Does the effect of lifestyle intervention for individuals with diabetes vary by food insecurity status? A preplanned subgroup analysis of the REAL HEALTH randomized clinical trial

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ABSTRACT

Introduction We aimed to test the effectiveness of a lifestyle intervention (LI) for individuals with food insecurity and type 2 diabetes.

Research design and methods Adults with type 2 diabetes, body mass index ≥25 kg/m² (or ≥23 kg/m² if Asian), hemoglobin A1c of 6.5%–11.5% (48–97 mmol/mol) and who were willing to lose 5%–7% bodyweight were enrolled in REAL HEALTH-Diabetes. This practice-based randomized clinical trial compared LI (delivered in person or by telephone) with medical nutrition therapy (MNT) on weight loss at 6 and 12 months. Two or more affirmative responses on the six-item US Department of Agriculture Food Security Survey Module indicated food insecurity.

In this prespecified subgroup analysis, we tested using linear mixed effects models whether the intervention effect varied by food security status.

Results Of 208 participants, 13% were food insecure. Those with food insecurity were more likely to be racial/ethnic minorities (p<0.001) and have lower education (p<0.001). LI, versus MNT, led to greater weight loss at 6 months (5.1% lost vs 1.1% lost; p=0.0005) and 12 months (5.1% lost vs 1.1% lost; p<0.0001) and 12 months (5.1% lost vs 1.1% lost; p<0.0001). LI, versus MNT, led to greater weight loss at 6 months (5.1% lost vs 1.1% lost; p=0.0005). The intervention effect was similar regardless of food security status (5.1% bodyweight lost vs 1.1% in food secure participants and 5.1% bodyweight lost vs 1.3% in food insecure participants at 6 months; 4.7% bodyweight lost vs 2.1% in food secure participants and 4.5% bodyweight lost vs 0.9% in food insecure participants at 12 months; p for interaction=0.99).

Conclusions The REAL HEALTH-Diabetes lifestyle intervention led to meaningful weight loss for individuals with food insecurity and type 2 diabetes.

Trial registration number NCT023220253.

The primary goal of type 2 diabetes management is to prevent complications of diabetes by controlling risk factors.1 However, the burden of diabetes and its complications does not fall evenly—individuals with lower socioeconomic status face increased risk of diabetes as well as worse diabetes outcomes.2 One potentially modifiable contributor to this is food insecurity, defined as limited access to sufficient food for a healthy, active life.3 Food insecurity is strongly associated with greater obesity and worse risk factor control in individuals with diabetes.4–9

Typical food insecurity interventions focus on increasing healthy food access by subsidizing or providing healthy food.10 11 While important, the clinical impact of these approaches is not yet known, and a prior randomized trial of a food pantry-based intervention did not result in changes in diabetes outcomes.12 Given the complexity of the relationship between food insecurity and clinical outcomes,3 13 additional interventions for individuals with food insecurity which could either be combined with food access interventions or serve as stand-alone approaches...
are needed. Lifestyle intervention (LI), which combines education and skill-building in order to facilitate diabetes self-management, could be one way to meet this need. In both ‘efficacy’ trials and in more ‘real-world’ applications, LIs have led to weight loss and improvements in diabetes control, risk factors for diabetes complications, and complications themselves. Further, among food insecure populations, education and skill-building approaches that focus on making best use of available resources, exemplified by Supplemental Nutrition Assistance Program Education (SNAP-Ed), a program that helps SNAP recipients and communities use farmers markets to address food insecurity, have been shown to be successful in improving diet quality.

Despite the promise of LI, however, few studies have examined LI for management of type 2 diabetes in individuals with food insecurity. Testing the effectiveness of LI in those with food insecurity is important, since food insecurity, for several reasons, is likely to present a barrier to weight loss that may render LI less effective. Mechanistically, food insecurity may act to thwart lifestyle change by incentivizing the consumption of highly processed, calorie-dense foods that are cheaper on a per-calorie basis than healthier alternatives. Further, food insecurity may impose competing demands and increase stress, which makes enacting behavior change more difficult. Finally, the time commitment required for LI may be prohibitively high for those with many competing demands, such as work and childcare. Alternatively, the skills taught during LI may help facilitate more effective diabetes self-management even in the face of challenging circumstances like food insecurity, if those interventions are appropriately adapted to the needs of individuals with lower socioeconomic status. For these reasons, we believe it is important to determine whether the effectiveness of LI varies by food security status.

