

# Combined Diet and Physical Activity Promotion Programs to Prevent Type 2 Diabetes Among Persons at Increased Risk: A Systematic Review for the Community Preventive Services Task Force

Ethan M. Balk, MD, MPH; Amy Earley, BS; Gowri Raman, MD, MS; Esther A. Avendano, BA; Anastassios G. Pittas, MD, MS; and Patrick L. Remington, MD, MPH

**Background:** Trials have shown efficacy of rigorous diet and physical activity promotion programs to reduce diabetes incidence and improve glycemic measures in adults at increased risk for type 2 diabetes.

**Purpose:** To evaluate diet and physical activity promotion programs for persons at increased risk for type 2 diabetes, primarily to reduce diabetes risk and decrease body weight and glycemia.

**Data Sources:** MEDLINE, the Cochrane Central Register of Controlled Trials, CAB Abstracts, Global Health, and Ovid HealthSTAR from 1991 through 27 February 2015, with no language restriction.

**Study Selection:** 8 researchers screened articles for single-group or comparative studies of combined diet and physical activity promotion programs with at least 2 sessions over at least 3 months in participants at increased risk for type 2 diabetes.

**Data Extraction:** 7 researchers extracted data on study design; participant, intervention, and outcome descriptions; and results and assessed study quality.

**Data Synthesis:** 53 studies (30 of diet and physical activity promotion programs vs. usual care, 13 of more intensive vs. less intensive programs, and 13 of single programs) evaluated 66

programs. Compared with usual care, diet and physical activity promotion programs reduced type 2 diabetes incidence (risk ratio [RR], 0.59 [95% CI, 0.52 to 0.66]) (16 studies), decreased body weight (net change,  $-2.2\%$  [CI,  $-2.9\%$  to  $-1.4\%$ ]) (24 studies) and fasting blood glucose level (net change,  $-0.12$  mmol/L [ $-2.2$  mg/dL] [CI,  $-0.20$  to  $-0.05$  mmol/L [ $-3.6$  to  $-0.9$  mg/dL]]) (17 studies), and improved other cardiometabolic risk factors. Evidence for clinical events was limited. More intensive programs were more effective.

**Limitations:** Wide variation in diet and physical activity promotion programs limited identification of features most relevant to effectiveness. Evidence on clinical outcomes and in children was sparse.

**Conclusion:** Combined diet and physical activity promotion programs are effective at decreasing diabetes incidence and improving cardiometabolic risk factors in persons at increased risk. More intensive programs are more effective.

**Primary Funding Source:** Centers for Disease Control and Prevention Community Preventive Services Task Force.

*Ann Intern Med.* 2015;163:437-451. doi:10.7326/M15-0452 [www.annals.org](http://www.annals.org)  
For author affiliations, see end of text.

This article was published online first at [www.annals.org](http://www.annals.org) on 14 July 2015.

Diabetes is a large and growing medical problem, and the costs to society are high and escalating. According to the latest figures from the Centers for Disease Control and Prevention (CDC), 29.1 million persons (9.3% of the U.S. population) have diabetes, and 1.7 million new cases are diagnosed annually (1). Worldwide, an estimated 387 million adults are living with diabetes, and this number is projected to increase to 592 million by 2035 (2). Prevalence of diabetes and related costs are expected to more than double in the next 25 years (3), given that in excess of 86 million Americans (37% of the adult population) are at risk for the disease (1). Effective prevention strategies are, therefore, crucial to slow the diabetes tide and its associated burden.

Nearly 9 out of 10 new diabetes cases are type 2 diabetes, which has a natural history characterized by a gradual increase in glycemia. Identification of persons at increased risk can enable the implementation of interventions to decrease the risk for progression to clinical diabetes. The American Diabetes Association has defined prediabetes as a high-risk category based on a glycemic level that does not meet criteria for diabetes but is too high to be considered normal (4). Persons with prediabetes progress to type 2 diabetes at a rate of about 5% to 10% per year without intervention (5).

Three large clinical trials from the United States (6), Finland (7), and China (8) have shown that the primary components of diabetes prevention in adults are weight loss and increased physical activity. In these trials, among persons at risk for type 2 diabetes, rigorous application of combined diet and physical activity promotion programs, with the goals of weight loss and increased physical activity, reduced risk for diabetes by 50% to 60% during the active intervention period (3 to 6 years). Although attenuated, the effect of the intervention can persist in the long term (9-11). The results of these trials are well-known; however, wide-scale implementation of combined diet and physical activity promotion programs in clinical and community-based settings has only recently begun and requires further expansion (12).

## See also:

Related articles . . . . . 452, 465

Editorial comment . . . . . 475

Web-Only  
Supplement

Combined diet and physical activity promotion programs aim to prevent type 2 diabetes among persons who are at increased risk for the disease. These programs actively encourage persons to improve their diet and increase physical activity by using trained providers in various settings who work with clients for at least 3 months, providing some combination of counseling, coaching, and extended support in multiple sessions (delivered in person or by other methods) related to diet and physical activity. Programs may also include many other features, including specialized counselors; a range in the number and frequency of sessions; different session types; and different diet, weight-loss, or exercise goals.

The purpose of this review was to assess the effectiveness of diet and physical activity promotion programs implemented in a wide range of clinical or community settings to reduce risk for new-onset diabetes among adults and children at risk for type 2 diabetes. The Community Preventive Services Task Force (Task Force) ([www.thecommunityguide.org](http://www.thecommunityguide.org)) used this review to update its guidance on diabetes prevention and to identify gaps in the evidence to inform future research. Potential effect modifiers, such as intensity and specificity of the programs, settings, and implementers, were evaluated. Furthermore, the potential benefit of the diabetes prevention programs extending to other cardiometabolic risk factors, such as overweight, high cholesterol level, and high blood pressure (BP), was also assessed.

## METHODS

This review was conducted in accordance with the methods of the Task Force (13, 14) and the highest standards for conducting systematic reviews (15, 16). We convened a panel of domain experts and stakeholders (Coordination Team) that, together with our Community Guide Technical Monitor and Task Force members, provided input on the protocol, feedback on the findings, conclusions, and evidence gaps.

### Data Sources

We searched MEDLINE, the Cochrane Central Register of Controlled Trials, CAB Abstracts, Global Health, and Ovid HealthSTAR from 1991 through 27 February 2015 with no language restrictions. Table 1 of the Supplement (available at [www.annals.org](http://www.annals.org)) shows the search strategy. We also screened reference lists of related systematic and narrative reviews and suggestions from the expert panel.

### Study Selection

We included randomized, controlled trials and prospective nonrandomized comparative studies with at least 30 participants per group, as well as prospective single-group intervention studies with at least 100 participants. The population of interest was adults or children at increased risk for type 2 diabetes (that is, with prediabetes) as determined by glycemic measures or diabetes risk assessment tools. We included studies of participants with the metabolic syndrome (who are at

increased risk for *both* diabetes and cardiovascular disease) and studies with participants who were chosen because they were at risk for *either* type 2 diabetes or cardiovascular disease. However, we excluded studies of participants with established type 2 diabetes or whose only risk factor was obesity or increased risk for cardiovascular disease (without explicit inclusion of participants with prediabetes). The implied or explicit intent of the diet and physical activity promotion programs had to be to prevent diabetes, and the programs had to include at least 2 contact sessions (in-person or virtual) over at least 3 months. Programs had to include both dietary and increased physical activity components and could be conducted in any outpatient setting. We allowed any type of advice to improve diet and increase physical activity (except for single-food or supplement dietary changes, such as addition of fish oil). We excluded interventions that included antidiabetic medications. The comparative studies had to include a usual care group (no active diet and physical activity promotion program) or a lower-intensity diet and physical activity promotion program (for example, one with fewer contact sessions or a more liberal diet). We required at least 6 months of follow-up for any of the following outcomes: incident diabetes, reversion to normoglycemia, body weight, glycemic measures (fasting glucose level, 2-hour glucose level after a 75-g oral glucose tolerance test, or hemoglobin A<sub>1c</sub> [HbA<sub>1c</sub>] level), all-cause death, diabetes-related clinical outcomes (such as cardiovascular events, end-stage renal disease, nephropathy, amputation, retinopathy, neuropathy, skin ulcers, or periodontitis), BP, and lipid levels (total, low-density lipoprotein [LDL], and high-density lipoprotein [HDL] cholesterol and triglycerides).

### Data Extraction and Quality Assessment

We screened titles and abstracts using Abstrackr (17). Eight researchers double-screened the abstracts after iterative training of all reviewers on the same batches of abstracts. Discordant decisions and queries were resolved at group meetings. Full-text articles were retrieved for all potentially relevant abstracts and re-screened by the same researchers.

Data from each study were extracted by 1 of 7 experienced methodologists and confirmed by a senior methodologist; the same methodologists assessed study quality. Data extraction was conducted in the Systematic Review Data Repository (18) and included elements for study design, including eligibility criteria, population characteristics, detailed descriptions of the diet and physical activity promotion programs and comparison interventions, outcomes, and results. We assessed the quality of each study by using 12 Community Guide quality-of-execution questions (see the footnotes of Table 2 of the Supplement, available at [www.annals.org](http://www.annals.org)) (14, 19). Per Community Guide protocol, we excluded studies with "limited quality of execution," defined as those with at least 5 major limitations.

### Data Synthesis and Analysis

All extracted data were placed into summary evidence tables (available in the supporting materials at

www.thecommunityguide.org/diabetes/combineddietandpa.html). Two studies that were conducted in children were not included in the meta-analyses and are reported separately. For outcomes with data from at least 3 comparative studies of diet and physical activity promotion versus usual care, we performed meta-analysis of the risk ratio (RR) or net change (20) using a profile likelihood random-effects model. For nonrandomized studies, we preferentially used results of adjusted analyses. Meta-analyses were conducted with the metaan package in Stata 13.1 (StataCorp). For the overall meta-analyses of incident diabetes and reversion to normoglycemia, we used data from the longest reported follow-up. For continuous outcomes, we used data closest to 1 year of follow-up and those from the longest follow-up. We evaluated differences in effect (for incident diabetes and weight only) using direct comparisons of different diet and physical activity promotion programs within studies, reported within-study subgroup analyses, and across-study metaregression (based on predetermined study setting and program features and using a random-effects model) across all programs. Incident diabetes and weight change were chosen for metaregression because of their relative importance in determining the effectiveness of diet and physical activity promotion programs. Metaregressions were conducted with the metareg package in Stata and were considered potentially significant if the *P* value was less than 0.10. For each outcome with at least 10 studies, we examined the possibility of publication bias with funnel plots and the Harbord test (for diabetes incidence) or the Egger test (for continuous outcomes) using the metabias and metafunnel packages in Stata (21).

### Role of the Funding Source

One member of the Coordination Team and our Technical Monitor are employed by the CDC; none of the Task Force members are. The Coordination Team, the Technical Monitor, and members of the Task Force participated in the formulation of the study questions and the development of the protocol but did not participate in the literature search, the determination of study eligibility, or data analysis or interpretation. The Coordination Team, the Technical Monitor, and CDC personnel were given an opportunity to provide feedback on the manuscript and the decision to submit the manuscript for publication. However, the research team retained final determination of the content and the decision to publish the manuscript.

