A within-trial cost-effectiveness analysis of primary care referral to a commercial provider for weight loss treatment, relative to standard care—an international randomised controlled trial

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INTRODUCTION

The prevalence of overweight and obesity is placing a substantial burden on health-care resources, even in developed countries.1 Overweight and obesity accounts for 44% of the global burden of type 2 diabetes mellitus, 23% of ischaemic heart disease and 7–41% of certain cancers.2 Therefore obesity management programmes that are both efficacious and cost effective are needed. Policy makers are increasingly seeking evidence of the cost effectiveness of interventions. It is important to know whether it is more cost effective to support and fund programmes already in place or subsidise others (including extant commercial weight loss programmes).

A partnership between primary-care providers and commercial organisations may be a practical approach, whereby participants can benefit from early lifestyle intervention for weight management. Observational data3,4 show that this approach has the potential to deliver weight management programmes at the necessary scale in a community setting and at potentially relatively low cost. Our recent 12-month randomised controlled trial (RCT) involving three countries (Australia, the United Kingdom and Germany) showed referral to a commercial weight-loss community intervention programme (Weight Watchers—commercial provider (CP)) produced greater weight loss compared with standard care (SC).5 Similar efficacy of this CP has been demonstrated in other RCTs.6,7 However, the cost effectiveness of CPs over SC has not been estimated. We calculated this using data from the above trial.8 Previous estimates of the cost of the CP have been done, but were small scale and used limited data.8

Our aim was to evaluate the cost effectiveness of a CP compared with conventional SC for both weight loss and quality of life (QOL). A societal perspective was also adopted as we have previously reported that those attending the CP had more frequent visits,5 which may have contributed to the success of the CP.

BACKGROUND: Due to the high prevalence of overweight and obesity there is a need to identify cost-effective approaches for weight loss in primary care and community settings.

OBJECTIVE: We evaluated the cost effectiveness of two weight loss programmes of 1-year duration, either standard care (SC) as defined by national guidelines, or a commercial provider (Weight Watchers) (CP).

DESIGN: This analysis was based on a randomised controlled trial of 772 adults (87% female; age 47.4 ± 12.9 years; body mass index 31.4 ± 2.6 kg m−2) recruited by health professionals in primary care in Australia, United Kingdom and Germany. Both a health sector and societal perspective were adopted to calculate the cost per kilogram of weight loss and the ICER, expressed as the cost per quality adjusted life year (QALY).

RESULTS: The cost per kilogram of weight loss was USD122, 90 and 180 for the CP in Australia, the United Kingdom and Germany, respectively. For SC the cost was USD138, 151 and 133, respectively. From a health-sector perspective, the ICER for the CP relative to SC was USD18 266, 12 100 and 40 933 for Australia, the United Kingdom and Germany, respectively. Corresponding societal ICER figures were USD31 663, 24 996 and 51 571.

CONCLUSION: The CP was a cost-effective approach from a health funder and societal perspective. Despite participants in the CP group attending two to three times more meetings than the SC group, the CP was still cost effective even including these added patient travel costs. This study indicates that it is cost effective for general practitioners (GPs) to refer overweight and obese patients to a CP, which may be better value than expending public funds on GP visits to manage this problem.

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METHODS

Clinical trial
This cost-effectiveness analysis used data from an RCT whereby overweight and obese adults were randomised to receive 12-month access to a CP or SC by a primary-care professional in Australia, the United Kingdom and Germany. Participants were recruited by their general practitioners (GPs) and randomised to one of the two groups. A full list of inclusion and exclusion criteria, as well as a description of the two intervention groups are in the report of the primary findings from the study. All participants were aged 18 years with a body mass index of 27–35 kg m⁻², and had at least one risk factor for obesity-related disease. Participants randomised to the CP group received vouchers to attend a weekly community CP meeting. Those randomised to SC received weight-loss advice delivered by a GP/primary care professional at their local medical practice. The frequency of these SC visits was at the discretion of the GP and the participant. The frequency of such visits was recorded, with GP visits only being counted for SC. GPs and primary-care professionals were provided with and encouraged to use relevant national clinical guidelines for weight management.