In this prespecified subgroup analysis of the REAL HEALTH-Diabetes pragmatic randomized clinical trial, we sought to determine whether an evidence-based LI, delivered in English or Spanish, in-person or via telephone, would result in clinically meaningful weight loss in individuals with type 2 diabetes both with and without food insecurity, and whether the magnitude of weight loss would vary by food security status.

RESEARCH DESIGN AND METHODS

Participants and study setting

A description of the REAL HEALTH-Diabetes randomized clinical trial design and participants has been previously published. In brief, individuals were eligible if they were adults (age ≥18 years) with type 2 diabetes, body mass index ≥25 kg/m² (or ≥23 kg/m² if of self-reported Asian ancestry), had hemoglobin A1c (HbA1c) of 6.5%–11.5% (48–97 mmol/mol), had blood pressure <160/100 mm Hg, and were willing to lose 5%–7% body-weight. Individuals were drawn from those with a medical provider in the Partners HealthCare clinical network. Individuals were excluded if they weighed more than 159 kg, spoke a language other than English or Spanish, were pregnant or planning to become pregnant, were currently enrolled in a weight loss program or medical nutrition therapy (MNT), or had a previous or planned bariatric procedure. Further, those who could not participate in the intervention owing to a medical condition that would interfere with the ability to engage in the LI or were taking medications likely to affect weight (eg, corticosteroids) were also excluded. Intervention and data collection visits were conducted either at one of three community health centers or at the Massachusetts General Hospital Diabetes Research Center, depending on the location of the patient’s residence.

Participants were recruited between January 2015 and July 2017, with 1-year outcomes analyzed from December 2018 to January 2020. All trial participants provided written informed consent. Prior to participant enrollment, the trial was registered at ClinicalTrials.gov.

INTERVENTION

REAL HEALTH-Diabetes randomized individuals in a 1:1:1 ratio to in-person group LI, telephone conference call group LI, or MNT. Randomization was accomplished using a computer-generated variable block size randomization scheme stratified by gender and clinical site. The LI was a complex behavioral weight loss intervention developed as a curated hybrid of key publicly available Look AHEAD and Diabetes Prevention Program (DPP) session materials and included the first 19 sessions from the original Look AHEAD LI. The LI addressed a variety of nutrition, activity, and behavioral topics and focused on building skills for self-monitoring, goal setting, stimulus control, stress management, problem solving, and relapse prevention. The session materials were culturally sensitive and had a Flesch Reading Ease score of 74.0 and a Flesch-Kincaid grade level of 5.3. However, the materials were not specifically adapted to address the needs of those with food insecurity or low socioeconomic status. Similar to Look AHEAD’s LI, participants were encouraged to purchase and use meal replacements in the form of shakes, bars or frozen entrees to replace one to two meals per day to promote weight loss. Unlike the original Look AHEAD LI, which was delivered largely in individual sessions supplemented by some group sessions, the REAL HEALTH LI sessions were delivered by registered dietitians in groups of 4–12 (stratified by English or Spanish delivery of materials) and were delivered at a decreasing cadence of 14 weekly sessions followed by 5 biweekly and 6 monthly sessions, for a total of 25 sessions over 1 year. Participants could also elect to take part in up to three individual sessions during the year. In-person and telephone conference call sessions were identical in number, duration (approximately 60–90 min) and content—the only difference between these sessions was the delivery format. Using a 60 min estimate, this means that planned contact for LI participants across 25 planned sessions was

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25 hours. The planned duration of each optional session was 45 min or 2.25 hours for three sessions.

