

Programmed shared medical appointments for weight management in primary care

An exploratory study in translational research



Garry Egger, John Stevens, Nerida Volker, Sam Egger

Background and objective

Programmed shared medical appointments (pSMAs) are an extension of shared medical appointments (SMAs) for managing chronic diseases. Acceptance and outcomes of a trial for weight management are reported here.

Method

Six programmed sessions in weight management were carried out in seven NSW primary healthcare centres.

Results

Seventy-seven per cent of 216 participants (64 male, 102 female) completed at least four of six pSMAs over 16–18 weeks ('adherers'). Eighty-five per cent of adherers approved of pSMAs, and 73% said they would prefer these over a standard clinical consult. Providers also rated the process highly. Overall mean weight loss over 12 months was 3.2% (95% confidence interval [CI]: 2.2, 4.3; $P < 0.001$). For adherers, mean weight loss was 4.3% for males (95% CI: 2.6, 6.1; $P < 0.001$) and 4.2% for females (95% CI: 2.7, 5.7; $P < 0.001$), with 46% of males and 35% of females maintaining clinically significant losses (>5%) after 12 months. pSMAs were calculated to be four times more cost effective and up to seven times more time effective than 1:1 consultations for weight loss.

Discussion

pSMAs are a popular, cost and time-effective adjunct to standard clinical management.

CHRONIC DISEASES are currently the leading cause of ill health and death in Australia.¹ Various approaches have been taken to manage these conditions,² most with limited success. However, these approaches still generally involve a conventional 1:1 model of medical consultation or, alternatively, group education without medical consultation. The need to increase access to services and improve outcomes makes the testing of alternative treatment modalities a priority.

Shared medical appointments (SMAs) are 'individual medical consultations carried out sequentially in a group of patients with similar concerns, with all listening and contributing'.³ These have been tested extensively in many countries^{4–8} and have been adopted as a process in the emerging discipline of lifestyle medicine.⁹

Our work with SMAs in primary care centres^{6–8} has led us to extend this into a form of 'programmed' SMA (pSMA), which potentially provides a platform for a range of interventions for chronic diseases and conditions.

pSMAs have been defined as 'a sequence of SMAs in a semi-structured form providing discrete educational input relating to a specific topic'.¹⁰ These allow for a set number of SMAs run in a sequenced 'active learning' format coordinated by an SMA facilitator (practice nurse/allied health professional [AHP]) with generic training in conducting SMAs and specific training in the disease topic, but with a doctor providing individual sequential consultations during part of the session, with participation and input of the group.

The following represents a trial of pSMAs for weight management in primary health care (PHC) centres that include general practitioners (GPs) in NSW.¹¹ Weight management was chosen for testing as obesity is currently a health problem worldwide, and it is a chronic condition for which PHC centres need to become more engaged.¹² Preliminary work has also indicated the potential value of such an approach.¹³ The primary aim of the trial was to measure the acceptability of pSMAs for best use in primary care. Secondary aims were to measure weight loss over a 12-month period and to assess cost and time effectiveness of the process.

Method

Ethics approval for this study was granted by Southern Cross University Ethics Committee (Renewal Approval Numbers ECN-14-268 and ECN-18-181).

Setting

Medical centres

PHC centres in the South Eastern Primary Health Network of NSW were invited to participate through a call for expressions of interest. The aim was completion of six pSMA sessions over 14–16 weeks in each of four pSMA groups (two all-male groups and two all-female groups) with 6–12 participants per group. A selection questionnaire was used to select centres on the basis of our previous work with such centres.

Patients

Overweight and obese (body mass index [BMI] 24–44 kg/m²) patients were recruited through GP referrals and other centre staff on the basis of perceived need and expressed interest. The only exclusions were no English language capability and extreme obesity (BMI >45 kg/m²) with complications. There was no charge for attendance.