Health economic evaluation
A health-care funder and societal perspective was adopted for the cost-effectiveness evaluation and included direct health costs to the government or patient arising from costs of intervention delivery and costs to patients for participation.

Costing data
Commercial programme. The costing for the CP group was based on a monthly payment plan, which included unlimited access to meetings and online electronic web tools. The cost of the referral visit to the CP was also included in the costing.

Standard care. For SC, the cost applied was that of a consultation lasting 20 minutes or less with a GP. In the United Kingdom, all consultations provided in the SC group were by a nurse.

Valuation of costs
The cost of the CP for each country was sourced directly from Weight Watchers International, Inc. (New York, NY, USA). Unit costs for all other intervention and conventional resources were obtained from the relevant governments/health-care authorities (Table 1). Costs were estimated for patient travel to attend either CP or SC consultations. This was based on the assumption that patients travelled within a 10-km radius to their intervention. The number of participant visits to either the CP or SC was recorded throughout the 1-year study.

Opportunity costs of employment were not considered because participants could attend their intervention outside working hours, during their lunch break, or on weekends. Childcare costs were not considered as children of any age are welcome at the CP meetings and can accompany their parent to an SC visit.

Australia. In Australia, medical care is priced by the Medicare Benefits Schedule (MBS) and Pharmaceutical Benefits Scheme (PBS). Professional attendances were considered as ‘Group A1—general practitioner attendances to which no other item applies (level B)’—MBS item 23.

United Kingdom. The cost of a GP referral and nurse consultation were calculated according to the Personal Social Services Research Unit (PSSRU).

Germany. The costs of a primary-care visit were calculated from 2007–2008 data (later data not available) and indexed using the geometric mean of the increase of GP costs 2007–2008, according to the Working Group ‘Methods in Health Economic Evaluation’.

Outcomes measured and statistical analyses
Weight loss and change in QOL were measured. While in Germany, bodyweight (in light clothes without shoes) was measured in GP practices with standard scales, in the other countries measurements were with a Tanita BC-418 segmental body composition analyser (Tanita Corporation of America, Arlington Heights, IL, USA). Participants were weighed on six occasions over 12 months (baseline, month 2, 4, 6, 9 and 12) and completed the Impact of Weight on Quality of Life Questionnaire-Lite (IWQOL-Lite) on three occasions (baseline, months 6 and 12).

All participants who completed a baseline assessment were included in an intention to treat analysis using last-observation carried forward. A utility score was derived using the algorithm described by Brazier et al.

Outcomes were analysed by linear regression with fixed effects for continuous normal data; intervention group (CP vs SC) and baseline measurement were used as the fixed effects.

Cost-effectiveness analysis
The cost-effectiveness analysis was performed for each country separately, incorporating differences of the cost of the commercial intervention, medical consultation fee structure, salary structures and other costing data.

An additional cost-effectiveness ratio (ICER) was calculated giving the net costs of CP relative to SC. The ICER represents the additional expenditure required to generate an additional unit of benefit, and was expressed as the cost per QALY. QALYs were calculated from the IWQOL-Lite result. A preference based single index was estimated for each country to provide country specific ICERs. This formula was used to calculate cost per QALY.

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\text{ICER} = \frac{\text{Cost of CP} - \text{Cost of SC}}{\text{Difference in QALYs}}
\]

An analysis was also performed to calculate the cost per kilogram of weight loss.

Sensitivity analysis
One-way sensitivity analyses were performed for the Australian site to include medication costs. An average costing for each subject was calculated in 3-month time periods so an annual cost could be estimated. Drug costs were obtained from the 2011 PBS pricing index—dispensed price for maximum quantity and units were based on the subject's self-reported data. All medications and dosages prescribed were collected at each of the individual subject's visits. The difference in total medication costs between CP and SC, plus the costs of implementing the intervention, divided by the difference in QALYs, gave an additional cost-effectiveness ratio.