Individuals randomized to MNT were referred to individual in-person sessions with a dietitian at their health center or another location of their choice as part of usual care (which is the current recommended standard of care). MNT proceeded per usual care, with an initial visit for evaluation with follow-up sessions as determined by the dietitian and the participant. While LI was provided without cost sharing, MNT could require visit copayments depending on the participant’s insurance coverage, as is typical in usual care. For MNT sessions, the planned duration was 60 min for the first session and 30 min for each additional session. As in usual care, there was no planned number of sessions for MNT, as that is determined in a dialogue between the registered dietitian and the participant. As an example, if an MNT participant attended one initial session and three follow-up sessions over 12 months, that would be 2.5 hours of contact.

**Food insecurity**

Food insecurity assessment occurred at the baseline study visit, prior to randomization. Food insecurity was assessed using the six-item US Department of Agriculture Food Security Survey Module with 12-month recall period.24 An example item is: ‘The food that I bought just didn’t last, and I didn’t have money to get more. Was that often, sometimes, or never true for you in the last 12 months?’24 In accordance with standard scoring, those with two or more affirmative responses were classified as food insecure.24 Based on prior work examining food insecurity prevalence in individuals with diabetes, including within the clinical network study participants were drawn from, we anticipated food insecurity prevalence of approximately 20%.8 9 25

In addition to categorizing participants by food security status, we also categorized individuals by household income level, using categories of <$30 000 per year, $30 000–$50 000 per year, $50 000–$100 000 per year, and ≥$100 000 per year.

**Outcomes**

The primary outcome for this study was weight loss, defined as per cent change in weight from baseline. A 3% decrease in weight from baseline was considered a clinically meaningful change. Weights were obtained using a calibrated scale in light clothing following a standardized protocol.21 The trained research assistant measuring the weights was masked to study arm. Another secondary outcome was HbA1c, which was analyzed at a clinical laboratory certified by the NGSP (formerly the ‘National Glycohemoglobin Standardization Program’). To reduce the risk of hypoglycemia with LI, the REAL HEALTH-Diabetes trial protocol reduced glucose-lowering medication proactively based on self-monitoring of blood glucose results, anticipated reduction in caloric intake, and use of meal replacements to enhance weight loss.21 Finally, we examined session attendance.

**Covariates and other data collection**

Information relating to sociodemographics, anthropometrics, and laboratory values was obtained during the baseline visit prior to randomization and at follow-up study visits.21

**Statistical analyses**

All analyses followed the intention-to-treat approach for intervention arm classification. Because prior analyses found that the in-person and telephone-delivered LI produced similar results,22 we combined these intervention arms for analysis, comparing those who received either format of LI with those who received MNT. To account for repeated measures within individuals and possible clustering within the study arms in which the LI was delivered, we used three-level mixed effect models with random effects terms for both the individual and intervention group. We examined changes at 6 and 12 months and used interaction terms (food security (or income category) by intervention modality) to examine whether intervention effect varied by food security status or income category. For these analyses, the food insecurity by intervention modality analysis was considered the primary analysis, with other analyses considered exploratory. Nine (4.3%) participants did not report income levels and were excluded from income analyses.

We conducted both unadjusted analyses and analyses that adjusted for possible confounders of the relationship between food insecurity (or income) and the outcomes. For the food insecurity models, the adjustment variables were age, gender, race/ethnicity, education, income, primary language, and health insurance. For the income models, we used the same set of adjustment variables minus income. Missing data were not imputed because missingness was less than 5% for any variable. All analyses were conducted in SAS V.9.4. A two-sided p value of ≤0.05 was considered statistically significant.

**RESULTS**

As previously described, 211 individuals enrolled in the trial and 3 (1.4%) withdrew, leaving 208 participants (98.6%) who were included in the intention-to-treat analyses.22 The mean age of participants was 61.7 (SD: 10.2) years, 55.3% were women, 76.9% were of non-Hispanic white race/ethnicity, and 8.2% spoke Spanish as their primary language (table 1). Thirteen per cent of participants reported food insecurity, with a similar proportion in the two study arms, as expected in a randomized trial. Overall, those who reported food insecurity were more likely to be women and of racial/ethnic minorities, and to have less education, lower income, and public health insurance.

