vs. those who withdrew between 6- and 12-month assessment

Baseline characteristics	6-Month assessment		12-month assessment	
	Completers N=166 ^a Mean (SD) or median [min, max]	Dropouts N=107 ^b Mean (SD) or median [min, max]	Completers N=88 ^c Mean (SD) or median [min, max]	Dropouts N=39 ^d Mean (SD) or median [min, max]
Demographics				
% Female	77 %	75 %	74 %	80 %
% Caucasian	87 %	90 %	90 %	74 %*
% Married	75 %	69 %	75 %	77 %
Senior high school or greater	61 %	57 %	58 %	62 %
Employed (FT, PT, casual)	61 %	63 %	59 %	62 %
% income >\$1,000/week	57 %	61 %	50 %	72 %
Age (years)	47.3 (12.1)	44.0 (11.1)*	49.3 (12.0)	45.9 (10.3)
Clinical outcomes				
BMI (kg/m ²)	36.7 (7.2)	37.3 (8.8)	36.0 (7.1)	37.0 (6.8)
Weight (kg)	102.0 (22.1)	104.0 (25.0)	102.0 (23.4)	100.0 (19.7)
Waist circumference (cm)	110.7 (17.1)	112.6 (15.9)	110.0 (17.1)	113.0 (19.2)
Total cholesterol (mmol/L)	5.3 (1.0)	5.5 (1.1)	5.3 (0.1)	5.3 (0.9)
HDL-cholesterol (mmol/L)	1.3 (0.3)	1.3 (0.3)	1.3 (0.3)	1.3 (0.3)
LDL-cholesterol (mmol/L)	3.2 (0.9)	3.4 (0.8)	3.2 (0.9)	3.1 (0.9)
Systolic BP (mmHg)	129.2 (17.3)	126.0 (18.6)	128.4 (16.2)	130.3 (20.1)
Diastolic BP (mmHg)	81.3 (10.2)	80.0 (11.8)	81.1 (9.70)	81.7 (11.0)
Behavioral outcomes				
Vegetables (serves/day)	2.0 [0–10]	2.0 [0–8]	2.0 [0–10]	2.0 [0–8]
Fruit (serves/day)	1.0 [0–4]	1.0 [0–4]	1.5 [0–4]	1.0 [0–4]*
MVPA (min/week)	77.5 [0–1260]	60.0 [0–954]	95.0 [0–820]	60.0 [0–1260]
Screen time (min/day)	183.2 [14.3–900]	192.0 [0–613.0]	197.1 [47.1–900]	167.1 [14.–557.1]
<i>MVPA</i> moderate-to-vigorous physical activity				
^a Income, n=164; cholesterol, diastolic, and systolic BP, n=161; waist circumference, n=157 HDL and LDL, n=133				
^b Weight and BMI, n=106; waist circumference, cholesterol, income, n=104; systolic and diastolic BP, n=101; HDL and LDL, n=84				
^c Income, n=86; waist circumference, n=85; cholesterol, diastolic, and systolic BP, n=84; HDL and LDL, n=74				
^d Waist circumference and cholesterol, n=38; HDL and LDL, n=34				
* p≤0.05				

in a range of community and clinical settings. A review of 16 such studies found that weight loss ranged from -1.0 to -8.6 kg, with the percentage of participants meeting the 5 % weight loss goal ranging from 11 to 64 % [57]. However, studies included in the review were predominantly group based/face to face, with only one including some telephone contact [63]. The magnitude of physical activity and dietary improvements observed in the OHP is broadly comparable to other telephone-delivered dissemination studies [24, 26, 64] as well as the original trial upon which it was based [30].

In addition to reporting on participant (effectiveness) outcomes, a number of factors related to external validity that are important to informing the broader evidence around dissemination (i.e., adoption, reach, and retention) were also assessed as part of the OHP evaluation. At the practice level, initial adoption of the program was moderate, with just over one quarter of potentially eligible practices taking up the program to date. This is in line with

the practice recruitment rate (i.e., 27.8 %) observed in the original Logan Healthy Living Program trial [29] as well as other primary care-based trials [65, 66]. Encouragingly, all adopting practices of the OHP continue to refer patients into the program and expressions of interest from other practices remain forthcoming. Ongoing resources in the form of additional practice visits and follow-up telephone calls as well as regular mailed feedback to GPs concerning patient outcomes have been key strategies for sustaining referrals.

