


# Providing food to patients in primary care to induce weight loss: a systematic literature review

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## ABSTRACT

**Purpose** Primary care is the ideal setting to promote weight management, warranting innovative ways to support patients. This systematic review aimed to determine whether providing food to patients in primary care can help to reduce body weight.

**Methods** Four databases were searched for studies that aimed to elicit weight loss by directly providing foodstuffs and/or supplements to patients in primary care settings. Interventions with adults of any gender or race were included. Interventions that involved other components such as exercise classes or education sessions were excluded. The methodological quality of each study was appraised using the Effective Public Health Practice Project Quality Assessment Tool for Quantitative Studies.

**Results** Four heterogeneous studies met the eligibility criteria, representing 476 adults. Two studies used meal-replacement products but differed in length and intensity, another study provided green tea and vitamin E supplementation, and the final study provided vouchers for use at a farmers' market hosted at a primary care clinic. Interventions ranged in length from 4 to 13 weeks. Three of the four studies observed weight loss in some form and all studies observed at least one other improvement in a health outcome such as waist circumference, blood pressure or fasting insulin levels.

**Conclusions** A small yet notable body of literature supports the concept of providing food to patients in primary care settings to support weight loss. Further, high-quality research is needed on the efficacy and cost-effectiveness of this approach to ultimately inform policy initiatives for primary care.

## INTRODUCTION

Diet influences the aetiology of 7 of the 10 most globally prevalent chronic conditions.<sup>1</sup> The avoidable negative impact of poor diet on excess body weight is also well recognised.<sup>2</sup> The WHO recommends all countries facilitate healthy eating through primary care settings, stating it is one of the 'best buys' for improving the health of societies and enabling people to live and age well.<sup>1</sup> Patients highly value discussions about diet during consultations with primary health professionals,<sup>3–7</sup> however, many health professions report significant

## What this paper adds

- ▶ Primary care interventions that involve food provision may support weight management in patients.
- ▶ Trialled interventions are diverse in length, intensity and approach.
- ▶ There is opportunity to further investigate the cost-effectiveness providing food to patients in primary care to support weight loss.

barriers to providing nutrition care including insufficient nutrition knowledge, low nutrition self-efficacy and competing priorities in consultations.<sup>8–10</sup> Consequently, only one-third of patients remember ever discussing nutrition during consultations,<sup>9</sup> warranting innovative strategies that enable and support patients to eat well and manage weight.

Novel approaches to supporting healthy eating in primary care have been trialled in many countries. In the USA, telephone counselling and mailed support material have shown a small positive effect on diet but only moderate sustainability.<sup>11</sup> In the UK, a primary care referral to a 12-month behavioural weight management programme has shown both effectiveness and cost-effectiveness in reducing weight in obese patients, but is time-intensive for patients.<sup>12</sup> In South America, group nutrition education sessions have been trialled to support large groups of patients simultaneously, but are challenged by the need for face-to-face schedules and limited clinical space.<sup>13</sup> There is opportunity to continue to explore feasible and effective strategies to support healthy eating in primary care.<sup>14</sup>

Providing food directly to patients in primary care is one promising approach to weight management that has been underexplored. Providing foodstuffs such as portion-controlled meals to overweight and obese adults outside of healthcare settings has revealed both clinically significant weight loss

and increased diet quality.<sup>15</sup> However, it is unclear whether this approach can be implemented in primary care where pragmatic considerations are needed, for example, identifying suitable patients, sourcing and storing foodstuffs.<sup>16</sup> Thus, we undertook a systematic review to determine whether primary care interventions involving the provision of food can support weight loss and improvements in associated health outcomes.

## METHODS

We undertook a systematic review following preferred reporting items for systematic reviews and meta-analyses guidelines.<sup>17</sup> We registered the review with the PROSPERO database (CRD42020175508).

### Search strategy

Peer-reviewed literature was searched in January 2020 with guidance from an experienced health librarian. We searched the following databases: CINAHL, MEDLINE, Embase and SCOPUS. The complete search strategy for each database is included as an online supplemental appendix. We cross-matched reference lists to identify additional studies for consideration. We imported all search results into EndNote and Covidence for study selection.