A further sensitivity analysis was performed including referral to allied health professionals in the SC group. An assumption was made that a proportion of GPs in the SC group referred their patients to a dietitian and/or psychologist for specific advice. A 1.1% referral rate for six consultations over a 1-year time period was applied, which was the probability of referral to an allied health professional during a GP visit in Australia in 2009–2010.

Scenario analysis
As the CP costs sourced in Table 1 reflect commercial pricing decisions and are financial costs, the programme costs were re-evaluated according to opportunity costs. The CP is identical across countries and the cost to deliver the intervention is similar. The cost-effectiveness consequences of reducing programme costs in all countries to the equivalent of the Weight Watchers NHS referral scheme (GBP 45 for 12 sessions) was examined. This is based on an attendance of 36 CP sessions per year (GBP 135–12 session cost multiplied by 3). The Weight Watchers NHS referral scheme was used as it is an existing system.

RESULTS

Clinical trial results
Baseline characteristics. Baseline characteristics of participants have been reported previously. The mean (s.d.) age of subjects was 47.4 (12.9) years, the mean body mass index was 31.4 (2.6) kg m⁻², and 87% were females; 65% had type 2 diabetes.

Weight loss. Both treatment groups lost weight but mean 12-month weight loss was significantly greater for CP than SC in all three countries (CP: -5.1 ± 0.3 kg vs SC: -2.3 ± 0.2 kg; P < 0.0001). Table 2 shows the mean weight loss per participant and country.
Quality of life. Seven hundred seventy-two participants commenced the study. However, only 744 participants completed an IWQOL-Lite questionnaire at the initial baseline visit. Of this latter group, 12 participants were excluded owing to incomplete questionnaires. A further 73 participants were excluded owing to missing data when the algorithm was applied for conversion to utility scores, leaving 659 participants who could be included in the intention to treat analysis. Four hundred forty-four participants completed the 12-month study. There was a significantly greater change in utility for CP than SC in Australia and the United Kingdom but not in Germany. A 0.021 and 0.015 improvement in utility score for the CP relative to SC was found in Australia and the United Kingdom, respectively, and a 0.009 improvement in utility score for the CP relative to SC was evident in Germany. When analysing completers only, there was a significantly greater change in utility for the CP than SC in Australia and Germany but not in the United Kingdom (owing to the small sample size) (Table 3). From a pooled analysis of all three countries the greater change in utility for CP than SC remained significant (results not shown).

Intervention costs

**Australia.** The average number of visits for each participant attending the CP or their GP was 33.0 and 10.7, respectively. Unit costs were summed for each resource measurement to obtain a total cost for each intervention (Table 1). The annual cost per patient was AUD 754 and AUD 373 for CP and SC, respectively. When including patient travel, the annual cost was AUD 1170 and AUD 508 for the respective programmes.

**United Kingdom.** The average number of visits for each patient attending the CP or their GP was 36.4 and 13.3, respectively. The annual cost per patient was GBP 269 and GBP 160 for CP and SC, respectively.
respectively. Including patient travel, costs were GBP 451 and GBP 226 for the respective programmes.

Germany. The average number of visits for each patient attending the CP or their GP was 23.1 and 11.3, respectively. The annual cost per patient was EUR 487 and EUR 217 for CP and SC, respectively. Including patient travel, costs were EUR 626 and EUR 284 for the respective programmes.

Cost effectiveness

Australia. The cost per patient increased during the 12-month study period for the CP group (Figure 1). Medication costs remained the same for the SC group. While not reaching statistical significance, the average medication cost per patient was EUR 28,826 (Table 4).

United Kingdom. The cost per patient was GBP 94 for the CP and SC, respectively. The ICER for the CP relative to SC was GBP 7378 (Table 4).