Facilitators

Nurses or AHP staff volunteered for training as facilitators for the program. Our goal was to obtain 1–2 facilitators per centre. Staff selection was vetted by the study investigators with PHC staff as having the required qualities for the role of an SMA facilitator as specified by Noffsinger.³

Doctors

Doctors were GPs with no specific background in SMAs or weight control. It was explained that while it may be beneficial for GPs to attend facilitator training, this was not compulsory, as the doctor in an SMA is required to conduct a standard medical consultation (with the possible exception of an intimate examination). It differs from a standard consultation as it is conducted in the presence, and with participation, of other patients. However, doctors were encouraged to familiarise themselves with up-to-date content material relating to

weight control on the facilitators' websites and in a text¹⁴ and guidelines provided.¹⁵

Procedure

Facilitator training

Selected facilitators underwent two days of face-to-face training. The first day of training focused on conducting SMAs in general (www.lifestylemedicine.org.au/shared-medical-appointments-workshop-for-practitioners-and-facilitators), and the second day focused on evidence-based weight management skills. Successful completion of training resulted in certification by the Australasian Society of Lifestyle Medicine (ASLM).

Program content

The program was designed as an 'active learning' initiative based on evidence-based weight loss information. The process of active learning involved all patients participating in the learning process rather than passively listening. It included the optional choice of meal replacements and possibility for triaging into additional psychological care, prescription medication and/or surgical intervention if required.¹⁶

Program structure

The program was conducted over six sessions, two weeks apart for the first two sessions, then monthly. The focus was on reducing 'volume' of energy input (where volume(i) = energy density × portion size × frequency of eating/drinking) and increasing 'volume' of energy expenditure (where volume(e) = intensity × duration × frequency of physical activity). Topics differed for males and females, although there was significant overlap. Each session lasted 90–120 minutes. For the first 20–40 minutes facilitators presented a pre-prepared audiovisual presentation (approximately 10 minutes in length) and held an open discussion on the topic for the session, before the doctor joined for one hour for individual sequential consultations. The facilitator spent 10–15 minutes at the end of the session summarising and answering questions. Participants were given a program handbook, weight loss goals and access to presentations and

online links. Each session was designed around three major actions, which were listed on a fridge magnet provided to patients after each session. As in standard SMAs, patients signed a confidentiality agreement and were advised in an introductory brief that confidential matters did not need to be raised, and that a standard 1:1 consultation could be used for this.

Measures

Participant and provider satisfaction were measured using a five-point Likert scale (1 = very negative; 5 = very positive).

Weight was taken by facilitators:

1. at baseline
2. after the 16–18-week intervention at the last SMA
3. from medical records from each centre (70%) after 12 months, within a two-month window (ie 11–13 months), or self-reports from follow-up telephone calls when these were not available.

Twelve-month weights were compared with beginning weights for participants who had not completed at least four sessions ('non-adherers') and whose weights were available from medical records after 12 months (42 of 54). Waist circumferences were also measured but are not reported here because of inconsistency in measures taken by facilitators and difficulty in assessing changes in women in particular. As well as Likert scales, limited semi-qualitative information was provided from semi-structured interviews with providers. Preliminary cost and time effectiveness estimates were made by comparing weight losses (per kilogram) with those from a comparative 1:1 study.¹⁷

Statistical analyses

Regression analyses with generalised estimating equation (GEE) adjustment for repeated observations of the same participants were used to estimate percentage changes in mean weight (kg) over the study period. For each GEE model, a Poisson distribution, log-link function and independent working correlation matrix were used. The dependent variable in each model was weight in kilograms. To conservatively

account for loss to follow-up, an additional analysis was performed estimating percentage change in weight from baseline to 12 months by substituting missing weight values with baseline observation carried forward (BOCF). Another analysis calculated the proportions of patients who achieved clinically important weight loss (>5%) from baseline to 12 months, both by using the observed data and more conservatively by assuming those with missing weight values at 12 months did not achieve clinically important weight loss.