### Study selection criteria

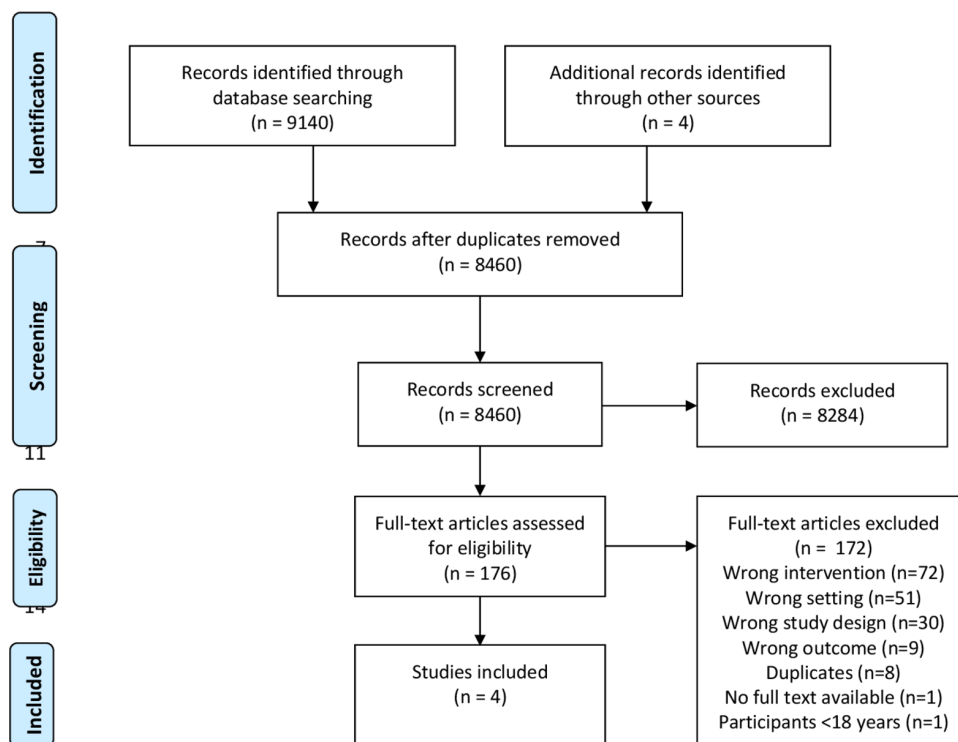
Figure 1 outlines study selection. We developed selection criteria using the population, intervention, comparison, outcome format.<sup>17</sup> We used the criteria to screen, in duplicate, the titles and abstracts of literature returned via database searching. After all duplicate and non-relevant

studies were excluded, we retrieved the remaining articles for full-text review. If insufficient information was found in the abstract, we considered the study again at the full-text review stage. We resolved all discrepancies through regular discussion.

Study selection criteria included: (1) participants: ambulatory patients aged >18 years of any gender, ethnicity and body mass index (BMI) receiving care in a primary care setting. Studies with participants in hospital, schools, aged care or lab settings were excluded. (2) Intervention: any intervention that involved the exclusive provision of foods, ingredients, oral nutrition support or food vouchers to participants. Multifaceted interventions that did not allow for the effect of food provision to be independently assessed were excluded (eg, food provision+exercise classes). (3) Comparison: baseline, usual care, no care or minimal care that does not involve intentional weight loss strategies or the provision of food. (4) Outcomes: studies that assessed change in weight were included. (5) Study design: articles were limited to published, original research studies including intervention designs (randomised controlled trials (RCTs), non-randomised) and pre-post studies. We included review articles on similar topics to full-text stage to screen the original studies within each review.

### Data extraction and synthesis

We developed a template for data extraction to collate information from included studies. Data extracted included study details, participant details, intervention, outcome measures and key findings. Data were extracted by one researcher and checked by a second,



**Figure 1** Study selection flowchart.

with discrepancies resolved through discussion. We only extracted data from applicable arms of studies. If studies reported SE or 95% CIs rather than SD, we calculated SD using the Cochrane Handbook-proposed method.<sup>18</sup> We deemed studies to have a positive overall outcome if a statistically significant reduction was reported for the weight outcome. A meta-analysis was not feasible due to the heterogeneity of included studies.