Compared with those receiving MNT, those receiving LI lost a greater percentage of baseline weight at 6 months (5.1% lost vs 1.1% lost; p<0.0001) and 12 months (4.7% lost vs 2.0% lost; p=0.0005).22 The intervention effect in the LI arm was similar regardless of food security status.
| Table 1  | Participant characteristics by food security status |
|----------|------------------------------------------------------|
|          | All   | Food secure | Food insecure | P value* |
| n        | 208   | 180         | 28            |         |
| Study arm, n (%) |       |             |               | 1.00    |
| Lifestyle intervention | 142 (68.3) | 123 (68.3) | 19 (67.9) |       |
| Medical nutrition therapy | 66 (31.7) | 57 (31.7)  | 9 (32.1)   |       |
| Age, mean (SD) | 61.7 (10.2) | 62.3 (9.9) | 58.0 (11.5) | 0.072  |
| Female, n (%) | 115 (55.3) | 93 (51.7)  | 22 (78.6)  | 0.008  |
| Race/ethnicity, n (%) |       |             |               | <0.0001 |
| Non-Hispanic white | 160 (76.9) | 148 (82.2) | 12 (42.9)  |       |
| Non-Hispanic black  | 9 (4.3)   | 9 (5.0)     | 0 (0.0)     |       |
| Hispanic/Latino     | 28 (13.5) | 14 (7.8)    | 14 (50.0)   |       |
| Non-Hispanic Asian  | 7 (3.4)   | 5 (2.8)     | 2 (7.1)     |       |
| Other               | 4 (1.9)   | 4 (2.2)     | 0 (0.0)     |       |
| Education, n (%)    |       |             |               |         |
| Less than high school diploma | 17 (8.2)   | 10 (5.6)    | 7 (25.0)    |       |
| High school diploma or GED | 43 (20.7) | 39 (21.7)  | 4 (14.3)    |       |
| 1–3 years of college | 60 (28.8) | 47 (26.1)  | 13 (46.4)   |       |
| 4 or more years of college or graduate school | 88 (42.3) | 84 (46.7)  | 4 (14.3)    |       |
| Annual income, n (%) |       |             |               | <0.0001 |
| <$30 000 | 48 (23.1) | 30 (16.7)   | 18 (64.3)    |       |
| $30 000–$50 000  | 21 (10.1) | 19 (10.6)   | 2 (7.1)      |       |
| $50 000–$100 000 | 68 (32.7) | 65 (36.1)   | 3 (10.7)     |       |
| ≥$100 000 | 62 (29.8) | 60 (33.3)   | 2 (7.1)      |       |
| Did not report | 9 (4.3)   | 6 (3.3)     | 3 (10.7)     |       |
| Insurance, n (%)    |       |             |               | 0.001   |
| Commercial | 88 (42.3) | 80 (44.4)   | 8 (28.6)     |       |
| Medicaid | 29 (13.9) | 25 (13.9)   | 4 (14.3)     |       |
| Medicare | 59 (28.4) | 55 (30.6)   | 4 (14.3)     |       |
| Medicare and Medicaid (‘dual eligible’) | 26 (12.5) | 16 (8.9)    | 10 (35.7)    |       |
| Self-pay | 6 (2.9)   | 4 (2.2)     | 2 (7.1)      |       |
| Spanish language, n (%) | 17 (8.2)   | 9 (5.0)     | 8 (28.6)     | 0.0004 |
| Baseline hemoglobin A1c | 7.7 (1.2)   | 7.7 (1.2)   | 8.0 (1.2)    | 0.25   |
| NGSP %, mean (SD)   | 61.1 (12.8)| 60.6 (12.7)| 63.7 (13.2)  |       |
| Baseline body mass index, kg/m², mean (SD) | 35.0 (5.4) | 35.0 (5.5) | 35.3 (4.7) | 0.75   |
| Baseline weight, kg, mean (SD) | 98.1 (18.9) | 98.9 (19.2) | 93.4 (16.1) | 0.11   |
| Baseline use of any glucose-lowering medication | 191 (91.8) | 163 (90.6) | 28 (100) | 0.14   |
| Baseline use of metformin | 166 (79.8) | 141 (78.3) | 25 (89.3) | 0.21   |
| Baseline use of insulin | 69 (33.2) | 53 (29.4) | 16 (57.1) | 0.005  |
| Baseline use of GLP-1 RA | 14 (6.7) | 13 (7.2) | 1 (3.6) | 0.70   |
| Baseline use of sulfonylurea | 65 (31.3) | 58 (32.2) | 7 (25.0) | 0.52   |
| Baseline use of DPP-4 | 9 (4.3) | 8 (4.4) | 1 (3.6) | 1.00   |
| Baseline use of SGLT2 | 6 (2.9) | 4 (2.2) | 2 (7.1) | 0.19   |