Results

Centres, facilitators, doctors and patients

Seven PHC centres in the South Eastern Primary Health Network of NSW were chosen from 15 responders to a call for expressions of interest. A total of 216 overweight and obese patients (83 men and 133 women) aged 35–85 years were recruited for 25 groups (mean 8.7 patients per group). Twelve nurses or AHP staff volunteered for training as facilitators for the program. This included practice nurses (8), a diabetes educator (1), an exercise physiologist (1), a dietitian (1) and a psychologist (1). Seven GPs also participated.

Of 216 patients starting the program, 166 (77%), completed at least four of six sessions (hereafter called

‘adherers’). Weight measurements were obtained at 12 months for 89% (n = 192) of participants (92% [n = 153] of all adherers and 78% [n = 39] of non-adherers).

The mean age of both male and female adherers was 57 years. The mean age of those failing to complete four or more sessions was 59 for males (n = 19) and 48 for females (n = 31). Both adherers and non-adherers claimed to have been ‘overweight’ for a mean of 16 years. Socioeconomic status of most participants was estimated by providers as being predominantly low-to-middle income. Mean patient weight loss expectation for male adherers was 21% of current weight; for female adherers it was 25% of current weight. This was similar for non-adherers at 25%.

Tables 1–3 show patient satisfaction with the program and program components.

The participating doctor (used by all) was rated highly as part of the program, confirming the notion of an SMA (Table 3).

Providers

Likert scale averages for evaluation of the program (1 = poor; 5 = great) are shown for 19 providers (12 AHP facilitators and seven GPs) in Table 4.

Common statements from providers about what they liked most about running pSMAs in weight control included:

Good change to the normal day – helps build more rapport with patients.

Seeing patients learning off each other is great.

It’s good to have time to cover topics and not repeat oneself.

Common statements about what they liked least included:

Financial concerns [for doctors]. (For example, is payment enough for the groups?)

No-shows and time required chasing people up.

Too many and too high level resources for some. These need to be simplified.

Body weight changes

Overall, participants lost an average of 3.2% (95% confidence interval [CI]: 2.2, 4.3; P <0.001) of their baseline weight at 12 months (Table 5). Male adherers lost 4.3% (95% CI: 2.6, 6.1; P <0.001) and female adherers lost 4.2% (95% CI: 2.7, 5.7; P <0.001) of their baseline weight at 12 months (Table 5; Figures 1 and 2). There were no significant changes among non-adherers.

Results obtained using BOCF were similar to analyses of the observed

Table 1. Mean participant satisfaction, preferences and recommendation

	Response		
	Men (n = 65)	Women (n = 102)	Total (n = 167)
Satisfaction	Mean Likert score		
<i>How do you rate the program you have attended here?</i>	4.4	4.2	4.3
<i>How useful has the program been for you?</i>	4.3	4.2	4.2
<i>How did the program compare with other weight loss methods you have tried?</i>	4.2*	3.2	3.7
Preferences	Yes/Maybe (%)		
<i>Would you recommend this form of weight loss program to others?</i>	51 (78%)*	90 (88%)	142 (85%)
<i>Would you prefer this type of program for weight control over a 1:1 session with your doctor?</i>	42 (65%)	83 (81%)	122 (73%)

*Men had tried far fewer weight loss methods than women.

data, suggesting the results were not substantively affected by loss to follow-up.

Weight losses of >5% maintained from baseline are considered clinically significant.¹⁷ In the current study, 46% (95% CI: 33.1, 58.5) of male and 35% (95% CI: 25.5, 44.8) of female adherers achieved this (Table 6).

Cost analysis

Using pSMAs for six sessions (10 participants/session), and a mean loss of 4.5 kg among adherers as reported here, with doctors' time 1.5 hours/session, and assuming payment for the doctor of approximately \$40 per person [pp] (current Medicare Benefits Schedule [MBS] item 23), and nurse \$10 pp (ie a total of approximately \$50 pp/session), the costs would be \$300 pp or approximately

\$67/kg/pp. The time required for the doctor would be 14 mins/kg/pp.