### Study quality assessment

Quality assessment was conducted using the Effective Public Health Practice Project Quality Assessment Tool for Quantitative Studies.<sup>19</sup> We selected this tool because it can assess validity of both randomised trial and pre-post designs, it has documented content and construct validity, and moderate inter-rater reliability.<sup>20</sup> The tool comprises five criteria which assess likelihood of selection bias, quality of study design, presence of confounding factors, validity and reliability of data collection, and withdrawals or dropouts. The constructs are scored as 'strong', 'moderate' or 'weak'. Each study is rated as an overview of its construct scores: 'strong' for no weak constructs; 'moderate' for one weak construct and 'weak' for two or more weak constructs. The tool considers intervention integrity and appropriateness of statistical analysis, though these do not contribute to overall study rating.<sup>19</sup> The methodological quality of articles was appraised in duplicate and discrepancies resolved through discussion. Inter-rater agreement was assessed using Cohen's kappa for construct scores within each study and across all included studies.

## RESULTS

The study selection process is illustrated in [figure 1](#). The initial database search identified 9140 articles. After duplicates were removed, we screened 8460 articles. The main reasons for excluding articles from further consideration were not being a relevant intervention (n=72) (which included multifaceted interventions, interventions not providing foodstuffs to participants and interventions not aiming to reduce weight by providing food); or not providing care in the primary care setting (n=51); or being cross-sectional in design without appropriate pre-measures and post-measures of weight (n=30). Four publications fulfilled the inclusion criteria.

### Characteristics of included studies

Characteristics of the four included studies are outlined in [table 1](#). Two of the studies were two-arm RCTs<sup>21 22</sup> and two were pre-post studies.<sup>23 24</sup> We excluded data from the control arm of one RCT because it was an ineligible comparison group.<sup>21</sup> The intervention arm was therefore treated in this review as a pre-post study.<sup>21</sup> Two of the studies were from the USA,<sup>21 23</sup> one from the UK<sup>24</sup> and one from Israel.<sup>22</sup> The studies collectively provided eligible baseline weight measures for 476 adults (range: n=22<sup>22</sup> to n=335<sup>24</sup>). Dropout between baseline and

follow-up ranged from 0%<sup>22</sup> to 38%.<sup>24</sup> Two studies only recruited overweight or obese participants<sup>21 24</sup> and two studies did not consider BMI when screening for potential participants.<sup>22 23</sup>

### Intervention description

One study provided participants with three sachets of green tea and a vitamin E supplement daily for 12 weeks,<sup>22</sup> taken in addition to the participants' usual diet. Two studies examined the effectiveness of providing meal-replacement shakes to facilitate low-energy diets.<sup>21 24</sup> Of these studies, one provided participants with three shakes per day for 30 days, replacing all meals with no extra food to be consumed.<sup>24</sup> The second meal replacement study provided two shakes per day for 12 weeks (replacing two meals, and allowing participants to prepare and consume one meal each day) followed by one shake per day for a further 10 weeks (replacing one meal, allowing participants to prepare and consume two meals each day).<sup>21</sup> The final study hosted a 'farmers' market' at the primary care clinic for 13 weeks and provided participants with vouchers (US\$10) to purchase fresh food for consumption as part of their usual diet and requested participants complete a goals worksheet to guide programme participation.<sup>23</sup>

### Results of individual studies

[Table 2](#) outlines the baseline and follow-up weight data for each study as well as other relevant primary and secondary outcome measures.

### Weight loss

Three of the four studies observed weight loss among participants. While some standard conversions could be made to align data where possible (eg, lbs to kg), all studies reported results using varied units and descriptors, making comparisons challenging. In the RCT that provided green tea and vitamin E supplementation, both the intervention and control groups observed significant reductions in weight (intervention mean 76.8±10.3kg at week 0 to mean 73.4±9.9kg at week 12; p=0.004; control mean 76.2±7.6kg at week 0 to mean 74.5±8.0kg at week 12; p=0.004), with no significant difference reported between groups.<sup>22</sup> In the pre-post study where community pharmacists delivered a meal replacement programme, participants lost a significant amount of weight during the active phase (12 weeks) (mean loss -4.90±0.30kg, p<0.0001).<sup>21</sup> Using intention-to-treat (ITT) analysis, the average percentage loss was 5.98% of initial body weight across the first 12 weeks of the study.<sup>21</sup> The second pre-post study that used a meal-replacement product reported a reduction in weight but failed to report if the weight loss was statistically significant.<sup>24</sup> In this study, the average weight loss among trial completers was 6.6kg with best results achieved by those participants who completed the 30-day trial and visited their doctor weekly (7.2kg weight loss).<sup>24</sup> In the farmers' market study, no significant