## RESULTS

Appendix Figure 1 (available at [www.annals.org](http://www.annals.org)) summarizes the search yield. Of 11 317 citations (plus articles from existing systematic reviews and suggestions from domain experts), 53 studies described 66 diet and physical activity promotion programs in 104 articles (6–11, 22–119). One additional study with 6 major limitations was excluded because of limited quality of execution (120). The included studies described 26

randomized and 4 nonrandomized comparisons of diet and physical activity promotion programs versus usual care, 12 randomized and 1 nonrandomized comparisons of 2 or more diet and physical activity promotion programs (3 of which also had usual care groups), and 13 single-group evaluations of diet and physical activity promotion programs. Thirty-three studies were of good quality (0 or 1 limitation), and 20 were of fair quality (2 to 4 limitations) (Table 2 of the Supplement). The most common limitations were poor descriptions of the study populations or intervention programs, problems with data measurement or interpretation, and high dropout rates. Although half of the studies (*n* = 27) analyzed all enrolled participants, 9 had rates of dropout or loss to follow-up greater than 20%.

The characteristics of the diet and physical activity promotion programs are summarized in Table 1, and details are provided in Tables 3 to 5 of the Supplement (available at [www.annals.org](http://www.annals.org)). All but 5 programs (in 4 studies) lasted at least 6 months. Programs offered a wide range of number of contact sessions (0 [virtual contacts only] to 72; median, 15), and most included both a core period (with frequent contact sessions) and a maintenance period (with less frequent contact). Except for 7 programs that were delivered entirely over the Internet or by video, telephone, or e-mail, programs used in-person individual or group sessions (or both) on diet or exercise (or both). Sessions were led by different combinations of trained diet counselors, including dietitians or nutritionists (among others); trained exercise counselors, including physical trainers (among others); nurses; physicians or psychologists; or trained laypersons. Many programs included specific weight-loss, diet, or physical activity goals (Table 1). Some included individually tailored plans for diet and physical activity.

Table 2 summarizes the participant characteristics, with details provided in Table 6 of the Supplement (available at [www.annals.org](http://www.annals.org)). Thirty (57%) studies were restricted to participants with prediabetes, of which 21 used standard diagnostic criteria; 12 (23%) studies included only participants at increased risk for diabetes on the basis of a risk score. More than three quarters of the studies included mostly overweight or obese participants, and most study participants were female and at least middle-aged. Two studies were conducted in adolescents at increased risk for type 2 diabetes; these studies were analyzed separately. None of the studies reported any long-term harms directly related to the diet and physical activity promotion programs.

### Incident Diabetes

Sixteen studies that compared diet and physical activity promotion programs versus usual care reported new-onset diabetes (6–9, 22–33); 2 studies each compared 2 programs with usual care. All but 3 were randomized trials (9, 22, 26). Incident diabetes was reported between 1 and 23 years from the start of the programs (Figure 1). Across studies, 0% (at 1 year) to 73% (at 23 years) of program participants developed diabetes. At all time points, program participants were

**Table 1.** Characteristics of Combined Diet and Physical Activity Promotion Programs

| Characteristic, by Category                                   | Value                |
|---|----------------------|
| <b>Median sessions (66 programs) (IQR; range), n</b>          |                      |
| Core  | 10 (6-16; 0*-72)     |
| Maintenance†  | 6 (1.5-12; 0*-24)    |
| Total   | 15 (6.5-24.5; 0*-72) |
| <b>Median program duration (66 programs) (IQR; range), mo</b> |                      |
| Core  | 6 (5-12; 1-60)       |
| Maintenance†  | 12 (7-18; 4-68)      |
| Total   | 12 (10-27; 3-72)     |
| <b>Program design (66 programs), n (%)‡</b>                   |                      |
| Nominally based on DPP or DPS                                 | 27 (41)              |
| <b>Weight-loss goal (66 programs), n (%)‡</b>                 |                      |
|   | 42 (64)              |
| <b>Diet intervention (66 programs), n (%)‡</b>                |                      |
| Individual sessions   | 40 (61)              |
| Group sessions  | 41 (62)              |
| Individual and group sessions                                 | 24 (36)              |
| Individually tailored diet plan                               | 16 (24)              |
| Diet goal   | 19 (29)              |
| Diet counselor  | 29 (44)              |
| <b>Physical activity intervention (67 programs), n (%)‡</b>   |                      |
| Individual sessions   | 41 (62)              |
| Group sessions  | 39 (59)              |
| Individual and group sessions                                 | 24 (36)              |
| Individually tailored exercise plan                           | 23 (35)              |
| Exercise goal   | 32 (48)              |
| Exercise counselor  | 18 (27)              |
| <b>Counselors (51 programs), n (%)‡</b>                       |                      |
| Dietitian   | 37 (73)              |
| Exercise therapist  | 26 (51)              |
| Nurse   | 15 (29)              |
| Layperson   | 13 (25)              |
| Physician   | 8 (16)               |
| Diabetes educator   | 3 (6)                |
| <b>Country (53 studies), n (%)</b>                            |                      |
| United States/Canada  | 22 (42)              |
| Western Europe/Australia                                      | 22 (42)              |
| Japan   | 3 (6)                |
| Middle-income§  | 6 (11)               |
| <b>Setting (41 studies), n (%)</b>                            |                      |
| Community   | 12 (29)              |
| Health care system  | 25 (61)              |
| Worksite  | 0 (0)                |
| Multiple  | 4 (10)               |
| <b>Location (53 studies), n (%)</b>                           |                      |
| Urban   | 25 (47)              |
| Regional  | 21 (40)              |
| Suburban  | 2 (4)                |
| Rural   | 1 (2)                |
| Mixed   | 4 (8)                |

DPP = Diabetes Prevention Program; DPS = Diabetes Prevention Study; IQR = interquartile interval.

\* In some programs, the contacts were by telephone, e-mail, Internet, or video only.

† 28 programs.

‡ Probably underestimated because of inadequate or unclear reporting in articles.

§ India (n = 3), Brazil (n = 1), China (n = 1), and Pakistan (n = 1).

**Table 2.** Characteristics of Study Participants

| Characteristic, by Category                                     | Value                       |
|---|-----------------------------|
| <b>Studies meeting eligibility criteria (53 studies), n (%)</b> |                             |
| Prediabetes, IGT, or IFG  | 30 (57)                     |
| By ADA/WHO criteria   | 21 (40)                     |
| At increased risk for diabetes (by risk score)                  | 12 (23)                     |
| Prediabetes or at increased risk for diabetes                   | 4 (8)                       |
| Prediabetes or at increased risk for cardiovascular disease     | 4 (8)                       |
| Metabolic syndrome, with or without prediabetes                 | 3 (6)                       |
| <b>Body weight (47 studies)</b>                                 |                             |
| Median of mean BMI (IQR; range), kg/m <sup>2</sup>              | 31.2 (28.1-33.6; 23.8-39.7) |
| <b>Hypertension (4 studies)</b>                                 |                             |
| Median participants (range), %                                  | 34.5 (30.6-50.0)            |
| <b>Female sex (39 studies)</b>                                  |                             |
| Median participants (IQR; range), %                             | 65.3 (50.3-73.9; 13.5-90.5) |
| <b>Age (39 studies)*</b>  |                             |
| Median of mean age (IQR; range), y                              | 53.6 (48-57; 43.1-65.0)     |
| <b>Median ethnicity, %†</b>                                     |                             |
| White (13 studies) (range)                                      | 74 (18-89)                  |
| Black/African American (10 studies) (range)                     | 18 (12-39)                  |
| Hispanic/Latino (8 studies) (range)                             | 13 (3-38)                   |
| East Asian (5 studies)  | 100                         |
| Southeast Asian (6 studies)                                     | 100                         |
| Asian/Pacific Islander (4 studies)                              | 4, 5, 15, and 17‡           |
| Native American (4 studies)                                     | 1, 3, 6, and 100‡           |
| <b>Median education level, %</b>                                |                             |
| Less than high school or equivalent (9 studies) (IQR; range)    | 14 (11-33; 5-64)            |
| High school or some college (20 studies) (IQR; range)           | 30 (21-48; 10-69)           |
| Bachelor's degree or equivalent (11 studies) (IQR; range)       | 28 (20-37; 14-52)           |
| Graduate degree or equivalent (4 studies)                       | 13, 15, 16, and 35‡         |

ADA = American Diabetes Association; BMI = body mass index; IFG = impaired fasting glucose; IGT = impaired glucose tolerance; IQR = interquartile interval; WHO = World Health Organization.

\* Excludes 2 studies in adolescents.

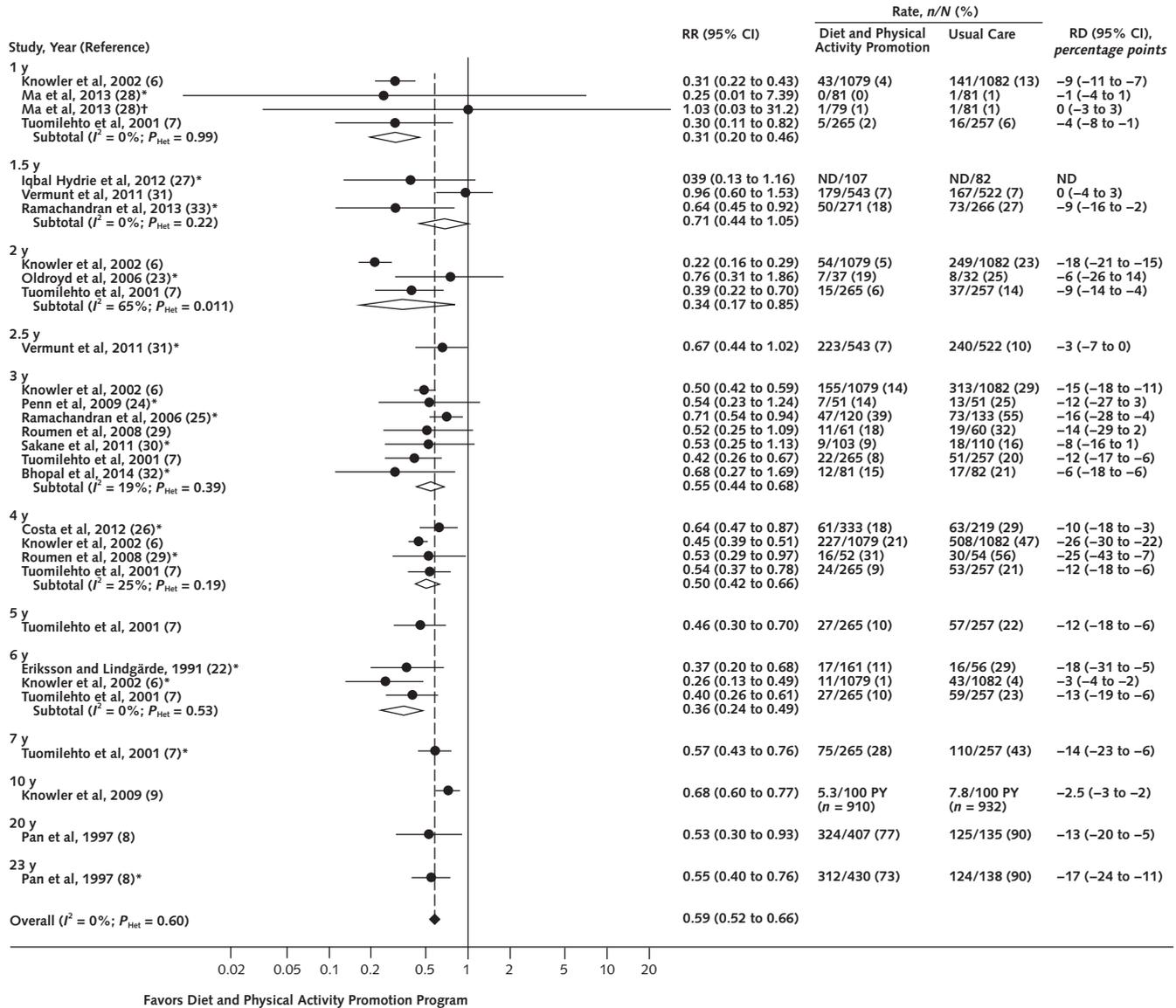
† Excludes studies with 0% of an ethnicity.