Discussion

To the best of our knowledge, the current study is the first to report results of pSMAs for chronic diseases and conditions in primary healthcare; in this case, the factor examined was weight control. Our findings of positive participant and provider satisfaction, clinically significant weight loss outcomes in approximately 40% of adherers over 12 months and a significantly reduced cost and time effectiveness all support the process. An adherence rate of 77%, together with willingness to recommend the program to others and high ratings on aspects of the program, add to this conclusion.

There are three commonly quoted participant advantages of SMAs:²

- peer support and contribution of peers to the consultation
- extra time with a more relaxed and focused physician
- the contribution of AHPs to a consultation.

Almost three in four of those involved in this study said they 'maybe/would' prefer this type of consultation (for weight control) over a conventional 1:1 session with their doctor.

All 19 providers (facilitators and doctors) involved in the study also expressed satisfaction with the process, because:

- it reduces repetition to patients
- there is less pressure on the doctor
- the doctor is not focused on a computer screen (patient notes are recorded by the facilitator).

All providers agreed they would 'definitely' be interested in running pSMAs for other chronic diseases.

Overall mean percentage weight loss at 12 months was modest at around 3.2% (although approximately 4.3% among adherers). However, as clinically significant weight loss is regarded to be >5% of baseline weight,¹⁶ the proportions of adherers achieving this (46% of men and 35% of women) are of interest. Even with the most conservative analysis combining adherers and non-adherers, and assuming those lost to follow-up did not lose weight, 34% of men and 26% of women achieved clinically significant weight loss.

One US review has reported that a total of 11–26 (mean = 18) intensive visits with a doctor on a 1:1 basis over one year results in an average 4–7 kg (mean = 5.5 kg) weight loss.¹⁷ At 30 minutes per visit (nine hours total), using reimbursement rates in 2019 (ie approximately \$75 per 30 mins), this would cost approximately \$1350 pp, or \$245/kg/pp, and take 98 mins/kg/pp. Comparative costs shown here are \$300 pp or approximately \$67/kg/pp and 14 mins/kg/pp. This is approximately four times more cost effective (approximately 27% of the cost of the conventional approach pp) and seven times more time effective (completed in approximately 14% of the time pp). Similarly, when compared

Table 2. Mean program component satisfaction ratings of participants

Program component	Mean Likert score		
	Men (n = 65)	Women (n = 102)	Total (n = 167)
Having time for asking questions	4.5	4.4	4.4
Seeing the doctor more relaxed than usual	4.3	4.3	4.2
Having the doctor's/staff's full attention	4.3	4.5	4.3
Getting support from others in the group	4.2	4.3	4.1
Contribution of other health professionals	4.4	4.3	4.2
Hearing experiences of other patients	4.4	4.3	4.2
Getting information from others	4.4	4.4	4.4

Table 3. Mean program materials enjoyment ratings of participants (1 = did not enjoy; 5 = enjoyed very much)

Material	Used material n (%)		Mean Likert score for users	
	Men (n = 65)	Women (n = 102)	Men (n = 65)	Women (n = 102)
Participating doctor	65 (100%)	102 (100%)	4.2	4.2
Program workbook	28 (43%)	44 (43%)	3.3	3.9
Ongoing tips	37 (57%)	38 (37%)	3.8	3.9
Fridge magnets	40 (62%)	40 (39%)	3.4	3.5
Website	25 (38%)	30 (29%)	2.4	3.4
Self-tests	21 (32%)	29 (28%)	3.4	3.3

Table 4. Mean Likert scale scores for provider satisfaction with the program (1 = poor; 5 = great)

	Facilitators (n = 12)	General practitioners (n = 7)
How do you rate the notion of pSMAs for weight control?	4.5	4.3
How do you rate the notion of pSMAs for other chronic diseases?	4.9	4.6
Rate your interest in running other programs such as this for chronic disease.	5.0	5.0
Rate your interest in getting training and accreditation as a lifestyle medicine practitioner for presenting different types of chronic disease in a pSMA program like this, where the individual programs are written (in a standard format) by experts in each area.	5.0	5.0

pSMA, programmed shared medical appointments

Table 5. Percentage change in mean weight (kg) from baseline to 12 months with 95% confidence intervals by all participants, sex and adherer status