Germany. The cost per patient was EUR 127 and EUR 94 for the CP and SC, respectively. The ICER for the CP relative to SC was EUR 28,826 (Table 4).

Sensitivity analysis

While not reaching statistical significance, the average medication cost per patient increased during the 12-month study period for the SC group (Figure 1). Medication costs remained the same for the CP group. The mean number of prescriptions was 1.9 per patient. These were for hypertension, cardiovascular disease, dyslipidemia, diabetes mellitus, thyroid disease, reflux, arthritis and musculoskeletal disease, and other conditions. No outliers who may have had the capacity to overwhelm any trends were evident. The highest costs were associated with proton pump inhibitors, cardiovascular disease and dyslipidemia.

When medications were included, the annual cost per patient was AUD 1200 and AUD 874 for the CP and SC, respectively. With patient travel, programme costs were AUD 1616 and AUD 1009 for respective interventions. Including medication costs resulted in a decrease in the ICER for the CP relative to SC to AUD 15,522.

Including the costs associated with referral to a health professional decreased the ICER to AUD 17,901, GBP 7117, and EUR 28,403.

Scenario analysis

If programme costs in Australia and Germany are equated to the UK Weight Watchers NHS referral scheme, the ICER for the CP relative to SC becomes dominant (more health benefit at a lower cost) in both countries. In the United Kingdom, the ICER is lowered to GBP 655. The cost per kilogram of weight loss for the CP becomes AUD 41, GBP 35 and EUR 46.

DISCUSSION

This analysis shows the CP is a cost-effective option, over 1 year, for weight management in persons within a body mass index range of 27–35 kg m

This analysis may be an effective programme for primary-care providers to refer patients. The efficacy of the CP may be in part attributable to the shared care approach as the subject was initially referred by their GP.

In Australia and the United Kingdom, the cost per kilogram of weight loss was lower for the CP than SC. In Germany, the cost per kilogram of weight loss was lower for SC. It is difficult to compare these data with other weight loss studies because of different costing methodologies used. Previous lifestyle intervention studies did not include indirect costs such as patient travel and vary in terms of duration of outcomes, reporting costs from USD 61.25 to USD 133 per kilogram of weight loss, and USD 6.4027 to USD 48 per pound lost.28 In contrast there is data of a programme with only AUD 7.30 per kilogram of weight loss.29 When we adjusted for the CP financial costs based on commercial pricing decisions, and costed according to economic prices, the cost per

Table 2. Mean weight loss (kg) between baseline and month 12 per participant, treatment group and country (using LOCF, and completers only)

|          | CP | s.d. | N   | SC | s.d. | N   | Coefficient<sup>b</sup> | P      |
|----------|----|------|-----|----|------|-----|--------------------------|--------|
| Australia| LOCF | -6.21 | 6.53 | 120 | -2.72 | 4.93 | 123                      | 3.49   |
|          | CC<sup>a</sup> | -7.75 | 6.48 | 78  | -4.20 | 5.11 | 63                       | 3.55   |
| United Kingdom | LOCF | -4.91 | 5.10 | 91  | -1.73 | 3.13 | 87                       | 3.18   |
|          | CC<sup>a</sup> | -7.52 | 5.28 | 27  | -4.46 | 3.97 | 17                       | 3.06   |
| Germany  | LOCF | -3.84 | 6.58 | 107 | -2.31 | 4.35 | 131                      | 1.53   |
|          | CC<sup>a</sup> | -5.40 | 7.01 | 71  | -2.71 | 5.00 | 88                       | 2.69   |

Abbreviations: CP, commercial provider; LOCF, last-observation carried forward; SC, standard care. <sup>a</sup>Complete case. <sup>b</sup>Coefficient is the change in weight (kg) for CP compared with SC.
kilogram of weight loss was lower for the CP compared with SC in all three countries. As the cost per kilogram of weight loss is not commonly used by decision makers, a cost per QALY gained was also calculated. As an example of health resources used by this demographic group, medication usage was estimated for the Australian sample. When medications and their usage were included, the cost per QALY was lowered. Regardless of this, and using the commonly accepted threshold of $50,000 per QALY, the CP proved to be cost effective from both a health sector and societal perspective in Australia.