Importantly, the OHP appears to be successfully targeting overweight/obese primary care patients from lower socioeconomic backgrounds who are often difficult to reach and engage in behavior change programs [67]. Participants and nonparticipants were similar across demographic variables, indicating that the program was successful in recruiting a representative sample, including a notable percentage from ethnic minority groups, a finding also reported in the GHS [28]. However, as

Table 3 | Mid-program outcomes for participants who completed the 6-month assessment

Outcomes	Number	Baseline Mean (SD) or median [min, max]	6 months Mean (SD) or median [min, max]	<i>p</i> value ^a
Clinical				
BMI (m/kg ²)	118	36.3 (6.0)	35.1 (6.8)	<0.001
Weight (kg)	118	99.9 (20.8)	96.6 (20.4)	<0.001
Waist circumference (cm)	101	108.7 (14.3)	104.5 (15.4)	<0.001
Total cholesterol (mmol/L)	114	5.4 (1.0)	5.1 (1.0)	<0.001
HDL-cholesterol (mmol/L)	91	1.30 (0.3)	1.34 (0.5)	0.279
LDL-cholesterol (mmol/L)	90	3.28 (0.8)	3.12 (0.9)	0.023
Systolic BP (mmHg)	106	130.0 (17.1)	126.7 (13.0)	0.019
Diastolic BP (mmHg)	105	81.2 (10.4)	78.8 (8.6)	0.004
Behavioral				
Vegetables (serves/day)	166	2 [0–10]	3 [0–10]	<0.001
Fruit (serves/day)	166	1 [0–4]	2 [0–4]	<0.001
MVPA (min/week)	165	75 [0–1260]	200 [0–1080]	<0.001
Screen time (min/day)	166	183.2 [14.3–900]	138.5 [0–849]	<0.001

MVPA moderate-to-vigorous physical activity

^a *p* for paired *t* tests (normal data) or Wilcoxon signed-rank test (non-normal)

in the GHS (82 %) [28], notably more females took part in the OHP. A recent systematic review also found that the majority of participants in diabetes translation programs were female (i.e., 74 %), with this rate being higher than in the original Diabetes Prevention Program research trial (68 % female) [57]. Similarly, the number of females taking part is slightly higher for the OHP, 74 %, than the Logan Healthy Living Program, 61 % [30]. For the OHP, the overselection of women occurred during the referral process when potential participants presented at primary care practices, as 73 % of referrals were for female patients. It may be that men are opting out of the program at this point and, thus, never being referred, or GPs are

simply referring more women, given that women are more likely to present for preventive care [68].

High withdrawal rates observed in the OHP (38 % attrition at 6 months) are reflective of the “real world” context and are in line with other dissemination studies [24, 25, 57], including the GHS, which reported 74 % attrition at the end of the 6-month intervention [64]. It may be that participants in dissemination studies with interventions delivered in applied settings do not perceive themselves as making the same level of commitment to complete a program as those who formally consent to participate in a controlled research study, especially when the program is offered free of charge. Further,

Table 4 | End-of-program outcomes for participants who completed the 12-month assessment

Outcomes	Number	Baseline Mean (SD) or median [min, max]	12 months Mean (SD) or median [min, max]	<i>p</i> value ^a
Clinical				
BMI (m/kg ²)	59	35.2 (6.0)	33.3 (6.1)	<0.001
Weight (kg)	59	97.4 (2.1)	92.0 (20.9)	<0.001
Waist circumference (cm)	53	106.9 (15.2)	102.2 (17.2)	0.001
Total cholesterol (mmol/L)	56	5.3 (0.9)	5.1 (1.0)	0.144
HDL-cholesterol (mmol/L)	46	1.27 (0.4)	1.35 (0.4)	0.072
LDL-cholesterol (mmol/L)	46	3.13 (0.9)	3.00 (0.8)	0.292
Systolic BP (mmHg)	53	127.3 (13.2)	125.7 (15.4)	0.350
Diastolic BP (mmHg)	53	80.6 (10.5)	76.4 (8.1)	0.004
Behavioral				
Vegetables (serves/day)	88	2 [0–10]	4 [0–10]	<0.001
Fruit (serves/day)	88	1.5 [0–4]	2 [0–3]	0.312
MVPA (min/week)	87	95 [0–820]	170 [0–1180]	0.004
Screen time (min/day)	87	191.1 [49.3–900]	150 [0–810]	<0.001