‡ Percentages among relevant studies.

less likely to develop diabetes. Across all studies, the summary RR for incident diabetes was 0.59 (95% CI, 0.52 to 0.66), with no statistical heterogeneity. The median risk difference across studies was -11 percentage points (interquartile interval, -16 to -5 percentage points). Funnel plot analysis did not find different effects between larger and smaller studies (Harbord test *P* = 0.27).

Both the U.S. DPP (Diabetes Prevention Program) study (6) and the Finnish DPS (Diabetes Prevention Study) (7) found statistically significantly larger effects in older participants, but although the latter found a non-significant effect in the youngest age group (<51 years), the former found statistically significant effects in all age groups. Neither study found differences by sex. The DPP found no difference by race or ethnicity, and

**Figure 1.** Random-effects model meta-analysis of RR of incident diabetes in at-risk participants in combined diet and physical activity promotion programs versus usual care.



The meta-analysis of the overall RR (black diamond) used data from the longest follow-up from each study, as indicated by the asterisks. Subgroup meta-analyses by follow-up time (open diamonds) were conducted for time points with data from ≥3 studies. ND = no data; P<sub>Het</sub> = chi-square P value of heterogeneity; PY = person-year; RD = risk difference; RR = risk ratio.

\* Included in overall meta-analysis.

† To avoid biased meta-analyses due to inclusion of correlated analyses, this comparison between the lower-intensity intervention and control was excluded.

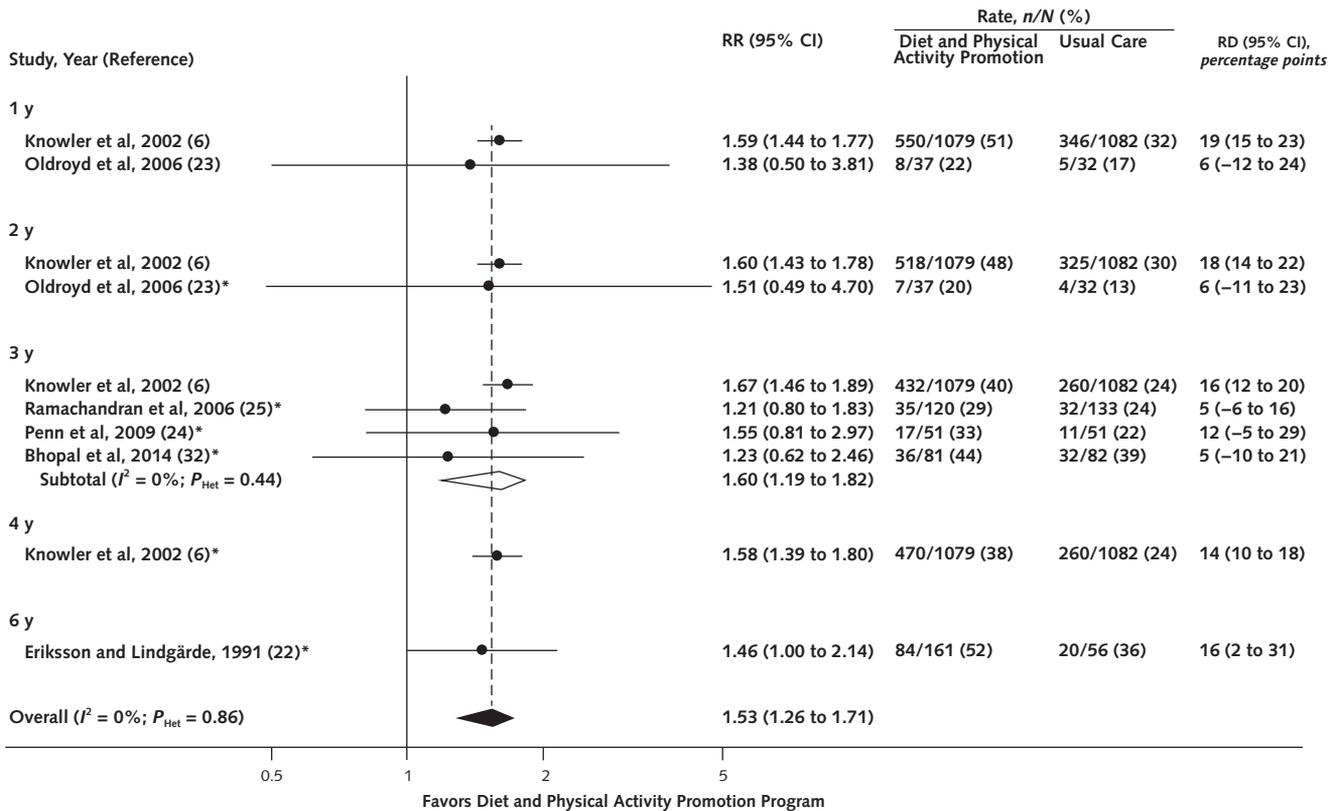
the DPS found no difference by educational attainment. The JDPP (Japan Diabetes Prevention Program) study reported a significant effect of diet and physical activity promotion programs among participants with baseline HbA<sub>1c</sub> levels of at least 5.7% compared with those with lower levels, but it did not provide a statistical analysis of the difference between subgroups (34).

No significant differences across studies were found by setting; number of sessions; program duration; whether the program was based on the DPP or DPS approach; or inclusion of a weight-loss goal, indi-

vidual or group diet or exercise sessions (analyzed separately), individually tailored diet plans, or diet or exercise counselors (analyzed separately). The 11 programs that included an individually tailored exercise plan (RR, 0.53 [CI, 0.45 to 0.63]) may have had a greater effect than the 5 that did not (RR, 0.67 [CI, 0.55 to 0.81]) (P = 0.070 for interaction).

Six studies directly compared more intensive versus less intensive diet and physical activity promotion programs (28, 45, 47, 48, 50, 56). Compared with less intensive programs, more intensive programs had

**Figure 2.** Random-effects model meta-analysis of RR of reversion to normoglycemia in at-risk participants in combined diet and physical activity promotion programs versus usual care.



The meta-analysis of the overall RR (black diamond) used data from the longest follow-up from each study, as indicated by the asterisks. Subgroup meta-analysis by follow-up time (open diamond) was conducted for the time point with data from >3 studies.  $P_{\text{Het}}$  = chi-square  $P$  value of heterogeneity; RD = risk difference; RR = risk ratio.

\* Included in overall meta-analysis.

more sessions (4 studies); weight-loss, diet, or exercise goals (3 studies); or a maintenance phase, more intensive diet and exercise plans, an exercise physiologist, individual contact sessions, or in-person (vs. DVD) sessions (1 study each). All 5 studies that reported at least 1 case of incident diabetes found lower incidence with a more intensive program (RR, 0.28 to 0.56), but this was statistically significant in only 1 study (50) (Appendix Figure 2, available at [www.annals.org](http://www.annals.org)).

### Reversion to Normoglycemia

Six studies (5 trials and 1 nonrandomized study) that compared diet and physical activity promotion programs versus usual care reported reversion to normoglycemia as early as 1 year from the start of the intervention (Figure 2) (6, 22-25, 32). Across studies, between 20% (at 2 years) and 52% (at 6 years) of program participants reverted to normoglycemia. At 3 years (4 studies) and across time points, the summary RRs for achievement of normoglycemia were statistically significant, with an overall summary RR of 1.53 (CI, 1.26 to 1.71) and no statistical heterogeneity. The median risk difference across studies was 12 percentage points (interquartile interval, 6 to 14 percentage points). No within-study subgroup differences were reported, and

no between-study subgroup differences were found. Three studies that directly compared more intensive versus less intensive programs (45, 47, 48) found effects favoring more intensive programs (RR, 1.58 to 2.11), 2 of which were statistically significant (47, 48) (Appendix Figure 3, available at [www.annals.org](http://www.annals.org)).

### Clinical Events

Three long-term studies reported all-cause mortality, 2 of which also reported cardiovascular mortality with no consistent pattern of results. The Da Qing study reported lower risk for all-cause death (hazard ratio [HR], 0.71 [CI, 0.51 to 0.99]) with diet and physical activity promotion after 23 years (10), but this effect was restricted to women and was not significant at earlier time points (HRs, 1.33 at 6 years and 0.96 at 20 years) (8). Knowler and colleagues (DPP study) (6) found no effect at 3 years (risk difference, -0.6 per 1000 person-years), and Uusitupa and coworkers (DPS) found no effect at 10 years (HR, 0.57 [CI, 0.21 to 1.58]) (105). Similar results were found for cardiovascular death, with significantly lower risk in the Da Qing study (HR, 0.59 [CI, 0.36 to 0.96]) at 23 years (10); this effect also was restricted to women and was not significant at earlier time points. The DPS found no significant effect on car-

diovascular death at 3 years (RR, 0.50 [CI, 0.09 to 2.73]) (105). The Da Qing study reported a reduction in severe retinopathy at the 20-year follow-up (HR, 0.53 [CI, 0.29 to 0.99]) (71). Limited evidence suggested no significant effects on other clinical outcomes, including cardiovascular events (78, 95, 105), nephropathy (71), and neuropathy (71), often due to a lack of power.