In the United Kingdom, the ICER level that the National Institute of Health and Clinical Excellence has adopted is £20–30,000 per QALY gained. Using this standard our results show the CP in the United Kingdom is a cost-effective programme from both health care and societal perspectives.

In Germany, obesity is not considered a disease in the health-care system. Therefore, obesity treatment is usually not undertaken in primary care. Comparing with cost-effective thresholds for pharmaceutical and surgical interventions in Europe (<£50,000), the CP is again cost effective when both perspectives were adopted.

The cost disparity of the CP and SC between countries had a large effect on the outcome, indicating that the results (cost per kilogram of weight loss and cost per QALY gained) are sensitive to the costs of the programme. In the United Kingdom, the cost to attend the CP was the lowest. When the CP costs were lowered to this value, the ICER for the CP relative to SC becomes dominant (more health benefit at a lower cost) in both Australia and Germany. Therefore the CP generated a better health outcome and cost less than SC. In the United Kingdom, this lowers the ICER to GBP 655.

It is possible that the SC group would have higher rates of referral to allied health due to greater contact with their GP. While attempting to compensate for such associated allied health costs in the SC group by applying a 1.1% referral rate, the only data available relate to the general population rather than specifically for an overweight and obese population, for which the referral rate is likely to be higher. Including costs associated with referral to allied health decreased the ICER in each country. As the CP subjects were not allowed to participate in other weight-loss programmes or seek other assistance for weight loss, these associated costs were not included for the CP group.

The cost of this early prevention approach targeting overweight and obese adults is low compared with the cost associated with the burden of type 2 diabetes. The average annual cost per person with type 2 diabetes in 2003 was AUD 5360 but prevention and intervention costs do vary considerably. As shown by previous lifestyle interventions, the degree of weight loss in the current study reduces the risk of type 2 diabetes in those at high risk. Despite weight regain being common, a so-called ‘metabolic memory effect’ has been shown in Diabetes Treatment and Prevention Outcome Studies and it could be anticipated that such metabolic benefit would occur with the weight loss achieved in this trial. Therefore, adopting cost-effective weight management programmes as demonstrated in this study, and an intervention that has the capacity to be large scale (CP), may be an effective way to reduce the prevalence of obesity associated diseases such as type 2 diabetes and their costs.

While a key strength of this analysis allows us to draw on the results of a RCT to perform a within-trial cost, it also acts as a limitation as the cost-effectiveness analysis is restricted to a 1-year time period. Importantly, the QALY metric is not all that applicable over a 1-year time horizon; however, it has been used in this study for demonstration purposes so that results can be compared with cost-effectiveness thresholds.

Another key strength of this study was the use of the societal perspective to take into consideration all intervention costs. This follows the recommended approach for cost-effectiveness analyses. However, as the available data from the RCT did not enable us to capture several cost-offsets including reduced rates of obesity-related disease, frequency of hospitalisations averted or

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**Table 3.** Change in utility scores (quality of life) between baseline and month 12 for CP compared with SC (using LOCF, and completers only)

| Analysis by country | Coefficient (to 3 decimal places) | 95% Confidence Interval | P | N |
|---------------------|----------------------------------|-------------------------|---|---|
| LOCF                |                                  |                         |   |   |
| Australia           | 0.021                            | 0.006 – 0.037           | 0.006 – 243 |   |
| United Kingdom      | 0.015                            | 0.001 – 0.028           | 0.033 – 178 |   |
| Germany             | 0.009                            | -0.006 – 0.024          | 0.222 – 238 |   |
| Completers Only     |                                  |                         |   |   |
| Australia           | 0.023                            | 0.003 – 0.044           | 0.027 – 141 |   |
| United Kingdom      | 0.020                            | -0.025 – 0.064          | 0.382 – 44  |   |
| Germany             | 0.021                            | 0.001 – 0.040           | 0.039 – 159 |   |

Abbreviations: CP, commercial provider; LOCF, last-observation carried forward; SC, standard care. *Coefficient is the change in utility score for CP compared with SC.