*P value from two-sample t-tests for continuous variables and Fisher’s exact tests for categorical variables.
DPP-4, dipeptidyl peptidase-4 inhibitor; GED, General Education Development certificate; GLP-1 RA, glucagon-like peptide 1 receptor agonist; NGSP, National Glycohemoglobin Standardization Program; SGLT2, sodium/glucose cotransporter 2 inhibitor.
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(5.1% bodyweight lost vs 1.1% in food secure participants and 5.1% bodyweight lost vs 1.3% in food insecure participants at 6 months; 4.7% bodyweight lost vs 2.1% in food secure participants and 4.5% bodyweight lost vs 0.9% in food insecure participants at 12 months) (table 2, figure 1). Interaction terms testing these differences in intervention effect were not statistically significant in either unadjusted (p=0.87) or adjusted (p=0.99) models. Change in HbA1c did not differ between the LI and MNT groups. HbA1c was 0.5% (5.7 mmol/mol) lower

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**Table 2** Per cent change in weight and hemoglobin A1c from baseline at 6 and 12 months

| Food security* | Change in weight | Change in hemoglobin A1c, NGSP % (mmol/mol) |
|----------------|-----------------|---------------------------------------------|
|                | 6 months | 12 months | 6 months | 12 months |          |          |
| **n Mean SD**  |          |          |          |          |          |          |
| Secure MNT     | 57 -1.1 3.7 | 55 -2.1 4.1 | 56 -0.2 (2.5) | 1.0 (11.3) | 54 -0.4 (3.9) | 1.3 (13.7) |
| LI             | 122 -5.1 5.3 | 122 -4.7 6.1 | 120 -0.6 (6.0) | 1.0 (11.5) | 122 -0.3 (3.8) | 1.1 (12.4) |
| Insecure MNT   | 9 -1.3 3.3 | 9 -0.9 3.8 | 9 -0.5 (5.8) | 1.7 (18.6) | 8 -0.5 (5.9) | 1.2 (13.4) |
| LI             | 19 -5.1 5.8 | 18 -4.5 6.0 | 19 -0.3 (3.6) | 0.8 (9.3) | 18 -0.1 (1.5) | 0.7 (8.1) |