### Body Weight and Glycemia

The 24 studies that compared diet and physical activity promotion programs versus usual care and reported weight change all found net weight loss with diet and physical activity promotion (6, 7, 9, 22-24, 27-33, 35-41, 52-55), ranging from  $-0.2\%$  to  $-10.5\%$  of initial body weight (summary net change,  $-2.2\%$  [CI,  $-2.9\%$  to  $-1.4\%$ ]); however, the studies had high statistical heterogeneity ( $I^2 = 89\%$ ;  $P < 0.001$ ) (Figure 3). Funnel plot analysis did not find different effects between larger and smaller studies (Egger test  $P = 0.51$ ). We used metaregression to test the same covariables examined for incident diabetes, and the only variable for which effects differed across studies was whether programs were based on the DPP or the DPS approach. The 12 programs based on either approach yielded a net change of  $-3.0\%$  (CI,  $-4.1\%$  to  $-1.9\%$ ) compared with  $-1.6\%$  (CI,  $-2.5\%$  to  $-0.6\%$ ) for the 13 other programs ( $P = 0.051$  for interaction). However, heterogeneity across studies remained high (residual  $I^2 = 95\%$ ). Across all 42 programs (not compared with usual care) (6, 7, 22, 23, 27-33, 35-51, 54-58), none of the factors explored by metaregression yielded statistically significant differences across studies. In contrast to the across-study analysis, 6 of the 10 studies that directly compared more intensive versus less intensive programs found statistically significantly greater weight loss with the more intensive programs (28, 35, 44, 45, 47-50, 56, 58) (Appendix Figure 4, available at [www.annals.org](http://www.annals.org)).

Eighteen studies that compared diet and physical activity promotion programs versus usual care reported glycemic outcomes (6-9, 23, 28-32, 35-40, 52, 53). Overall, such programs improved measures of glycemia. Across studies, at follow-up durations closest to 1 year, fasting glucose level had a summary net change of  $-0.12$  mmol/L ( $-2.2$  mg/dL) (CI,  $-0.20$  to  $-0.05$  mmol/L [ $-3.6$  to  $-0.9$  mg/dL]) (17 studies;  $I^2 = 77\%$ ), 2-hour glucose level improved by  $-0.48$  mmol/L ( $-8.6$  mg/dL) (CI,  $-0.86$  to  $-0.17$  mmol/L [ $-15.5$  to  $-3.1$  mg/dL]) (11 studies;  $I^2 = 87\%$ ), and HbA<sub>1c</sub> level improved by  $-0.08\%$  (CI,  $-0.12\%$  to  $-0.04\%$ ) (8 studies;  $I^2 = 0\%$ ) (Table 7 of the Supplement, available at [www.annals.org](http://www.annals.org)). Funnel plot analysis found no significant small-study effect for fasting glucose level (Egger test  $P = 0.54$ ), but smaller studies were more likely to have large net reductions in 2-hour glucose level ( $P = 0.003$ ). However, studies reporting significant effects on fasting glucose level were no more likely to report 2-hour glucose results than those with nonsignificant effects ( $P = 0.21$ ). Across 8 studies that compared more intensive versus less intensive programs (28, 43-45, 48-50, 56) (Table 8 of the Supplement, available at [www.annals.org](http://www.annals.org)),

the median net change in fasting glucose level was  $-0.11$  mmol/L ( $-2.0$  mg/dL) (range,  $-0.20$  to  $0.17$  mmol/L [ $-3.6$  to  $3.0$  mg/dL]), favoring more intensive programs; however, the difference was statistically significant in only 1 study (56). Among 4 studies (44, 45, 48, 50), the median net change in 2-hour glucose level was  $-0.37$  mmol/L ( $-6.7$  mg/dL) (range,  $-0.6$  to  $-0.2$  mmol/L [ $-11$  to  $-3.6$  mg/dL]), favoring more intensive programs; the difference was significant in 2 studies (48, 50). None of these studies reported on HbA<sub>1c</sub> level.

Across the 31 diet and physical activity promotion programs (not compared with usual care) in 24 studies that reported on fasting glucose level (6-9, 23, 28-32, 36-39, 43-46, 48-50, 52, 53, 56), results differed on the basis of whether individual diet sessions and diet counselors were included. After adjustment for follow-up duration, programs with individual diet sessions ( $n = 25$  of 31) or diet counselors ( $n = 22$  of 31) yielded larger decrements in fasting glucose level (individual sessions:  $-0.24$  vs.  $-0.02$  mmol/L [ $-4.3$  vs.  $-0.4$  mg/dL] [ $P = 0.020$ ]; counselors:  $-0.25$  vs.  $-0.07$  mmol/L [ $-4.5$  vs.  $-1.3$  mg/dL] [ $P = 0.034$ ]).

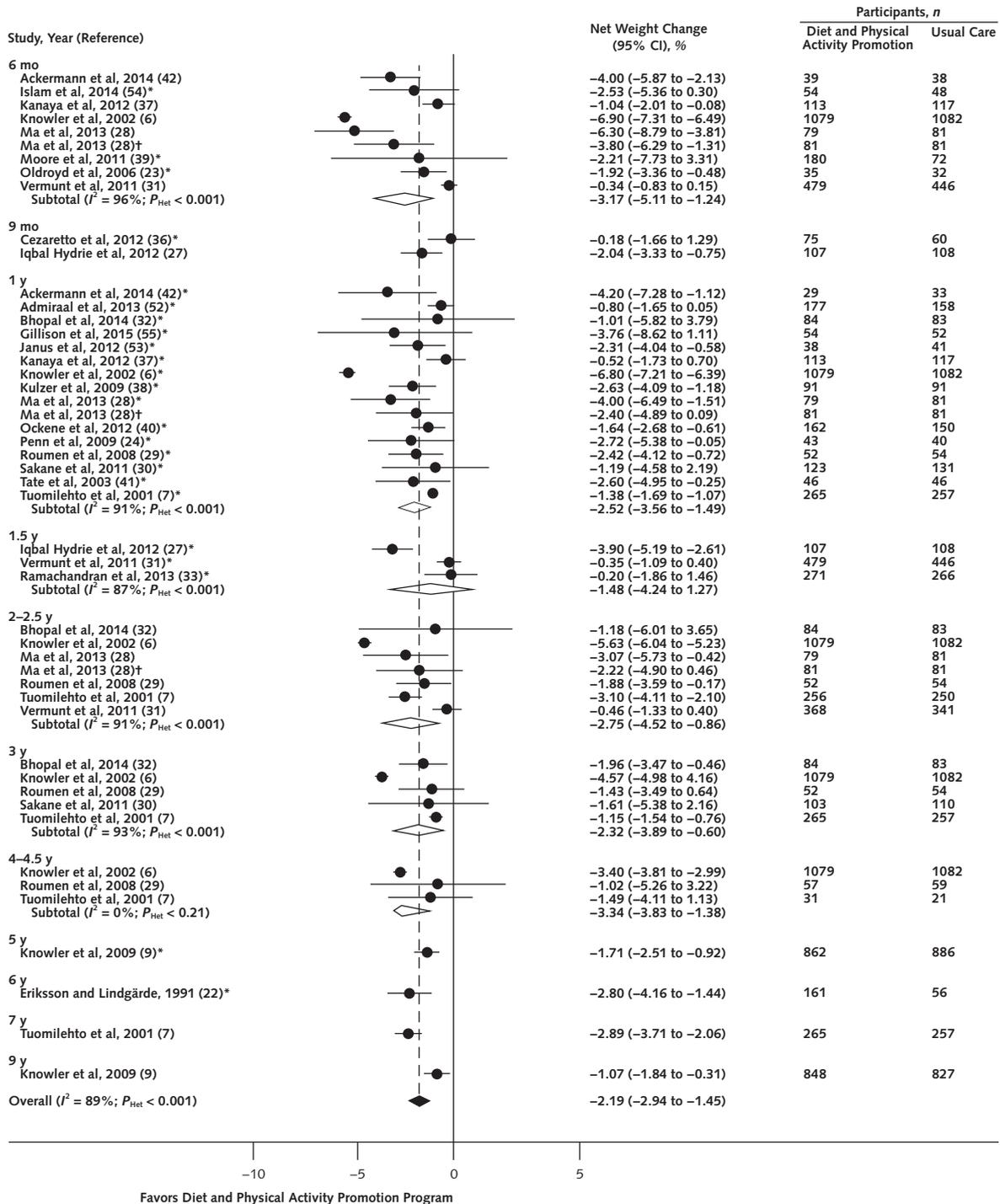
### Blood Pressure and Lipid Levels

Across 17 studies comparing diet and physical activity promotion programs versus usual care (6, 7, 9, 22, 23, 28, 29, 32, 33, 35-39, 52-54), at follow-up durations closest to 1 year, the programs improved systolic BP (net change,  $-1.6$  mm Hg [CI,  $-2.7$  to  $-0.5$  mm Hg];  $I^2 = 45\%$ ) and diastolic BP (net change,  $-1.6$  mm Hg [CI,  $-2.5$  to  $-0.8$  mm Hg];  $I^2 = 73\%$ ) (Table 9 of the Supplement, available at [www.annals.org](http://www.annals.org)). We found no evidence of small-study effects (Egger test  $P = 0.51$  for systolic BP and 0.83 for diastolic BP). Across 14 studies (7, 9, 22, 23, 28, 29, 33, 35-39, 52, 53), the programs also statistically significantly improved total cholesterol levels (net change,  $-0.05$  mmol/L [ $-1.8$  mg/dL] [CI,  $-0.12$  to  $-0.002$  mmol/L [ $-4.6$  to  $-0.1$  mg/dL]]) (12 studies;  $I^2 = 0\%$ ), LDL cholesterol levels (net change,  $-0.09$  mmol/L [ $-3.3$  mg/dL] [CI,  $-0.17$  to  $-0.01$  mmol/L [ $-6.4$  to  $-0.3$  mg/dL]]) (8 studies;  $I^2 = 0\%$ ), HDL cholesterol levels (net change,  $0.03$  mmol/L [ $1.2$  mg/dL] [CI,  $0.02$  to  $0.05$  mmol/L [ $0.7$  to  $1.7$  mg/dL]]) (12 studies;  $I^2 = 0\%$ ), and triglyceride levels (net change,  $-0.07$  mmol/L [ $-6.5$  mg/dL] [CI,  $-0.14$  to  $-0.02$  mmol/L [ $-12.7$  to  $-1.8$  mg/dL]]) (13 studies;  $I^2 = 38\%$ ) (Table 10 of the Supplement, available at [www.annals.org](http://www.annals.org)). No evidence of small-study effects was found (Egger test  $P = 0.17$  for total cholesterol level, 0.75 for HDL cholesterol level, and 0.12 for triglyceride level).

### Virtual Programs

Five studies evaluated programs that were conducted via Web tools, social networking, e-mail, text messaging, video, or a combination of these, with no in-person sessions (28, 33, 41, 42, 88). One study (28) found smaller but still significant improvements from baseline in weight ( $-5\%$  vs.  $-7\%$ ) and fasting glucose level ( $-0.15$  vs.  $-0.23$  mmol/L [ $-2.7$  vs.  $-4.2$  mg/dL]) with a DVD compared with an in-person program. Two studies (41, 42) found effects on weight loss similar to

**Figure 3.** Random-effects model meta-analysis of net percentage change in weight (from baseline) in at-risk participants in combined diet and physical activity promotion programs versus usual care.