MVPA moderate-to-vigorous physical activity

^a *p* for paired *t* tests (normal data) or Wilcoxon signed-rank test (non-normal)

nonresearch organizations, which often emphasize service delivery over evaluation, may be less likely to follow-up participants as extensively and systematically (due to staffing and budgeting constraints) as is typical in controlled research trials [69]. This was the case in the current study, where OHP participant follow-up protocols were much less stringent than in the precursor trial.

A recent review of attrition in weight loss trials showed that there were no consistent demographic, weight, or health behavior profiles that were associated with program dropout [70]. In the OHP, those who completed the program vs. those who dropped out were largely similar, except that younger and heavier participants were more likely to withdraw. Interestingly, those who withdrew after the 6-month assessment achieved less weight loss but self-reported larger behavioral changes from the start of the program to the mid-program assessment compared to those who remained in the program until the end-of-program assessment. Further clarification is needed to understand this finding. In any instance, promoting regular self-monitoring by participants of both weight and behavior can improve consistency between perceptions of changes made and actual behavioral and weight change progress [71].

Evaluation of OHP implementation, including the number and duration of calls completed, shows that the primary care organization largely followed evidence-based program delivery protocols, demonstrating that the program was able to be implemented with fidelity in the “real world”. It is important to note that resource constraints did not allow for more detailed quality assurance procedures (e.g., audiotaping and coding call content). Higher dropout rates in the beginning of the program indicate that the ability to implement the full 12-month intervention to all participants was challenging. This is an important issue given evidence from two recent systematic reviews of telephone-delivered physical activity and/or dietary behavior change interventions that indicate that delivery of longer term interventions (i.e., of at least 6-months duration) is associated with improved outcomes [9, 14]. It suggests that other modalities for providing ongoing intervention contacts should be evaluated [72].

Although nonrandomized, single-group pre-post designs are common in dissemination studies [24, 25, 40], the lack of a comparison group is a limitation in this study. However, evidence from Australian population-based prospective studies indicates an overall population trend to gain weight, with an average gain of 1.8 kg over 5 years in adults aged 18–65 years [8]. Further, there is evidence that those who are overweight/obese are more likely to continue to gain weight over a 5-year period [73]. Thus, it is a reasonable assumption that in the absence of the OHP, participants would have continued to gain weight.

Another limitation of the study is that behavioral outcomes were self-reported and collected by staff delivering the OHP, as was participant satisfaction data. This limitation is mitigated, to some extent, by corroborative objective data from GP-measured clinical outcomes. In our dissemination context, the complete standardization of data collection procedures was not feasible. However, all clinical outcomes (i.e., weight, waist circumference, blood pressure, and cholesterol) were collected at baseline and follow-up time points by the same GP or practice nurse for each participant. It was not feasible to have a standardized method (such as type of weight scale) across practices for collecting these outcomes. Within-person change was our primary outcome of interest; therefore, any error engendered by data collection procedures was likely to be consistent within individuals and, thus, not likely to threaten validity of outcomes obtained. It is also important to consider that this study reports on a complete analysis of a small number of participants, with our analyses showing that those who dropped out experienced poorer weight loss outcomes.

Summary and implications

Although small by dissemination study standards, findings from the OHP provide further support to a small but growing body of research that demonstrates that evidence-based lifestyle/weight loss interventions can be translated into practice and achieve outcomes, perhaps even with more representative samples, consistent with those observed in the original randomized trials. As previously described [34] strong and ongoing partnerships between the academic/research and primary care/community entities remain a key to both successful program implementation and the type of rigorous evaluation reported here. Future studies need to consider costs to deliver and cost effectiveness to further the evidence needed to inform future uptake into practice.

Acknowledgments: We thank the participating general practices in the Logan area and their patients as well as the staff of the Greater Metro South Brisbane Medicare Local. In particular, we would like to acknowledge the dedicated work of the accredited dietician telephone counselors, Jana Husak, Sarah Gruber, Natalie Cross, and Ashleigh Davidson and the support provided by Sherron Madden and Amada Adams.

Conflict of interest: None.

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