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**Table 4.** Costs per kilogram of weight loss per group and country and ICERs (cost per QALY) for the CP compared with SC from both a health-care funder and societal perspective

| Country                | Cost per kilogram of weight loss—including travel | Cost per kilogram of weight loss—including travel | Excluding travel | Including travel | 95% CI | ICER | 95% CI |
|------------------------|--------------------------------------------------|--------------------------------------------------|------------------|------------------|--------|------|--------|
| Australia (AUD)        | $121                                             | $137                                             | $188             | $187             | $18085 | $10239, $68841 | $31 349 | $17 723, $11 290 |
| (USD)                  | $122                                             | $138                                             | $190             | $189             | $18 266 | $10 342, $68 721 | $31 663 | $17 900, $11 443 |
| United Kingdom (GBP)   | £55                                              | £92                                              | £92              | £131             | £7 378  | £3851, £42 698 | £15 242 | £7 986, £89 286 |
| (USD)                  | £90                                              | £151                                             | £151             | £215             | £12 100 | £6316, £70 025 | £24 996 | £13 098, £146 429 |
| Germany (EUR)          | €127                                             | €94                                              | €163             | €127             | €28 826 | €10 890, Dominated | €36 318 | €13 561 Dominated |
| (USD)                  | €180                                             | €133                                             | €231             | €180             | €40 933 | €15 464, Dominated | €51 571 | €19 257, Dominated |

Abbreviations: AUD, Australian dollar; EUR, Euro; GBP, Great Britain pound; ICER, incremental cost-effectiveness ratio; SC, standard care. At the time of writing (early 2011), 1 AUD was 1.01 USD; 1 GBP was 1.64 USD; 1 EUR was 1.42 USD according to XE currency exchange (http://www.xe.com).
reduction in sick leave, the true cost effectiveness of the intervention programme may have been underestimated.

While it was assumed that no patient required time off from work to attend the CP or GP consultations, this information was not specifically captured. A large benefit of the CP and GP consultations is that they are available out of work hours, at lunchtimes and on weekends, minimising the opportunity costs of employment. An assumption may also be made that a percentage of patients are commenced on other weight loss assistance therapies by their GP in a SC real life setting, but such treatments were not included in our analysis. The limitations of the trial design meant that other costs such as food products endorsed by the CP and out of pocket expenses for the SC group (for example, allied health professional consultations) were ignored.

For the Australian sensitivity analysis medications were grouped together and it was not possible to determine whether there was a change in costs relating to obesity or disease-specific drugs. However, as reported by the Counterweight Programme in the United Kingdom, the range of prescribing areas being affected by obesity was greater than expected. An increased prescription rate was found for categories such as gastrointestinal, infections, skin conditions, antihistamines, hypnotics, and drugs used in the treatment of nausea and vertigo. Most clinicians would not typically associate these with obesity. Therefore, there is a wide range of clinical conditions for which obesity is a contributory factor.

In Australia, despite the GP-treated (SC) group showing an increase in median medication prescription over the 12-month period, it did not reach significance. This may reflect better control of risk factors by GPs as SC patients were required to see their primary-care professional on a more regular basis. However, despite this increase in prescriptions, they had no better control of risk factors for obesity-related conditions at the end of the 12-month study. Patients in the CP group had significantly greater total to HDL cholesterol ratio than in the SC group and serum insulin was lower, and weaker evidence existed of improvements in glucose, and HDL and LDL cholesterol. Small reductions in blood pressure were noted for both treatment groups. All these improvements may be related to, or influenced by, degree of weight loss. As the patients were overweight or obese with one or more additional risk factors for obesity-related disease, a slightly higher average number of medication scripts per person may be expected, and consequently a higher mean medication cost per patient. Results from a random Australian community sample of 3015 respondents with a mean age of 45.3 years found that 1411 (46.8%) were taking one or more prescribed medications.40 In this trial, it was 1.9 prescribed medications per patient.