| Income†       | Change in weight | Change in hemoglobin A1c, NGSP % (mmol/mol) |
|---------------|-----------------|---------------------------------------------|
|               | 6 months | 12 months | 6 months | 12 months |          |          |
| <$30 000 MNT  | 17 -1.0 3.5 | 16 -1.1 3.6 | 17 -0.2 (2.6) | 1.3 (14.3) | 16 -0.3 (3.4) | 1.0 (11.1) |
| LI            | 31 -3.3 4.9 | 30 -3.2 6.6 | 31 -0.4 (4.4) | 0.7 (7.9) | 30 -0.3 (2.8) | 0.8 (9.0) |
| $30 000–$50 000 MNT | 3 -1.8 1.4 | 3 -1.6 1.3 | 3 -0.1 (0.7) | 0.6 (6.4) | 3 -0.0 (0.4) | 0.7 (7.4) |
| LI            | 18 -5.4 4.2 | 18 -4.3 5.7 | 18 -0.6 (6.5) | 1.0 (11.2) | 18 0.0 (0.1) | 1.2 (13.5) |
| $50 000–$100 000 MNT | 17 -1.2 3.5 | 17 -2.2 3.5 | 16 0.1 (1.2) | 1.0 (11.0) | 16 -0.1 (1.4) | 1.1 (11.6) |
| LI            | 50 -5.4 5.8 | 51 -5.0 5.7 | 48 -0.6 (6.8) | 1.3 (13.7) | 51 -0.6 (6.0) | 1.2 (12.8) |
| ≥$100 000 MNT | 26 -0.9 4.3 | 25 -1.9 4.6 | 26 -0.4 (4.6) | 1.1 (12.3) | 25 -0.5 (5.2) | 1.5 (16.1) |
| LI            | 36 -6.1 4.9 | 35 -5.5 5.4 | 36 -0.5 (5.3) | 1.0 (11.0) | 35 -0.3 (2.8) | 1.1 (12.2) |

*P value for food security by intervention interaction was 0.99 for weight change and 0.27 for HbA1c change in the model adjusted for age, gender, race/ethnicity, education, primary language, income, and health insurance.
†Participants (4.3%) who did not report income are excluded from these analyses; p value for income category by intervention interaction was 0.69 for weight change and 0.29 for HbA1c change in the model adjusted for age, gender, race/ethnicity, education, primary language, and health insurance.
HbA1c, hemoglobin A1c; LI, lifestyle intervention; MNT, medical nutrition therapy; NGSP, National Glycohemoglobin Standardization Program.
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than baseline in the lifestyle group at 6 months vs 0.3% (3.0 mmol/mol) lower in the MNT group (p=0.20). HbA1c was 0.3% (3.5 mmol/mol) lower with LI vs 0.4% (4.1 mmol/mol) lower with MNT at 12 months (p=0.63). In the LI group, change in HbA1c did not vary by food security status. HbA1c was 0.6% (6.0 mmol/mol) lower than baseline in food secure vs 0.3% (3.6 mmol/mol) lower in food insecure participants at 6 months. HbA1c was 0.3% (3.8 mmol/mol) lower in food secure vs 0.1% (1.5 mmol/mol) lower than food insecure at 12 months. The p values for the difference between food secure and food insecure participants were 0.26 in unadjusted models and 0.28 in adjusted models.

Those with lower income were more likely to be women and of racial/ethnic minorities, to have less education, and to be food insecure (online supplemental eTable 1). The intervention effect on the percentage of body-weight lost was also similar regardless of income category (table 2, figure 2). The intervention effect was nominally smaller in the lowest income category (3.3% lost for LI vs 1.0% lost for MNT in those with income <$30 000 a year, compared with the overall average of 5.6% lost vs 1.1% loss). However, these interactions were not statistically significant in either unadjusted (p=0.72) or adjusted (p=0.68) models. The intervention effect on HbA1c also did not significantly differ by income category (table 2).

Examining differences in attendance in LI sessions (online supplemental eTable 2), those who were food insecure attended approximately three fewer intervention sessions at both 6 and 12 months compared with food secure individuals, but this difference was not statistically significant at 6 (p=0.06) or 12 (p=0.17) months. The average attendance in LI in the food secure group was 14.8 out of 19 sessions in the first 6 months (14.8 hours of contact received relative to 19 hours planned) and 18.6 sessions out of 25 planned (18.6 hours received relative to 25 hours planned) at 12 months. For food insecure LI participants, the average attendance was 11.8 out of 19 sessions (11.8 hours received relative to 19 hours planned) in the first 6 months and 15.7 out of 25 sessions at 12 months (15.9 hours received relative to 25 hours planned). Optional sessions attended are described in online supplemental eTable 3. In the MNT arm, those with food insecurity attended 1.6 fewer visits (p=0.009) at 6 months and 2.9 fewer visits (p=0.0002) at 12 months than food secure participants. For food secure MNT participants, the mean attendance was 2.5 sessions (1.75 hours received) at 6 months and 4.0 sessions (2.5 hours received) at 12 months. For food insecure MNT participants, visit attendance was 0.9 sessions (0.9 hours received) at 6 months and 1.1 sessions (1.05 hours received) at 12 months. Visit attendance did not vary significantly by income in either arm (online supplemental eTable 2).