Group	Baseline		12 months observed data				12 months baseline observation carried forward [†]			
	n (%)	Mean weight (kg)	n (%)	Mean weight (kg)	% change from baseline (95% CI)	P value	n (%)	Mean weight (kg)	% change from baseline (95% CI)	P value
All participants	216 (100%)	105.4	192 (89%)	102.2	-3.2 (-4.3, -2.2)*	<0.001	216 (100%)	101.7	-3.5 (-4.3, -2.7)	<0.001
Sex										
Men	83 (100%)	114.1	78 (94%)	110.6	-3.1 (-4.6, -1.5)	<0.001	83 (100%)	110.1	-3.6 (-4.8, -2.3)	<0.001
Women	133 (100%)	99.9	114 (86%)	96.5	-3.4 (-4.8, -2.0)	<0.001	133 (100%)	96.4	-3.4 (-4.4, -2.5)	<0.001
Completed ≥4 sessions										
Adherers	166 (100%)	105.2	153 (92%)	100.7	-4.3 (-5.4, -3.1)*	<0.001	166 (100%)	100.4	-4.5 (-5.4, -3.7)	<0.001
Non-adherers	50 (100%)	105.8	39 (78%)	108.1	0.4 (-1.7, 2.6)*	0.684	50 (100%)	105.8	0.0 (-1.0, 1.0)	0.974
Sex and completion of ≥4 sessions										
Male adherers	64 (100%)	113.5	59 (92%)	108.5	-4.3 (-6.1, -2.6)	<0.001	64 (100%)	108.0	-4.8 (-6.3, -3.4)	<0.001
Female adherers	102 (100%)	100.1	94 (92%)	95.9	-4.2 (-5.7, -2.7)	<0.001	102 (100%)	95.7	-4.3 (-5.4, -3.3)	<0.001
Male non-adherers	19 (100%)	116.4	19 (100%)	117.0	0.6 (-0.7, 1.9)	0.382	19 (100%)	117.0	0.6 (-0.7, 1.9)	0.382
Female non-adherers	31 (100%)	99.3	20 (65%)	99.6	0.3 (-3.6, 4.4)	0.875	31 (100%)	98.8	-0.4 (-1.9, 1.0)	0.554

*Adjusted for sex

[†]For each participant with missing weight measurements, their baseline value was used to fill in missing values at 12 months.

P values are for test of whether percentage change in weight is significantly different

CI, confidence interval

with meta-analyses of commercial weight loss programs, the outcomes shown here over one year are up to 10 times more cost effective,¹⁸ and retention levels are greater (77%, compared with <70%).¹⁹ It is acknowledged, however, that 1:1 and

SMA visits may be totally different when comparing the quality of care. Therefore, further work will be necessary to tease out these cost comparisons. These findings must also be considered preliminary to reflect any dynamic changes presenting to

practice following the SMA process. More specific economic research is required here.

The current study was funded by a Primary Health Network (PHN) grant and did not rely on the MBS. The case for pSMAs may become more compelling in

a capitation-funded system such as New Zealand, the National Health Service (NHS) in the UK, or the Health Care Homes model currently being trialled in Australia, where bundled payments provide greater certainty of funding as well as greater scope for multidisciplinary

involvement. Cost effectiveness may also come from a lower burden of disease and decreased impact of healthcare costs in the future, which can only be determined by a long-term economic study. The SMA process covers the three main components of quality in healthcare ('effective', 'safe'

and 'client-centred') as identified in a recent *New England Journal of Medicine* Catalyst survey.²⁰

The primary limitation of this exploratory study was the single-arm design involving patients who may be more motivated to lose weight than typical overweight and obese patients. Similarly, while adherers lost considerably more weight than non-adherers, these differences in weight loss may be at least partly due to adherers being inherently more motivated than non-adherers. Despite these limitations, the degree of weight loss observed in this exploratory study is encouraging and warrants further investigation in a clinical trial randomising patients to SMA or conventional 1:1 clinical care. When designing such a trial, it is worth noting that post-hoc evaluations with providers and some participants suggest that outcomes might be improved further by changes to the program such as: fortnightly, rather than monthly, sessions; more specific up-skilling of doctors to ensure greater knowledge in evidence-based aspects of weight control; and better targeting of participants.