The meta-analysis of the overall net percentage change in weight (black diamond) used data from follow-up durations closest to 1 y, as indicated by the asterisks. Subgroup meta-analyses by follow-up time (open diamonds) were conducted for time points with data from  $\geq 3$  studies.  $P_{\text{Het}}$  = chi-square  $P$  value of heterogeneity; RR = risk ratio.

\* Included in overall meta-analysis.

† To avoid biased meta-analyses due to inclusion of correlated analyses, this comparison between the lower-intensity intervention and control was excluded.

those in studies with in-person sessions (−3% to −5% from baseline). One study in India (33) found that an intervention relying on text messages was effective compared with usual care, with lower diabetes incidence over 2 years (18% vs. 27%; HR, 0.64 [CI, 0.45 to 0.92]) and statistically significant net differences in HDL cholesterol and triglyceride levels but not weight, BP, or total cholesterol level. However, the fifth study (88), which was done in adolescents, found no effect on weight, although this was also true for a similar program with group sessions.

### Programs in Adolescents

Two studies were conducted in adolescents. In the study by Savoye and associates (102), adolescents who participated in twice-weekly group sessions were significantly more likely to revert to normoglycemia, lose weight, and have lower fasting glucose levels and BP compared with a control group, but there was no change in lipid profile, except triglyceride levels. None developed diabetes during the 6-month follow-up. The study by Patrick and colleagues (88) evaluated 3 programs (Web, Web and text message, and Web and group session programs) and reported no difference in weight loss compared with a control group or between the more intensive and less intensive interventions after 6 and 12 months. The study did not report incident diabetes or fasting glucose outcomes.

## DISCUSSION

Across a wide spectrum of diet and physical activity promotion programs, there is strong evidence of effectiveness in reducing new-onset diabetes. Among 16 studies, participants in these programs were consistently about 40% less likely to develop diabetes, but this outcome was evaluated in a minority (30%) of studies. Such programs also increase the likelihood of reversion to normoglycemia and improve diabetes and cardiometabolic risk factors, including overweight, high blood glucose level, high BP, and abnormal lipid profile. The effectiveness of these programs in reducing cardiovascular disease, diabetes-related complications, and death is yet to be determined because few studies reported these outcomes.

During protocol development, we searched MEDLINE and the Cochrane Database of Systematic Reviews for pertinent systematic reviews but found none that was sufficiently up-to-date and that evaluated the breadth of outcomes and range of analyses evaluated in the current review. The most comprehensive review was a health technology assessment by Gillett and coworkers (121), whose search was conducted in 2011 but also included diet or exercise interventions (not in combination); 9 randomized trials were included. An updated search found 3 similar but more restrictive reviews published since 2013, which focused on narrower subsets of studies in adults. Schellenberg and associates (122) included 9 randomized trials of diet and physical activity promotion programs that had at least 1 other component. Dunkley and colleagues

(123) included 25 studies (11 randomized trials) of programs that explicitly translated previous efficacy trials into community settings, but they also included studies of a broader population (such as obese or sedentary persons). Aguiar and coworkers (124) included only 8 studies (5 randomized trials) of diet and physical activity promotion interventions that included both aerobic and resistance training. The latter 2 reviews found effects on weight loss similar to those in our review (123, 124), and Aguiar and coworkers also found effects on fasting glucose levels similar to those in our review. In metaregression, Dunkley and colleagues found larger changes in weight with better alignment with lifestyle intervention attributes (123).

Evidence suggests that higher-intensity programs lead to greater weight loss and reduction in new-onset diabetes. Although the evaluated programs differed from each other too much to draw firm conclusions about the unique contributions of specific components, results from 12 studies that directly compared programs showed that persons who participated in more intensive programs (based on such features as number of sessions, individual sessions, and additional personnel) lost more weight and were less likely to develop diabetes. Effects on diabetes risk were similar across studies that compared programs with control groups; therefore, no differences based on differences in their programs could be ascertained. However, across all studies, programs that provided individual (vs. group) diet sessions resulted in greater reductions in fasting glucose levels, as did programs that used diet counselors (vs. no diet counselors). Programs based on the DPP study or the DPS (which were more intensive than many other programs) resulted in greater weight loss. More information on virtual delivery will be useful to increase the reach of effective programs.

On the basis of evidence from 2 of the larger studies (the U.S. DPP study and the Finnish DPS), findings seem to be applicable to wide populations (in Western countries) across race and ethnicity, socioeconomic status, risk factor status, and other demographic features. Except in 2 studies, all programs were conducted in adults; therefore, our results may not apply to children and adolescents. However, the benefit of diet and physical activity promotion programs is probably applicable to younger persons at risk for type 2 diabetes because adults and children share the mechanisms of the disease. Although most diabetes cases in children are type 1 diabetes, nearly all cases that develop from prediabetes are type 2 diabetes. Key aspects of the pathophysiology of type 2 diabetes are similar in persons of all ages; thus, the programs are likely to be effective regardless of age, assuming that they are effective at changing children's diet and physical activity. The one in-person program conducted in adolescents had effectiveness similar to that in programs conducted in adults; however, the other study of various virtual programs in adolescents found no effect on weight.

Additional studies comparing diet and physical activity promotion programs versus usual care (no program) will probably not change the overall conclusion

about the effectiveness of such programs, except those in children and adolescents and, possibly, in specific populations or settings with gaps in data. However, several areas would benefit from future research. Because the available programs were highly heterogeneous and included many features, all of which probably interacted with each other, we were unable to explain the observed heterogeneity by whether programs included specific features. Furthermore, despite often protracted descriptions of the interventions, articles often did not clearly identify who led them or what the goals were or provide other details so that the intervention could be reproduced. Future studies that compare specific program features are needed to clarify which features (for example, individual vs. group sessions, few vs. many sessions, or differently trained counselors) optimize the effectiveness of the programs and which are less critical. The most effective way to structure the maintenance phase to help program participants maintain their improvements is also unclear. In addition, with the proliferation of mobile devices and applications, the effectiveness of virtual programs needs to be investigated further. Of note, long-term follow-up of existing community-based programs is needed to evaluate the durability of the programs' effects and their effects on clinical outcomes. Although this review did not specifically address participant attrition, a better understanding of typical attrition rates is needed to understand the reasons program participants drop out and to develop methods to retain them.

In conclusion, combined diet and physical activity promotion programs are effective in reducing new-onset diabetes, increasing reversion to normoglycemia, and improving diabetes and cardiometabolic risk factors in persons at increased risk for type 2 diabetes. Programs are effective across a wide range of features, but more intensive interventions seem to be more effective. Further research is needed to discern which specific program features are most important.

From Brown University School of Public Health, Providence, Rhode Island; Tufts Medical Center and Mapi Group, Boston, Massachusetts; and University of Wisconsin, Madison, Wisconsin.

**Disclaimer:** The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the CDC or the Community Preventive Services Task Force.

**Acknowledgment:** The authors thank the members of the Community Preventive Services Task Force for their contributions and assistance with developing the protocol and suggestions on assessing the body of evidence, particularly the Community Guide Technical Monitor (David P. Hopkins, MD, MPH [Centers for Disease Control and Prevention, Atlanta, Georgia]) and the Coordination Team panel of stakeholders (Ann Albright, PhD, RD [Centers for Disease Control and Prevention, Atlanta, Georgia]; Elizabeth B. Daniels, PhD, RN [Centers for Disease Control and Prevention, Atlanta, Georgia]; Judith Fradkin, MD [National Institute of Diabetes and Digestive and Kidney Diseases, National Institutes of Health, Bethesda,

Maryland]; Todd S. Harwell, MPH [Public Health and Safety Division, State of Montana Department of Public Health and Human Services, Helena, Montana]; Matt Longjohn, MD, MPH [YMCA of the USA, Chicago, Illinois]; Melinda Maryniuk, RD, MEd, CDE [Joslin Diabetes Center, Boston, Massachusetts]; Nicolaas P. Pronk, PhD [HealthPartners, Minneapolis, Minnesota, and Harvard School of Public Health, Boston, Massachusetts]; and Patrick L. Remington, MD, MPH [School of Medicine and Public Health, University of Wisconsin, Madison, Wisconsin]). They also thank other members of their research team who participated in various stages of the review (Denish Moorthy, MBBS, MS [Micronutrients, SPRING Project, Washington, DC]; Fadi Obeid, MD [Tufts Medical Center, Boston, Massachusetts]; Rebecca Persson, BA [Boston University School of Public Health, Boston, Massachusetts]; and Katrin Uhlig, MD, MS [Tufts University School of Medicine, Boston, Massachusetts]), as well as technical editors Kate W. Harris, BA (The Community Guide, Atlanta, Georgia), and Krista Hopkins Cole, MPH (Karna, Atlanta, Georgia).

**Financial Support:** By contract 200-2012-53787 from the CDC.

**Disclosures:** Ms. Earley reports a grant from the Centers for Disease Control and Prevention during the conduct of the study. Authors not named here have disclosed no conflicts of interest. Disclosures can also be viewed at [www.acponline.org/authors/icmje/ConflictOfInterestForms.do?msNum=M15-0452](http://www.acponline.org/authors/icmje/ConflictOfInterestForms.do?msNum=M15-0452).

**Reproducible Research Statement:** *Study protocol:* No protocol document is available, but interested readers may contact Dr. Balk to receive copies of working documents (e-mail, [ethan\\_balk@brown.edu](mailto:ethan_balk@brown.edu)). *Statistical code and data set:* Available from Dr. Balk (e-mail, [ethan\\_balk@brown.edu](mailto:ethan_balk@brown.edu)).

**Requests for Single Reprints:** Ethan Balk, MD, MPH, Brown University School of Public Health, Box G-S121-8, Providence, RI 02912; e-mail, [ethan\\_balk@brown.edu](mailto:ethan_balk@brown.edu).

Current author addresses and author contributions are available at [www.annals.org](http://www.annals.org).

## References

- Centers for Disease Control and Prevention (CDC). National Diabetes Statistics Report: Estimates of Diabetes and Its Burden in the United States, 2014. Atlanta, GA: U.S. Department of Health and Human Services; 2014.
- International Diabetes Federation. IDF Diabetes Atlas. Sixth Edition. Brussels, Belgium: International Diabetes Federation; 2014. Accessed at [www.idf.org/diabetesatlas](http://www.idf.org/diabetesatlas) on 3 May 2015.
- Huang ES, Basu A, O'Grady M, Capretta JC. Projecting the future diabetes population size and related costs for the U.S. *Diabetes Care*. 2009;32:2225-9. [PMID: 19940225] doi:10.2337/dc09-0459
- American Diabetes Association. Diagnosis and classification of diabetes mellitus. *Diabetes Care*. 2010;33 Suppl 1:S62-9. [PMID: 20042775] doi:10.2337/dc10-S062
- Gerstein HC, Santaguida P, Raina P, Morrison KM, Balion C, Hunt D, et al. Annual incidence and relative risk of diabetes in people with various categories of dysglycemia: a systematic overview and meta-analysis of prospective studies. *Diabetes Res Clin Pract*. 2007;78:305-12. [PMID: 17601626]
- Knowler WC, Barrett-Connor E, Fowler SE, Hamman RF, Lachin JM, Walker EA, et al; Diabetes Prevention Program Research Group.