**Table 1** Characteristics of included studies examining the effectiveness of providing food in primary care for weight loss, in order of recency

First author, year, country	Relevant aim	Intervention	Comparison	Participants	Study length	Dropouts	Relevant outcomes assessed	Fidelity
Bryce, 2017, <sup>23</sup> USA	To examine the impact of a farmers' market with fresh fruits and vegetables at reducing weight.	US\$10–15/visit voucher for farmers' market (up to four visits); maximum US\$45 available across study period	Baseline	74 participants with uncontrolled type 2 diabetes, low income	13 weeks	12.2% (n=9)	Weight (lbs), HbA1c (%), BP (mm Hg)	41 participants attended all four market visits, 55 participants completed the health goals sheet
Narotzki, 2013, <sup>22</sup> Israel	To examine the effectiveness of daily green tea and vitamin E consumption at reducing weight.	3×1.5g green tea per day (to be brewed in 240 mL water)+400 IU vitamin E+exercise	CON group (vitamin E placebo+exercise)	22 participants (n=11 intervention) aged 61–80 years	12 weeks	0% (n=0)	Weight (kg), BP (mm Hg), HbA1c (%), blood lipid profile, waist circumference (cm)	INT: green tea 93±8.3%, vitamin E supplement 95±5.6% and exercise 88±15.6% CON: placebo supplement 94± 6.6% and exercise 92±9.3%
Ahrens, 2003, <sup>21</sup> USA	To examine the effectiveness of a community pharmacist-facilitated meal replacement programme at reducing weight.	2 shakes per day for 12 weeks (INT) followed by 1 shake per day for 10 weeks (maintenance)	Baseline (CON arm did not meet review criteria due to dietitian input)	88 participants (n=45 intervention) aged 35–65 years, BMI 25–32 kg/m <sup>2</sup>	12 weeks+10 weeks	38.6% (n=34)	Weight (kg), BP (mm Hg), blood lipid profile, waist circumference (cm)	Assessed but NR
Isaac, 1984, <sup>24</sup> UK	To examine the effectiveness of a GP-facilitated meal replacement programme at reducing weight.	3 shakes per day in place of all foods		335 overweight or obese participants, 20–68 years	30 days	35.2% (n=118)	Weight (kg), BP (mm Hg), acceptability (including side effects)	88 participants did not fully comply with the intervention

.BMI, body mass index; BP, blood pressure; CON, control; GP, general practitioner; HbA1c, glycated haemoglobin; INT, intervention; NR, not reported.

**Table 2** Results of included studies examining the effectiveness of providing food in primary care to reduce weight in patients, in order of recency

First author, year, country	Weight results	Significance	Other outcome measures			
			BP	Blood lipids	Waist circumference	HbA1c
Bryce, 2017, <sup>23</sup> USA	Baseline: 208.3 lbs (94.68 kg) Follow-up: 209.0 lbs (95 kg) Percentage change: +0.3	No difference between baseline and follow-up observed (p>0.05)	No significant change from baseline to follow-up for SBP (p=0.70) or DBP (p=0.17)	NA	NA	Significant reduction in mean HbA1c from 9.54% to 8.83% (t-score=3.54, p=0.001)
Narotzki, 2013, <sup>22</sup> Israel	Baseline: INT: 76.8±10.3kg CON: 76.2±7.6kg Follow-up: INT: 73.4±9.9kg CON: 74.5±8.0kg Percentage change: INT: -4.4 CON: -2.2	No difference between groups at follow-up (p>0.05); both groups observed significant weight loss (INT mean -3.4 kg, p=0.004; CON mean -1.7 kg, p=0.004)	No significant difference between the groups at follow-up for SBP (p=0.409) or DBP (p=0.675); no significant changes within either group	No significant difference between groups at follow-up for TG (p=0.233), LDL-C (p=0.98) or HDL-C (p=0.671); no significant changes within either group	No significant difference between groups at baseline or follow-up (p>0.05). INT men and women observed a significant reduction (men: -3.8±3.4%, p=0.026; women: -11.2±3.1%, p=0.005)	No significant difference between the groups at baseline or follow-up (p>0.05); no significant change within either group
Ahrens, 2003, <sup>21</sup> USA	Baseline: 81.9±11.1 kg 12 weeks: 77.0±10.2 kg Percentage change: -6	Mean change -4.9 kg, p<0.0001	Mean change 12 weeks SBP -5.91 mm Hg (-4.56%), DBP -4.16 mm Hg (-5.79%). Significance of changes NR	Mean change 12 weeks TC -11.95 mg/dL (-5.87%); TG -0.08 mg/dL (-1.73%); LDL-C -8.86 mg/dL (-7.08%); HDL-C -0.19 mg/dL (-0.36%). Significance of changes NR	Mean change 12 weeks -5.31 cm (-5.96%). Significance of change NR	NA
Isaacs, 1984, <sup>24</sup> UK	Baseline: 88.93±15.5 kg Follow-up: 82.27±14.9 kg Percentage change: -7.5	Significance of change NA	Those who completed the trial and had initial DBP >90 mm Hg observed a significant reduction in mean BP (baseline 158/101, follow-up 145/89, p<0.0001)	NA	NA	NA