Chronic diseases in primary healthcare are unlikely to be managed fully through a single treatment modality. A procedure such as that tested here (pSMAs) could help a significant proportion (perhaps the majority) of patients with chronic diseases or ailments at the non-critical end of an affected population. If achieved at scale, pSMAs could enhance the contribution of PHC centres to population health outcomes.

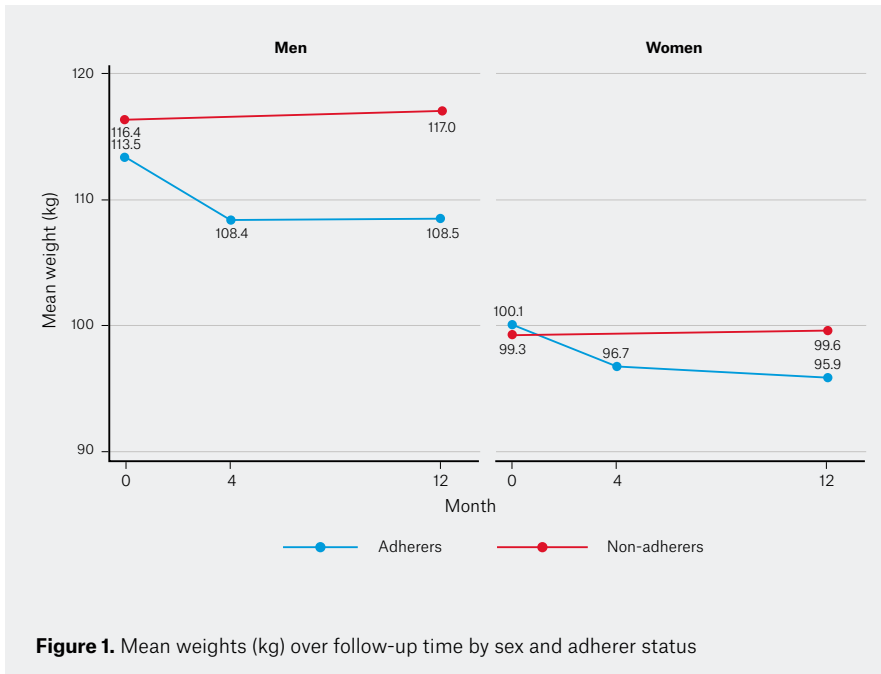


Figure 1. Mean weights (kg) over follow-up time by sex and adherer status

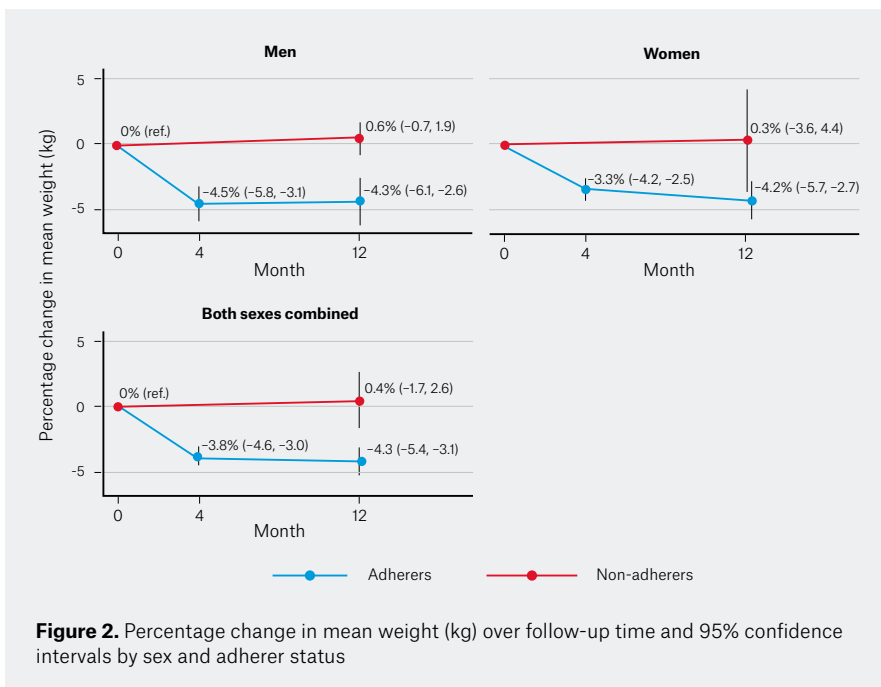


Figure 2. Percentage change in mean weight (kg) over follow-up time and 95% confidence intervals by sex and adherer status

Implications for general practice

- pSMAs are a series of SMAs that provide an alternative to conventional 1:1 clinical care.
- This exploratory study of pSMAs in weight loss shows they are highly popular among both participants and providers, with significant clinical outcomes after 12 months and cost and time efficiencies over conventional care.
- pSMAs provide an effective adjunct approach to chronic disease management in primary care.

Authors

Garry Egger AM, MPH, PhD, FASLM, Board Member, Australasian Society of Lifestyle Medicine (ASLM), Vic; Director, Centre for Health Promotion and Research, NSW; Adjunct Professor, Southern Cross University, NSW. egggergj@ozemail.com.au

John Stevens PhD, RN, Adjunct Associate Professor, School of Health Sciences, Southern Cross University, NSW; Board Member, Australasian Society of Lifestyle Medicine (ASLM), Vic; Director, Lifestyle Medicine Programs, NSW

Nerida Volker BSc, Grad Dip Diet, PGrad Dip Hlth Prom, PhD, Service Development and Performance Manager, South Eastern NSW PHN (Coordinare), NSW
Sam Egger BSc, MBIostat, Biostatistician, Cancer Research Division, Cancer Council, NSW

SENSW Weight Management Trial Group

Includes the following SENSW Australia medical practices: Bombala St., Surgery, Cooma; Cringila Medical Centre; Milton Medical Centre; North Nowra Medical Centre; OCHRE Medical Centre, Wollongong; Russel Vale Family Medical and Acupuncture Practice, Wollongong; Sapphire Medical Clinic, Merimbula
Competing interests: None.

Funding: This study was funded by Coordinare, South East NSW, Primary Health Network.

Provenance and peer review: Not commissioned, externally peer reviewed.

Acknowledgements

We wish to acknowledge Coordinare, South East NSW, Primary Health Network, for funding this project; Energize Health Club and Martin Dunkerly for website development and contributions to the study.