DISCUSSION

In this preplanned subgroup analysis of a pragmatic randomized trial, we found that the LI led to clinically meaningful weight loss in food insecure individuals. Further, although there were small variations in per cent weight lost by income category, the differences across groups were neither statistically significant nor clinically meaningful. There was no evidence that factors that

Figure 2 Percentage weight loss at 6 and 12 months by intervention arm (LI, lifestyle intervention; MNT, medical nutrition therapy) by category of annual income (k indicates thousands of dollars). P value for interaction term testing whether the intervention effect varies among subgroups was 0.72 in unadjusted models and 0.69 in models adjusted for age, gender, race/ethnicity, education, primary language, income, and health insurance.
may confound the relationship between food insecurity (or low income) and weight loss (eg, race/ethnicity or education) affected the results observed.

The major contribution of this study is to examine whether the effect of an LI for weight loss varied by food security status in individuals with type 2 diabetes. To our knowledge, whether an LI is similarly effective in those with, versus without, food insecurity has not previously been tested. However, there have been several prior studies addressing weight loss among low-income individuals with type 2 diabetes. A randomized controlled trial (RCT) that combined lifestyle and psychosocial intervention strategies in a low-income sample with diabetes found only modest weight loss (1.2 kg difference compared with usual care). A community-based diabetes self-management intervention for low-income African–American participants did not find changes in weight. This study did report improvements in HbA1c. African–American participants did not find changes in sample. Prior estimates of the patient population from anticipated prevalence of food insecurity in the included trials (RCT) that combined lifestyle and psychosocial intervention strategies in a low-income sample with diabetes found only modest weight loss (1.2 kg difference compared with usual care). A community-based diabetes self-management intervention for low-income African–American participants did not find changes in weight. This study did report improvements in HbA1c at 6 months, but these were not sustained at 12 or 18 months. A community health worker-led intervention for African–American women, primarily with lower income, found a modest difference in weight lost (1.35 kg lost vs 0.39 kg, respectively; p=0.046), using an educational mailings comparison group. A pilot RCT for African–American individuals with diabetes that used dietitian-led counseling sessions with weekly peer-support telephone calls, in a sample where 72% had an annual household income <$20,000, did not find a significant effect on weight loss. None of the studies mentioned examined whether intervention effect varied by food security status. Further, studies that are restricted to individuals with lower income are unable to determine whether intervention effects vary in lower, compared with higher, income groups.

Two other studies that did not focus on weight loss but did examine food insecurity are worth a discussion. A previous study of a diabetes self-management intervention, which included dietary advice but was not focused on weight loss, did find that food insecure participants showed similar (or larger) improvements in diabetes management self-efficacy and HbA1c, compared with food secure participants. A previous randomized trial of a food pantry-based intervention providing healthy food and diabetes management support for food insecure individuals found improvements in fruit and vegetable intake, but no change in HbA1c or weight.

In the REAL HEALTH trial, because of anticipated weight loss which could lead to hypoglycemia if glucose-lowering medications were not adjusted, trial protocols emphasized reduction of glucose-lowering medications when appropriate. Thus the lack of difference in glycemic control between MNT and LI was anticipated. For the purpose of this subgroup analysis, the key finding is that similar HbA1c reductions were observed regardless of food security status (or income category).