BP, blood pressure; CON, control; DBP, diastolic blood pressure; HbA1c, glycated haemoglobin; HDL-C, high-density lipoprotein cholesterol; INT, intervention; LDL-C, low-density lipoprotein cholesterol; NA, not assessed; NR, not reported; SBP, systolic blood pressure; TC, total cholesterol; TG, triglycerides.

change in body weight was reported between baseline and 13 weeks (p=0.45).<sup>23</sup>

**Health outcomes**

All studies reported secondary health outcomes. In the RCT that provided green tea and vitamin E supplementation, a significant decrease in fasting insulin levels was observed between baseline and week 12 for both groups (intervention -22.2±25.5%, p=0.016; control -25.2±28.5%, p=0.015).<sup>22</sup> A reduction in fasting glucose (intervention -5.9±5.6%, p=0.005; control +4.9±9.0%, p>0.05) and waist circumference for both men and women (intervention men -3.8±3.4%, p=0.026; intervention women -11.2±3.1%, p=0.005, respectively) was seen in the intervention group.<sup>22</sup> Blood pressure, C reactive protein and serum lipids did not change throughout the study in either group or between groups.<sup>22</sup> Dietary intakes

(ie, macronutrient distribution, vitamin E intake, vitamin C intake and kilojoule intake) also did not change significantly within or between groups.

In the pre-post study where community pharmacists delivered a meal replacement programme, improvements were observed (but no statistical testing reported) at the end of 12 weeks (weight loss phase) and 22 weeks (weight maintenance phase) for waist circumference (-5.31 cm and -8.08 cm, respectively), systolic blood pressure (-5.91 mm Hg and -8.26 mm Hg) and diastolic blood pressure (-4.16 mm Hg and -5.97 mm Hg).<sup>21</sup> No improvements were observed by the end of the study for triglyceride levels (-0.08 mg/dL and 0.00 mg/dL, respectively), total cholesterol (-11.95 mg/dL and 4.74 mg/dL, respectively), high-density lipoprotein cholesterol (-0.19 mg/dL and 4.44 mg/dL, respectively) or

**Table 3** Quality assessment of included studies using the Effective Public Health Policy Projects Quantitative

Author (year)	Intervention integrity						Analysis				Rating			
	Selection design	Study design	Confounders	Blinding	Data collection methods	Withdrawal and dropouts	Participants received allocated intervention	Was consistency of intervention measured	Likelihood of contamination	Unit of allocation		Unit of analysis	Appropriate statistical methods	ITT
Ahrens <i>et al</i> (2003) <sup>21</sup>	Weak	Strong	Strong	Weak	Moderate	Weak	Unclear	Unclear	Unclear	Individual	Individual	Yes	Unclear	Weak
Bryce <i>et al</i> (2017) <sup>23</sup>	Weak	Moderate	Weak	Weak	Weak	Strong	80%–100%	Yes	Yes	Practice	Individual	Yes	No	Weak
Isaacs and Parry (1984) <sup>24</sup>	Moderate	Moderate	Moderate	Weak	Weak	Moderate	80%–100%	Yes	Unclear	Practice	Individual	Yes	Unclear	Weak
Narotzki <i>et al</i> (2013) <sup>22</sup>	Weak	Strong	Strong	Weak	Strong	Strong	80%–100%	Yes	Unclear	Individual	Individual	Yes	Yes	Weak