References

1. Australian Institute of Health and Welfare. Australia's health 2018: In brief. Cat. no. AUS 222. Canberra: AIHW, 2018.
2. National Health Performance Authority. Healthy Communities: GP care for patients with chronic conditions in 2009–2013. Sydney: NHPA, 2014.
3. Noffsinger EB. The ABCs of group visits: An implementation manual for your practice. New York: Springer, 2013.
4. Edelman D, McDuffie JR, Oddone E, Gierisch JM, Nagi A, Williams JW Jr. Shared medical appointments for chronic medical conditions: A systematic review. Durham, NC: Evidence-based Synthesis Program (ESP) Center Durham Veterans Affairs Healthcare System, 2012.
5. Housden L, Wong ST, Dawes M. Effectiveness of group medical visits for improving diabetes care: A systematic review and meta-analysis. CMAJ 2013;185(13):E635–44. doi: 10.1503/cmaj.130053.
6. Egger G, Binns A, Cole M-A, et al. Shared medical appointments – An adjunct for chronic disease management in Australia? Aust Fam Physician 2014;43(3):151–54.
7. Stevens JA, Cole M-A, Binns A, Dixon J, Egger G. A user assessment of the potential for shared medical appointments in Australia. Aust Fam Physician 2014;43(11):804–07.
8. Stevens JA, Dixon J, Binns A, Morgan B, Richardson J, Egger G. Shared medical appointments for Aboriginal and Torres Strait Islander men. Aust Fam Physician 2016;45(6):425–29.
9. Sayburn A. Lifestyle medicine: A new medical speciality? BMJ 2018;363(1):k4442. doi: 10.1136/bmj.k4442.
10. Egger G, Stevens J, Ganora C, Morgan B. Programmed shared medical appointments: A novel procedure for chronic disease management. Aust J Gen Practice 2018;47(1-2):71–75.
11. Egger G, Stevens J, Ganora C. Establishing proof of concept for clinical processes in lifestyle medicine. Am J Lifestyle Med 2018. doi: 10.1177/1559827617750896.
12. World Health Organization. Obesity and overweight. Geneva: WHO, 2018. Available at www.who.int/news-room/fact-sheets/detail/obesity-and-overweight [Accessed 22 August 2019].
13. Tunay M, Kurdak H, Ozcan S, Ozdemir C, Özer ZY. Family physician-led group visits for lifestyle modification in women with weight problems: A pilot intervention and follow-up study. Obes Facts 2018;11:1–14. doi: 10.1159/000486133.
14. Egger G, Binns A. The expert's weight loss guide. Sydney, NSW: Allen and Unwin, 2001.
15. National Health and Medical Research Council. Clinical practice guidelines for the management of overweight and obesity in adults, adolescents and children in Australia. Melbourne: National Health and Medical Research Council, 2013.
16. National Institute for Health and Care Excellence. Weight management: Lifestyle services for overweight or obese adults. London: NICE, 2014.
17. Ard J. Obesity in the US: What is the best role for primary care? BMJ 2015;350:g7846. doi: 10.1136/bmj.g7846.
18. Finkelstein EA, Kruger E. Meta- and cost-effectiveness analysis of commercial weight loss strategies. Obesity (Silver Spring) 2014;22(9):1942–51. doi: 10.1002/oby.20824.

Table 6. Proportions of patients who achieved clinically important weight loss (>5%) from baseline to 12 months

Group	Achieved clinical weight loss at 12 months			
	Observed data		All participants*	
	Proportion	% (95% CI)	Proportion	% (95% CI)
All participants	63/192	32.8% (26.2, 39.5)	63/216	29.2% (23.1, 35.2)
Sex				
Men	28/78	35.9% (25.3, 46.5)	28/83	33.7% (23.6, 43.9)
Women	35/114	30.7% (22.2, 39.2)	35/133	26.3% (18.8, 33.8)
Completed ≥4 sessions				
Adherers	60/153	39.2% (31.5, 47.0)	60/166	36.1% (28.8, 43.5)
Non-adherers	3/39	7.7% (0.0, 16.1)	3/50	6.0% (0.0, 12.6)
Sex and completion of ≥4 sessions				
Male adherers	27/59	45.8% (33.1, 58.5)	27/64	42.2% (30.1, 54.3)
Female adherers	33/94	35.1% (25.5, 44.8)	33/102	32.4% (23.3, 41.4)
Male non-adherers	1/19	5.3% (0.0, 15.3)	1/19	5.3% (0.0, 15.3)
Female non-adherers	2/20	10.0% (0.0, 23.1)	2/31	6.5% (0.0, 15.1)

*Patients who had missing weight values at 12 months were treated as having not achieved clinical weight loss.
CI, confidence interval

19. McEvedy SM, Sullivan-Mort G, McLean SA, Pascoe MC, Paxton SJ. Ineffectiveness of commercial weight-loss programs for achieving modest but meaningful weight loss: Systematic review and meta-analysis. *J Health Psych* 2017;22(12):1614–27. doi: 10.1177/1359105317705983.
20. NEJM Catalyst. Buzz report: Addressing the problems of quality measurement. Waltham, MA: NEJM Group, 2018.

correspondence ajgp@racgp.org.au