- Reduction in the incidence of type 2 diabetes with lifestyle intervention or metformin. *N Engl J Med.* 2002;346:393-403. [PMID: 11832527]
7. Tuomilehto J, Lindström J, Eriksson JG, Valle TT, Hämäläinen H, Ilanne-Parikka P, et al; Finnish Diabetes Prevention Study Group. Prevention of type 2 diabetes mellitus by changes in lifestyle among subjects with impaired glucose tolerance. *N Engl J Med.* 2001;344:1343-50. [PMID: 11333990]
  8. Pan XR, Li GW, Hu YH, Wang JX, Yang WY, An ZX, et al. Effects of diet and exercise in preventing NIDDM in people with impaired glucose tolerance. The Da Qing IGT and Diabetes Study. *Diabetes Care.* 1997;20:537-44. [PMID: 9096977]
  9. Knowler WC, Fowler SE, Hamman RF, Christophi CA, Hoffman HJ, Brenneman AT, et al; Diabetes Prevention Program Research Group. 10-year follow-up of diabetes incidence and weight loss in the Diabetes Prevention Program Outcomes Study. *Lancet.* 2009;374:1677-86. [PMID: 19878986] doi:10.1016/S0140-6736(09)61457-4
  10. Li G, Zhang P, Wang J, An Y, Gong Q, Gregg EW, et al. Cardiovascular mortality, all-cause mortality, and diabetes incidence after lifestyle intervention for people with impaired glucose tolerance in the Da Qing Diabetes Prevention Study: a 23-year follow-up study. *Lancet Diabetes Endocrinol.* 2014;2:474-80. [PMID: 24731674] doi:10.1016/S2213-8587(14)70057-9
  11. Lindström J, Ilanne-Parikka P, Peltonen M, Aunola S, Eriksson JG, Hemiö K, et al; Finnish Diabetes Prevention Study Group. Sustained reduction in the incidence of type 2 diabetes by lifestyle intervention: follow-up of the Finnish Diabetes Prevention Study. *Lancet.* 2006;368:1673-9. [PMID: 17098085]
  12. Albright AL, Gregg EW. Preventing type 2 diabetes in communities across the U.S.: the National Diabetes Prevention Program. *Am J Prev Med.* 2013;44:S346-51. [PMID: 23498297] doi:10.1016/j.amepre.2012.12.009
  13. Briss PA, Zaza S, Pappaioanou M, Fielding J, Wright-De Agüero L, Truman BI, et al. Developing an evidence-based Guide to Community Preventive Services—methods. The Task Force on Community Preventive Services. *Am J Prev Med.* 2000;18:35-43. [PMID: 10806978]
  14. Zaza S, Wright-De Agüero LK, Briss PA, Truman BI, Hopkins DP, Hennessy MH, et al. Data collection instrument and procedure for systematic reviews in the Guide to Community Preventive Services. Task Force on Community Preventive Services. *Am J Prev Med.* 2000;18:44-74. [PMID: 10806979]
  15. Institute of Medicine. Finding What Works in Health Care: Standards for Systematic Reviews. Washington, DC: National Academies Press; 2011.
  16. Agency for Healthcare Research and Quality. Methods Guide for Effectiveness and Comparative Effectiveness Reviews. AHRQ publication no. 10(12)-EHC063-EF. Rockville, MD: Agency for Healthcare Research and Quality; 2012.
  17. Wallace BC, Small K, Brodley CE, Lau J, Trikalinos TA. Deploying an interactive machine learning system in an evidence-based practice center: Abstract. In: Proceedings of the 2nd ACM SIGHIT International Health Informatics Symposium, Miami, Florida, 28-30 January 2012. New York: Association for Computing Machinery; 2012: 819-24. doi:10.1145/2110363.2110464
  18. Agency for Healthcare Research and Quality. Systematic Review Data Repository Web site. Accessed at <http://srdhrq.gov> on 3 May 2015.
  19. The Community Guide. Data Abstraction Form. Atlanta, GA: The Community Guide. Accessed at [www.thecommunityguide.org/methods/abstractionform.pdf](http://www.thecommunityguide.org/methods/abstractionform.pdf) on 3 May 2015.
  20. Hardy RJ, Thompson SG. A likelihood approach to meta-analysis with random effects. *Stat Med.* 1996;15:619-29. [PMID: 8731004]
  21. Harbord RM, Harris RJ, Sterne JAC. Updated tests for small-study effects in meta-analyses. *Stata J.* 2009;9:197-210.
  22. Eriksson KF, Lindgärde F. Prevention of type 2 (non-insulin-dependent) diabetes mellitus by diet and physical exercise. The 6-year Malmö feasibility study. *Diabetologia.* 1991;34:891-8. [PMID: 1778354]
  23. Oldroyd JC, Unwin NC, White M, Mathers JC, Alberti KG. Randomised controlled trial evaluating lifestyle interventions in people with impaired glucose tolerance. *Diabetes Res Clin Pract.* 2006;72:117-27. [PMID: 16297488]
  24. Penn L, White M, Oldroyd J, Walker M, Alberti KG, Mathers JC. Prevention of type 2 diabetes in adults with impaired glucose tolerance: the European Diabetes Prevention RCT in Newcastle upon Tyne, UK. *BMC Public Health.* 2009;9:342. [PMID: 19758428] doi:10.1186/1471-2458-9-342
  25. Ramachandran A, Snehalatha C, Mary S, Mukesh B, Bhaskar AD, Vijay V; Indian Diabetes Prevention Programme (IDPP). The Indian Diabetes Prevention Programme shows that lifestyle modification and metformin prevent type 2 diabetes in Asian Indian subjects with impaired glucose tolerance (IDPP-1). *Diabetologia.* 2006;49:289-97. [PMID: 16391903]
  26. Costa B, Barrio F, Cabré JJ, Piñol JL, Cos X, Solé C, et al; DEPLAN-CAT Research Group. Delaying progression to type 2 diabetes among high-risk Spanish individuals is feasible in real-life primary healthcare settings using intensive lifestyle intervention. *Diabetologia.* 2012;55:1319-28. [PMID: 22322921] doi:10.1007/s00125-012-2492-6
  27. Iqbal Hydrie MZ, Basit A, Shera AS, Hussain A. Effect of intervention in subjects with high risk of diabetes mellitus in Pakistan. *J Nutr Metab.* 2012;2012:867604. [PMID: 22888411] doi:10.1155/2012/867604
  28. Ma J, Yank V, Xiao L, Lavori PW, Wilson SR, Rosas LG, et al. Translating the Diabetes Prevention Program lifestyle intervention for weight loss into primary care: a randomized trial. *JAMA Intern Med.* 2013;173:113-21. [PMID: 23229846] doi:10.1001/2013.jamainternmed.987
  29. Roumen C, Corpeleijn E, Feskens EJ, Mensink M, Saris WH, Blaak EE. Impact of 3-year lifestyle intervention on postprandial glucose metabolism: the SLIM study. *Diabet Med.* 2008;25:597-605. [PMID: 18445174] doi:10.1111/j.1464-5491.2008.02417.x
  30. Sakane N, Sato J, Tsushita K, Tsujii S, Kotani K, Tsuzaki K, et al; Japan Diabetes Prevention Program (JDPP) Research Group. Prevention of type 2 diabetes in a primary healthcare setting: three-year results of lifestyle intervention in Japanese subjects with impaired glucose tolerance. *BMC Public Health.* 2011;11:40. [PMID: 21235825] doi:10.1186/1471-2458-11-40
  31. Vermunt PW, Milder IE, Wielaard F, de Vries JH, van Oers HA, Westert GP. Lifestyle counseling for type 2 diabetes risk reduction in Dutch primary care: results of the APHRODITE study after 0.5 and 1.5 years. *Diabetes Care.* 2011;34:1919-25. [PMID: 21775759] doi:10.2337/dc10-2293
  32. Bhopal RS, Douglas A, Wallia S, Forbes JF, Lean ME, Gill JM, et al. Effect of a lifestyle intervention on weight change in south Asian individuals in the UK at high risk of type 2 diabetes: a family-cluster randomised controlled trial. *Lancet Diabetes Endocrinol.* 2014;2:218-27. [PMID: 24622752] doi:10.1016/S2213-8587(13)70204-3
  33. Ramachandran A, Snehalatha C, Ram J, Selvam S, Simon M, Nanditha A, et al. Effectiveness of mobile phone messaging in prevention of type 2 diabetes by lifestyle modification in men in India: a prospective, parallel-group, randomised controlled trial. *Lancet Diabetes Endocrinol.* 2013;1:191-8. [PMID: 24622367] doi:10.1016/S2213-8587(13)70067-6
  34. Sakane N, Sato J, Tsushita K, Tsujii S, Kotani K, Tominaga M, et al; Japan Diabetes Prevention Program (JDPP) Research Group. Effect of baseline HbA1c level on the development of diabetes by lifestyle intervention in primary healthcare settings: insights from subanalysis of the Japan Diabetes Prevention Program. *BMJ Open Diabetes Res Care.* 2014;2:e000003. [PMID: 25452854] doi:10.1136/bmjdc-2013-000003
  35. Ackermann RT, Finch EA, Brizendine E, Zhou H, Marrero DG. Translating the Diabetes Prevention Program into the community. The DEPLOY Pilot Study. *Am J Prev Med.* 2008;35:357-63. [PMID: 18779029] doi:10.1016/j.amepre.2008.06.035
  36. Cezaretto A, Siqueira-Catania A, de Barros CR, Salvador EP, Ferreira SR. Benefits on quality of life concomitant to metabolic improvement in intervention program for prevention of diabetes mel-