An important observation in this study is the lower than anticipated prevalence of food insecurity in the included sample. Prior estimates of the patient population from which the study sample was drawn, including a study in the same clinical network, indicate a food insecurity prevalence of greater than 20%, while only 13% of REAL HEALTH-Diabetes participants were food insecure. This suggests that there may be barriers to enrollment in LI among groups who experience food insecurity, including the length of the program and other competing demands, along with food insecurity itself. Food insecure individuals also attended approximately three fewer LI sessions, although this difference was not statistically significant, and attended significantly fewer MNT sessions. The benefits seen in this study, in which participants did not incur out-of-pocket costs for LI, suggest that if LI is more widely implemented it should be offered without cost-sharing requirements, as these are likely only to reduce participation, further increase the financial burden of diabetes, and lead to cost-related non-adherence and deterioration of glycemic control. Future work may address this barrier by combining interventions. For example, an intervention might combine directly addressing food insecurity (via food or medically tailored meal provision) with LI. By providing nutrition support, such a combined intervention may make the benefits of LI available to individuals who may otherwise not feel able to commit to lifestyle change. At the same time, the skills learned during LI could help sustain dietary modification as individuals transition off of food support interventions.

The findings of this study have important implications for diabetes management among vulnerable patients. Because this LI was effective regardless of food security status or income category, it may be offered in clinics, such as health centers, that serve populations with low income and/or high food insecurity prevalence. Because LI is an effective alternative to MNT (which is often a benefit covered by health insurance), state Medicaid programs and private health insurers should consider adding LIs as covered benefits—much as Medicare has recently done in allowing coverage for DPP-derived services in certain contexts. Given the ease of administration of the telephonic intervention, such an approach may be a particularly attractive strategy among individuals with transportation barriers or schedules that preclude inperson sessions.

The results of this study should be interpreted in light of several limitations. Although this was a prespecified subgroup analysis, the study did not randomize on the basis of food insecurity or income. Nevertheless, since treatment assignment was made without regard to food security status, food security status cannot confound the comparisons made between the LI and MNT group, within levels of food security status. Further, adjustment for known confounders did not change the finding of no effect modification. Next, the LI was offered to participants without cost sharing, whereas MNT was not covered as part of the study, and therefore copays and deductibles could have precluded some food insecure participants from receiving the recommended efficacious dose of MNT. However, this is unlikely to have affected
the study results as food secure individuals, who should face fewer financial barriers to MNT, saw weight loss in the MNT arm that was similar to that seen in food insecure participants. Next, as noted above, the prevalence of food insecurity was lower in the study sample than would be expected in the background population. Given the effect estimates were not greater than a prespecified clinically meaningful threshold across the subgroups, the observation of similar effects regardless of food insecurity status (or income level) is not likely to be affected by low power, but this type of selection into the trial may reduce the generalizability of the results. Specifically, those who, owing to food insecurity, felt incapable of committing to an LI and thus did not enroll may have experienced less intervention benefit if they had enrolled. Next, we do not present information on possible mediators of intervention effect, such as gain of knowledge or changes in attitudes or behaviors. That will be an important direction for future work. Another limitation is that, owing to the small sample of food insecure individuals, we did not have sufficient power to stratify analyses by LI modality (telephone vs inperson). Finally, this study was conducted in a single clinical network in eastern Massachusetts; thus, results may not be generalizable throughout the USA. The limitations are balanced by several strengths. This was a pragmatic but rigorous randomized trial of an LI adapted for group implementation in primary care and community health centers. The intervention was conducted in English and Spanish, and the subgroup analyses reported here were prespecified.

We found that an LI was effective for weight loss in individuals with type 2 diabetes and food insecurity and/or low income who enrolled in the program. However, barriers to enrolling in LI remain a concern. New research that combines the ability to support those experiencing food insecurity in making lifestyle changes that modify their disease trajectory is an important next step that may help us to reduce socioeconomic disparities in diabetes.

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Competing interests DJW reports serving on a data monitoring committee for Novo Nordisk. LMD serves on the advisory boards for Omada Health, Jana Care, and WW International. SJC reports employment of a close family member by Johnson & Johnson.

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