ITT, intention to treat.

low-density lipoprotein cholesterol levels (−8.86 mg/dL and 1.15 mg/dL, respectively).<sup>21</sup>

The second pre–post study that used a meal-replacement product saw nearly two-thirds of the participants with initially hypertension became normotensive (45 of the 73 patients with an initial diastolic blood pressure recording above 90 mm Hg), but no other changes in health outcomes were assessed.<sup>24</sup> In the farmers’ market study, a significant decrease in glycated haemoglobin (HbA1c) was observed 3 months after the 13-week study period (mean 9.54% to mean 8.83%, t-score 3.54, p=0.001) but no significant changes were reported for systolic or diastolic blood pressure.<sup>23</sup>

### Attrition and safety

Attrition and side effects varied between the studies. In the RCT that provided green tea and vitamin E supplementation, compliance with green tea and vitamin E consumption was 93±8.3% and 95±5.6%, respectively. In the control group, the placebo supplement compliance was 94±6.6%. The level of vitamin E intake, 400 IU/day, was based on literature that less than 1000 IU/day is safe in healthy individuals.<sup>22</sup> In the study that used community pharmacists to deliver a meal replacement programme, a dropout rate was reported for the total study rather than the arm that was included for review, as 23% for the active weight loss phase (two shakes per day), and a further 19% during the weight maintenance phase (one shake per day).<sup>21</sup> Major reasons reported for attrition were time constraints and loss to follow-up, while only two participants reported intolerance to the meal-replacement shakes.<sup>21</sup>

The *Modifast* product used in the second pre–post study that used a meal replacement was rated on average ‘tolerable’, but side effects, although, generally minor and transient, were reported by 110 participants (32.8%) and significantly affected their ability to complete the trial.<sup>24</sup> Side effects were reported by 21.7% of those who completed the trial successfully, 30.7% of those who lapsed during the trial (26.3%, n=88) and 42.4% among those who did not complete the trial (25.3%, n=118).<sup>24</sup> In the farmers’ market study, most participants (63.1%, n=41) attended the required four market visits and of these, 84.6% (55 of 65) completed their health goals sheet and received a US\$5 incentive voucher.<sup>23</sup> No adverse events or concerns were reported for hosting the farmers’ market at the primary care clinic.<sup>23</sup>

### Quality appraisal

Table 3 describes the methodological quality of each study. All studies were rated as ‘weak’.<sup>21–24</sup> Common weaknesses related to blinding of participants and outcome assessors,<sup>21–24</sup> use of non-validated tools for data collection or lack of data collection tool description,<sup>23 24</sup> and potential selection bias.<sup>21–23</sup> Most studies poorly described the intervention and steps taken to avoid contamination.<sup>21 22 24</sup> While statistical methods appeared appropriate for all study designs, there was inconsistent reporting of

results within studies,<sup>21 24</sup> poor descriptions of the study groups,<sup>21 23 24</sup> and low use or poor application of ITT analysis.<sup>21 23 24</sup> Agreement for overall study ratings was 100%. Inter-rater reliability for assessment of constructs within studies was moderate to substantial ( $\kappa=0.44-0.72$ ). Overall construct agreement was moderate ( $\kappa=0.58$ ).

## DISCUSSION

This review explored the concept of providing food to patients in primary care as a means of supporting weight loss. Primary care is regarded as an ideal setting to support healthy eating and weight management,<sup>1</sup> but notable barriers often prevent health professionals addressing nutrition and weight during routine care.<sup>8-10</sup> The body of literature on this topic is notably small, with only four studies eligible for this review.<sup>21-24</sup> There is clear opportunity for further research that advances the understanding gained from these four studies and ultimately inform policy initiatives for providing food to support weight loss and improvements in other health outcomes.