- litus. *Qual Life Res.* 2012;21:105-13. [PMID: 21538199] doi:10.1007/s11136-011-9919-2
37. Kanaya AM, Santoyo-Olsson J, Gregorich S, Grossman M, Moore T, Stewart AL. The Live Well, Be Well study: a community-based, translational lifestyle program to lower diabetes risk factors in ethnic minority and lower-socioeconomic status adults. *Am J Public Health.* 2012;102:1551-8. [PMID: 22698027] doi:10.2105/AJPH.2011.300456
38. Kulzer B, Hermanns N, Gorges D, Schwarz P, Haak T. Prevention of diabetes self-management program (PREDIAS): effects on weight, metabolic risk factors, and behavioral outcomes. *Diabetes Care.* 2009;32:1143-6. [PMID: 19509014] doi:10.2337/dc08-2141
39. Moore SM, Hardie EA, Hackworth NJ, Critchley CR, Kyrios M, Buzwell SA, et al. Can the onset of type 2 diabetes be delayed by a group-based lifestyle intervention? A randomised control trial. *Psychol Health.* 2011;26:485-99. [PMID: 20945253] doi:10.1080/08870440903548749
40. Ockene IS, Tellez TL, Rosal MC, Reed GW, Mordes J, Merriam PA, et al. Outcomes of a Latino community-based intervention for the prevention of diabetes: the Lawrence Latino Diabetes Prevention Project. *Am J Public Health.* 2012;102:336-42. [PMID: 22390448] doi:10.2105/AJPH.2011.300357
41. Tate DF, Jackvony EH, Wing RR. Effects of Internet behavioral counseling on weight loss in adults at risk for type 2 diabetes: a randomized trial. *JAMA.* 2003;289:1833-6. [PMID: 12684363]
42. Ackermann RT, Sandy LG, Beauregard T, Coblitz M, Norton KL, Voita D. A randomized comparative effectiveness trial of using cable television to deliver diabetes prevention programming. *Obesity (Silver Spring).* 2014;22:1601-7. [PMID: 24740868] doi:10.1002/oby.20762
43. Cole RE, Boyer KM, Spanbauer SM, Sprague D, Bingham M. Effectiveness of prediabetes nutrition shared medical appointments: prevention of diabetes. *Diabetes Educ.* 2013;39:344-53. [PMID: 23589326] doi:10.1177/0145721713484812
44. Dunbar JA, Davis-Lameloise N, Philpot B, Reddy P, Bunker S, Heistaro S, et al. Sustained gains from a diabetes prevention program and the role of telephone support. *Int J Diabetes Mellit.* 2010;2:95-100.
45. Gagnon C, Brown C, Couture C, Kamga-Ngande CN, Hivert MF, Baillargeon JP, et al. A cost-effective moderate-intensity interdisciplinary weight-management programme for individuals with prediabetes. *Diabetes Metab.* 2011;37:410-8. [PMID: 21489843] doi:10.1016/j.diabet.2011.01.003
46. Jiang L, Manson SM, Beals J, Henderson WG, Huang H, Acton KJ, et al; Special Diabetes Program for Indians Diabetes Prevention Demonstration Project. Translating the Diabetes Prevention Program into American Indian and Alaska Native communities: results from the Special Diabetes Program for Indians Diabetes Prevention demonstration project. *Diabetes Care.* 2013;36:2027-34. [PMID: 23275375] doi:10.2337/dc12-1250
47. Kosaka K, Noda M, Kuzuya T. Prevention of type 2 diabetes by lifestyle intervention: a Japanese trial in IGT males. *Diabetes Res Clin Pract.* 2005;67:152-62. [PMID: 15649575]
48. Liao D, Asberry PJ, Shofer JB, Callahan H, Matthys C, Boyko EJ, et al. Improvement of BMI, body composition, and body fat distribution with lifestyle modification in Japanese Americans with impaired glucose tolerance. *Diabetes Care.* 2002;25:1504-10. [PMID: 12196418]
49. Nilsen V, Bakke PS, Gallefoss F. Effects of lifestyle intervention in persons at risk for type 2 diabetes mellitus—results from a randomised, controlled trial. *BMC Public Health.* 2011;11:893. [PMID: 22117618] doi:10.1186/1471-2458-11-893
50. Saito T, Watanabe M, Nishida J, Izumi T, Omura M, Takagi T, et al; Zensharen Study for Prevention of Lifestyle Diseases Group. Lifestyle modification and prevention of type 2 diabetes in overweight Japanese with impaired fasting glucose levels: a randomized controlled trial. *Arch Intern Med.* 2011;171:1352-60. [PMID: 21824948] doi:10.1001/archinternmed.2011.275
51. Sepah SC, Jiang L, Peters AL. Translating the Diabetes Prevention Program into an online social network: validation against CDC standards. *Diabetes Educ.* 2014;40:435-443. [PMID: 24723130]
52. Admiraal WM, Vlaar EM, Nierkens V, Holleman F, Middelkoop BJ, Stronks K, et al. Intensive lifestyle intervention in general practice to prevent type 2 diabetes among 18 to 60-year-old South Asians: 1-year effects on the weight status and metabolic profile of participants in a randomized controlled trial. *PLoS One.* 2013;8:e68605. [PMID: 23894322] doi:10.1371/journal.pone.0068605
53. Janus ED, Best JD, Davis-Lameloise N, Philpot B, Hernan A, Bennett CM, et al; Melbourne Diabetes Prevention Study research group. Scaling-up from an implementation trial to state-wide coverage: results from the preliminary Melbourne Diabetes Prevention Study. *Trials.* 2012;13:152. [PMID: 22929458] doi:10.1186/1745-6215-13-152
54. Islam NS, Zanolwiak JM, Wyatt LC, Kavathe R, Singh H, Kwon SC, et al. Diabetes prevention in the New York City Sikh Asian Indian community: a pilot study. *Int J Environ Res Public Health.* 2014;11:5462-86. [PMID: 24852392] doi:10.3390/ijerph110505462
55. Gillison F, Stathi A, Reddy P, Perry R, Taylor G, Bennett P, et al. Processes of behavior change and weight loss in a theory-based weight loss intervention program: a test of the process model for lifestyle behavior change. *Int J Behav Nutr Phys Act.* 2015;12:2. [PMID: 25592314] doi:10.1186/s12966-014-0160-6
56. Katula JA, Vitolins MZ, Rosenberger EL, Blackwell CS, Morgan TM, Lawlor MS, et al. One-year results of a community-based translation of the Diabetes Prevention Program: Healthy-Living Partnerships to Prevent Diabetes (HELP PD) Project. *Diabetes Care.* 2011;34:1451-7. [PMID: 21593290] doi:10.2337/dc10-2115
57. Penn L, Ryan V, White M. Feasibility, acceptability and outcomes at a 12-month follow-up of a novel community-based intervention to prevent type 2 diabetes in adults at high risk: mixed methods pilot study. *BMJ Open.* 2013;3:e003585. [PMID: 24227871] doi:10.1136/bmjopen-2013-003585
58. Weinstock RS, Trief PM, Cibula D, Morin PC, Delahanty LM. Weight loss success in metabolic syndrome by telephone interventions: results from the SHINE Study. *J Gen Intern Med.* 2013;28:1620-8. [PMID: 23843020] doi:10.1007/s11606-013-2529-7
59. Absetz P, Valve R, Oldenburg B, Heinonen H, Nissinen A, Fogelholm M, et al. Type 2 diabetes prevention in the "real world": one-year results of the GOAL Implementation Trial. *Diabetes Care.* 2007;30:2465-70. [PMID: 17586741]
60. Absetz P, Oldenburg B, Hankonen N, Valve R, Heinonen H, Nissinen A, et al. Type 2 diabetes prevention in the real world: three-year results of the GOAL lifestyle implementation trial. *Diabetes Care.* 2009;32:1418-20. [PMID: 19401442] doi:10.2337/dc09-0039
61. Bouchard DR, Baillargeon JP, Gagnon C, Brown C, Langlois MF. Impact of health professionals' contact frequency on response to a lifestyle intervention with individuals at high risk for diabetes. *Diabetes Res Clin Pract.* 2012;96:129-34. [PMID: 22245692] doi:10.1016/j.diabetes.2011.12.019
62. Carr DB, Utzschneider KM, Boyko EJ, Asberry PJ, Hull RL, Kodama K, et al. A reduced-fat diet and aerobic exercise in Japanese Americans with impaired glucose tolerance decreases intra-abdominal fat and improves insulin sensitivity but not beta-cell function. *Diabetes.* 2005;54:340-7. [PMID: 15677490]
63. Critchley CR, Hardie EA, Moore SM. Examining the psychological pathways to behavior change in a group-based lifestyle program to prevent type 2 diabetes. *Diabetes Care.* 2012;35:699-705. [PMID: 22338102] doi:10.2337/dc11-1183
64. De la Rosa A, Tahsin B, Sanghani R, Pikelny I, Fogelfeld L, Stroger JH Jr. Detecting and managing metabolic syndrome: a feasibility study in a general medicine clinic. *Ethn Dis.* 2008;18:S16-8.
65. Delahanty LM, Peyrot M, Shrader PJ, Williamson DA, Meigs JB, Nathan DM; DPP Research Group. Pretreatment, psychological, and behavioral predictors of weight outcomes among lifestyle intervention participants in the Diabetes Prevention Program (DPP). *Diabetes Care.* 2013;36:34-40. [PMID: 23129133] doi:10.2337/dc12-0733