Missing data were dealt with from an intention to treat perspective using last-observation carried forward. Despite a mixed model or multiple imputation approach now being commonly used to deal with missing data, the last-observation carried forward method was adopted as it was more conservative than the mixed model results previously reported in an interim analysis for weight loss of this RCT. This would imply a higher cost per kilogram of weight loss for the CP relative to SC.

The models by which IWQOL-Lite values are transformed into utilities has its weaknesses, and such a mapping exercise is second preference compared with the direct use of the EuroQol Europe or the AQLQ Australia, for example. But, the IWQOL-Lite is an obesity-specific measure of QOL sensitive to obesity-related QOL effects, and it performs well in terms of conventional psychometric criteria in obese populations. Despite the concerns with this mapping approach, it is possible to produce a robust model for predicting SF-6D index values from the IWQOL-Lite.

It has been argued that there is too much reliance on health professionals such as GPs to treat overweight and obesity. This is in part because GPs do not have additional or alternative resources to manage this issue beyond a standard GP consultation. The ability to be able to refer to cost-effective commercial programmes may assist. However, due to the out of pocket cost it may still be beyond the financial reach of a substantial proportion of the population, particularly those who need it most.68 Thus, governments could consider funding cost-effective commercial programmes in preference to GP visits for managing overweight and obesity. In Australia, for example, the addition of items to Medicare and private health insurance, providing financial incentives to better treatment, could help support individuals in changing their lifestyle behaviours.

This study provides data from three different countries and consistent results. The cost per kilogram of weight loss was lower for the CP versus SC in Australia and the United Kingdom. When adjusting for the CP financial costs based on commercial pricing decisions, and costing according to economic prices, the cost per kilogram of weight loss was lower for the CP compared with SC in all three countries. The CP is cost effective when assessed by the commonly accepted threshold of a cost <$50 000 per QALY. Importantly, despite participants in the CP group attending on average three meetings per month in the United Kingdom and Australia, and two meetings per month in Germany, compared with only one appointment per month for SC group, the CP remained cost effective when including these added patient travel costs.

CONFLICT OF INTEREST

NRF and IDC have received research grants for other clinical trials funded by Sanofi-Aventis, Allergan, Eli Lilly and Novo Nordisk. IDC was a board member for the SCOUT trial, and currently for the EXSCEL trial, and has received payment for lectures from iNovo Pharmaceuticals, Eisai Pharmaceuticals, Pfizer Australia and Servier Laboratories (Australia). SAJ has received research grants for other clinical trials from Sanofi-Aventis and Coca Cola, and is a member of the Tanita Medical Advisory Board and receives a fee for nutrition articles and lectures for Rosemary Conley Enterprises. HH is on the Advisory Board for Weight Watchers International and has received payment for lectures from Sara Lee, Novartis, Sanofi-Aventis and Bristol-Myers Squibb.

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AUTHOR CONTRIBUTIONS

The author’s responsibilities were as follows: NRF: acquisition of data, analysis and drafting of the paper. ADO, RS: analysis of data. CH, SBW, RH: health economic contribution for German specific data and critical review of the paper; ALA: health economic contribution for UK specific data and critical review of the paper. SAJ, HH: study design and conception, obtained funding and critical review of the paper. IDC: study design and conception, obtained funding and writing the paper. All authors read and approved the final paper.

TRIAL REGISTRATION

This study is registered with the International Standard Register of Clinical Trials, ISRCTN 85485463.

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