The use of meal replacements to initiate weight loss in overweight patients has potential in the general practice setting for patients who desire structured weight loss programmes. Some potentially relevant studies were not included in our review due to the heavy reliance on education.<sup>25 26</sup> Significant weight loss and other improvements in biomarkers of disease were observed for both meal replacement studies. Commercial meal-replacement products have been shown in one review to be a convenient, often low-cost means of supplying nutritionally dense, yet low-energy meal options, which when combined with low-fat self-selected meals, significantly improves weight loss and provides a safe and effective method for maintaining that weight loss.<sup>27</sup> While the review did not restrict studies to those conducted in primary care, a recognised challenge of incorporating meal replacements into usual care is low tolerance and acceptability.<sup>27</sup> Further research is required to investigate the suitability and likely uptake of meal replacements in primary care clinics and the potential cost-effectiveness for reducing weight-related morbidity.

In the green tea and vitamin E supplement study, improved exercise capacity was followed by a significant reduction in body weight and fasting insulin levels in all participants.<sup>22</sup> Even though both the intervention and control groups lost weight in the trial, the small sample size means an effect may still be possible. Polyphenolic compounds present in green tea catechins are thought to influence sympathetic nervous system activity, increasing energy expenditure and promoting the oxidation of fat, which may explain the weight loss.<sup>28</sup> Additionally, caffeine which is naturally present in green tea also influences sympathetic nervous system activity, and may act synergistically to increase energy expenditure and fat oxidation.<sup>28 29</sup> In a recent systematic review of the effects of green tea with 1945 participants for 12–13 weeks, weight loss

was small and statistically insignificant, and viewed by the authors as unlikely to be clinically important.<sup>29</sup> Further studies are required to investigate whether the positive outcomes may stem from green tea or vitamin E and the potential benefits and cost-effectiveness of green tea and vitamin E supplementation to initiate weight loss among overweight/obese patients in primary care.

Fruit and vegetable markets are currently being investigated in Australia as a novel approach to support fruit and vegetable consumption among disadvantaged populations and those affected by restrictions imposed from the coronavirus pandemic.<sup>30</sup> Although the Bryce *et al* study did not observe significant weight loss as a result of vouchers for a local farmers' market being incorporated into routine care, a significant decrease in the HbA1C concentrations was reported, suggesting that the intervention may have had overall benefits.<sup>23</sup> However, a notable limitation is that the cost of implementing this intervention and the sustainability of the intervention were not examined. While the study in this review did not actually measure change in food intake as a result of farmers' market vouchers, the benefits of increased fruit and vegetable consumption from shopping at local farmers' markets have been reported in other similar studies with specific populations such as low-income patients with type 2 diabetes.<sup>31</sup> Modest financial vouchers provided to low-income adults in urban areas have been shown to be an effective incentive to shop at farmers' markets.<sup>32 33</sup> Financial hardship is recognised as one key reason for poor diets (ref).<sup>34</sup> Given these challenges, the findings of the farmers' market study demonstrate the multifaceted potential benefit of a food prescription programme for disadvantaged populations to contribute to improved health outcomes. Future studies should investigate the impact of food provision on measures of food security, access and affordability as well as dietary quality and acceptability.

The acceptability and unintended consequences of providing food to patients in primary care warrant consideration. This review excluded studies that had an educational focus, which could provide an ethical barrier to providing foodstuffs. Future studies should clearly describe the components of interventions to better investigate the impact on outcome measures. Side effects such as constipation were apparent in one of the studies that used meal-replacement products,<sup>24</sup> but were monitored more carefully in the other.<sup>21</sup> However, patient dropout from the meal replacement study that used pharmacists was high: ranging 28% in the weight loss phase to 42% during weight maintenance.<sup>21</sup> High dropout is not uncommon in weight loss studies due to the challenge of behaviour change.<sup>35</sup> Attrition can be reduced through structured, planned meals that require little participant choice; though ironically, long term this may result in

flavour fatigue and dropout.<sup>36</sup> High levels of dropout in weight loss studies highlight the need to employ appropriate ITT analyses, meaning dropouts are not excluded from final measures of effectiveness. Both studies in this review that used meal replacements employed ITT analyses for weight loss<sup>21 24</sup> though also reported non-ITT analyses for weight loss,<sup>21 24</sup> weight maintenance<sup>21</sup> and other health outcomes<sup>21 24</sup> which reflects efficacy rather than effectiveness. Overall, a small yet notable body of literature supports the concept of providing food to patients in primary care settings to support weight loss. Further, high-quality research is needed on the efficacy and cost-effectiveness of this approach to ultimately inform policy initiatives for primary care.

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