66. The Diabetes Prevention Program. Design and methods for a clinical trial in the prevention of type 2 diabetes. *Diabetes Care*. 1999;22:623-34. [PMID: 10189543]
67. Crandall J, Schade D, Ma Y, Fujimoto WY, Barrett-Connor E, Fowler S, et al; Diabetes Prevention Program Research Group. The influence of age on the effects of lifestyle modification and metformin in prevention of diabetes. *J Gerontol A Biol Sci Med Sci*. 2006;61:1075-81. [PMID: 17077202]
68. Eriksson J, Lindström J, Valle T, Aunola S, Hämäläinen H, Ilanne-Parikka P, et al. Prevention of type II diabetes in subjects with impaired glucose tolerance: the Diabetes Prevention Study (DPS) in Finland. Study design and 1-year interim report on the feasibility of the lifestyle intervention programme. *Diabetologia*. 1999;42:793-801. [PMID: 10440120]
69. Eriksson KF, Lindgärde F. No excess 12-year mortality in men with impaired glucose tolerance who participated in the Malmö Preventive Trial with diet and exercise. *Diabetologia*. 1998;41:1010-6. [PMID: 9754818]
70. Gilis-Januszewska A, Szybinski Z, Kissimova-Skarbek K, Piwonska-Solska B, Pach D, Topor-Madry R, et al. Prevention of type 2 diabetes by lifestyle intervention in primary health care setting in Poland: Diabetes in Europe Prevention using Lifestyle, physical Activity and Nutritional intervention (DE-PLAN) project. *Br J Diabetes Vasc Dis*. 2011;11:198-203.
71. Gong Q, Gregg EW, Wang J, An Y, Zhang P, Yang W, et al. Long-term effects of a randomised trial of a 6-year lifestyle intervention in impaired glucose tolerance on diabetes-related microvascular complications: the China Da Qing Diabetes Prevention Outcome Study. *Diabetologia*. 2011;54:300-7. [PMID: 21046360] doi:10.1007/s00125-010-1948-9
72. Haffner S, Temprosa M, Crandall J, Fowler S, Goldberg R, Horton E, et al; Diabetes Prevention Program Research Group. Intensive lifestyle intervention or metformin on inflammation and coagulation in participants with impaired glucose tolerance. *Diabetes*. 2005;54:1566-72. [PMID: 15855347]
73. Hosper K, Deutekom M, Stronks K. The effectiveness of "Exercise on Prescription" in stimulating physical activity among women in ethnic minority groups in the Netherlands: protocol for a randomized controlled trial. *BMC Public Health*. 2008;8:406. [PMID: 19077190] doi:10.1186/1471-2458-8-406
74. Ilanne-Parikka P, Eriksson JG, Lindström J, Peltonen M, Aunola S, Hämäläinen H, et al; Finnish Diabetes Prevention Study Group. Effect of lifestyle intervention on the occurrence of metabolic syndrome and its components in the Finnish Diabetes Prevention Study. *Diabetes Care*. 2008;31:805-7. [PMID: 18184907] doi:10.2337/dc07-1117
75. Katula JA, Vitolins MZ, Morgan TM, Lawlor MS, Blackwell CS, Isom SP, et al. The Healthy Living Partnerships to Prevent Diabetes study: 2-year outcomes of a randomized controlled trial. *Am J Prev Med*. 2013;44:S324-32. [PMID:23498294]doi:10.1016/j.amepre.2012.12.015
76. Kyrios M, Moore SM, Hackworth N, Buzwell SA, Crafti N, Critchley C, et al. The influence of depression and anxiety on outcomes after an intervention for prediabetes. *Med J Aust*. 2009;190:S81-5. [PMID: 19351299]
77. Laatikainen T, Dunbar JA, Chapman A, Kilkkinen A, Vartiainen E, Heistaro S, et al. Prevention of type 2 diabetes by lifestyle intervention in an Australian primary health care setting: Greater Green Triangle (GGT) Diabetes Prevention Project. *BMC Public Health*. 2007;7:249. [PMID: 17877832]
78. Li G, Zhang P, Wang J, Gregg EW, Yang W, Gong Q, et al. The long-term effect of lifestyle interventions to prevent diabetes in the China Da Qing Diabetes Prevention Study: a 20-year follow-up study. *Lancet*. 2008;371:1783-9. [PMID: 18502303] doi:10.1016/S0140-6736(08)60766-7
79. Lindström J, Louheranta A, Mannelin M, Rastas M, Salminen V, Eriksson J, et al; Finnish Diabetes Prevention Study Group. The Finnish Diabetes Prevention Study (DPS): lifestyle intervention and 3-year results on diet and physical activity. *Diabetes Care*. 2003;26:3230-6. [PMID: 14633807]
80. Lindström J, Eriksson JG, Valle TT, Aunola S, Cepaitis Z, Hakumäki M, et al. Prevention of diabetes mellitus in subjects with impaired glucose tolerance in the Finnish Diabetes Prevention Study: results from a randomized clinical trial. *J Am Soc Nephrol*. 2003;14:S108-13. [PMID: 12819313]
81. Lindström J, Peltonen M, Eriksson JG, Aunola S, Hämäläinen H, Ilanne-Parikka P, et al; Finnish Diabetes Prevention Study (DPS) Group. Determinants for the effectiveness of lifestyle intervention in the Finnish Diabetes Prevention Study. *Diabetes Care*. 2008;31:857-62. [PMID: 18252900] doi:10.2337/dc07-2162
82. Lindström J, Peltonen M, Eriksson JG, Ilanne-Parikka P, Aunola S, Keinänen-Kiukaanniemi S, et al; Finnish Diabetes Prevention Study (DPS). Improved lifestyle and decreased diabetes risk over 13 years: long-term follow-up of the randomised Finnish Diabetes Prevention Study (DPS). *Diabetologia*. 2013;56:284-93. [PMID: 23093136] doi:10.1007/s00125-012-2752-5
83. Makrilakis K, Liatis S, Grammatikou S, Perrea D, Katsilambros N. Implementation and effectiveness of the first community lifestyle intervention programme to prevent type 2 diabetes in Greece. The DE-PLAN study. *Diabet Med*. 2010;27:459-65. [PMID: 20536519] doi:10.1111/j.1464-5491.2010.02918.x
84. Makrilakis K, Grammatikou S, Liatis S, Kontogianni M, Perrea D, Dimosthenopoulos C, et al. The effect of a non-intensive community-based lifestyle intervention on the prevalence of metabolic syndrome. The DEPLAN study in Greece. *Hormones (Athens)*. 2012;11:316-24. [PMID: 22908064]
85. Merriam PA, Tellez TL, Rosal MC, Olendzki BC, Ma Y, Pagoto SL, et al. Methodology of a diabetes prevention translational research project utilizing a community-academic partnership for implementation in an underserved Latino community. *BMC Med Res Methodol*. 2009;9:20. [PMID: 19284663] doi:10.1186/1471-2288-9-20
86. Oldroyd JC, Unwin NC, White M, Imrie K, Mathers JC, Alberti KG. Randomised controlled trial evaluating the effectiveness of behavioural interventions to modify cardiovascular risk factors in men and women with impaired glucose tolerance: outcomes at 6 months. *Diabetes Res Clin Pract*. 2001;52:29-43. [PMID: 11182214]
87. Orchard TJ, Temprosa M, Barrett-Connor E, Fowler SE, Goldberg RB, Mather KJ, et al; Diabetes Prevention Program Outcomes Study Research Group. Long-term effects of the Diabetes Prevention Program interventions on cardiovascular risk factors: a report from the DPP Outcomes Study. *Diabet Med*. 2013;30:46-55. [PMID: 22812594] doi:10.1111/j.1464-5491.2012.03750.x
88. Patrick K, Norman GJ, Davila EP, Calfas KJ, Raab F, Gottschalk M, et al. Outcomes of a 12-month technology-based intervention to promote weight loss in adolescents at risk for type 2 diabetes. *J Diabetes Sci Technol*. 2013;7:759-70. [PMID: 23759410]
89. Penn L, White M, Lindström J, den Boer AT, Blaak E, Eriksson JG, et al. Importance of weight loss maintenance and risk prediction in the prevention of type 2 diabetes: analysis of European Diabetes Prevention Study RCT. *PLoS One*. 2013;8:e57143. [PMID: 23451166] doi:10.1371/journal.pone.0057143
90. Perreault L, Ma Y, Dagogo-Jack S, Horton E, Marrero D, Crandall J, et al; Diabetes Prevention Program. Sex differences in diabetes risk and the effect of intensive lifestyle modification in the Diabetes Prevention Program. *Diabetes Care*. 2008;31:1416-21. [PMID: 18356403] doi:10.2337/dc07-2390
91. Perreault L, Kahn SE, Christophi CA, Knowler WC, Hamman RF; Diabetes Prevention Program Research Group. Regression from pre-diabetes to normal glucose regulation in the diabetes prevention program. *Diabetes Care*. 2009;32:1583-8. [PMID: 19587364] doi:10.2337/dc09-0523
92. Perreault L, Pan Q, Mather KJ, Watson KE, Hamman RF, Kahn SE; Diabetes Prevention Program Research Group. Effect of regression from prediabetes to normal glucose regulation on long-term reduction in diabetes risk: results from the Diabetes Prevention Program Outcomes Study. *Lancet*. 2012;379:2243-51. [PMID: 22683134] doi:10.1016/S0140-6736(12)60525-X
93. Ramachandran A, Snehalatha C, Mary S, Selvam S, Kumar CK, Seeli AC, et al. Pioglitazone does not enhance the effectiveness of lifestyle modification in preventing conversion of impaired glucose

- tolerance to diabetes in Asian Indians: results of the Indian Diabetes Prevention Programme-2 (IDPP-2). *Diabetologia*. 2009;52:1019-26. [PMID: 19277602] doi:10.1007/s00125-009-1315-x
94. Ramachandran A, Arun N, Shetty AS, Snehalatha C. Efficacy of primary prevention interventions when fasting and postglucose dysglycemia coexist: analysis of the Indian Diabetes Prevention Programmes (IDPP-1 and IDPP-2). *Diabetes Care*. 2010;33:2164-8. [PMID: 20519663] doi:10.2337/dc09-1150
95. Ratner R, Goldberg R, Haffner S, Marcovina S, Orchard T, Fowler S, et al; Diabetes Prevention Program Research Group. Impact of intensive lifestyle and metformin therapy on cardiovascular disease risk factors in the diabetes prevention program. *Diabetes Care*. 2005;28:888-94. [PMID: 15793191]
96. Rautio N, Jokelainen J, Oksa H, Saaristo T, Peltonen M, Niskanen L, et al; FIN-D2D Study Group. Socioeconomic position and effectiveness of lifestyle intervention in prevention of type 2 diabetes: one-year follow-up of the FIN-D2D project. *Scand J Public Health*. 2011;39:561-70. [PMID: 21622677] doi:10.1177/1403494811408482
97. Rautio N, Jokelainen J, Oksa H, Saaristo T, Peltonen M, Niskanen L, et al. Participation, socioeconomic status and group or individual counselling intervention in individuals at high risk for type 2 diabetes: one-year follow-up study of the FIN-D2D-project. *Prim Care Diabetes*. 2012;6:277-83. [PMID: 22868007] doi:10.1016/j.pcd.2012.07.002
98. Rautio N, Jokelainen J, Oksa H, Saaristo T, Peltonen M, Puolijoki H, et al. Family history of diabetes and effectiveness of lifestyle counselling on the cardio-metabolic risk profile in individuals at high risk of type 2 diabetes: 1-year follow-up of the FIN-D2D project. *Diabet Med*. 2012;29:207-11. [PMID: 21781153] doi:10.1111/j.1464-5491.2011.03388.x
99. Rautio N, Jokelainen J, Saaristo T, Oksa H, Keinänen-Kiukaanniemi S, Peltonen M, et al; FIN-D2D Writing Group. Predictors of success of a lifestyle intervention in relation to weight loss and improvement in glucose tolerance among individuals at high risk for type 2 diabetes: the FIN-D2D project. *J Prim Care Community Health*. 2013;4:59-66. [PMID: 23799691] doi:10.1177/2150131912444130
100. Roumen C, Feskens EJ, Corpeleijn E, Mensink M, Saris WH, Blaak EE. Predictors of lifestyle intervention outcome and dropout: the SLIM study. *Eur J Clin Nutr*. 2011;65:1141-7. [PMID: 21587283] doi:10.1038/ejcn.2011.74
101. Saaristo T, Moilanen L, Korpi-Hyövälti E, Vanhala M, Saltevo J, Niskanen L, et al. Lifestyle intervention for prevention of type 2 diabetes in primary health care: one-year follow-up of the Finnish National Diabetes Prevention Program (FIN-D2D). *Diabetes Care*. 2010;33:2146-51. [PMID: 20664020] doi:10.2337/dc10-0410
102. Savoye M, Caprio S, Dziura J, Camp A, Germain G, Summers C, et al. Reversal of early abnormalities in glucose metabolism in obese youth: results of an intensive lifestyle randomized controlled trial. *Diabetes Care*. 2014;37:317-24. [PMID: 24062325] doi:10.2337/dc13-1571
103. Snehalatha C, Mary S, Selvam S, Sathish Kumar CK, Shetty SB, Nanditha A, et al. Changes in insulin secretion and insulin sensitivity in relation to the glycemic outcomes in subjects with impaired glucose tolerance in the Indian Diabetes Prevention Programme-1 (IDPP-1). *Diabetes Care*. 2009;32:1796-801. [PMID: 19587369] doi:10.2337/dc09-0676
104. Swanson CM, Bersoux S, Larson MH, Aponte-Furlow RT, Flatten SS, Olsen CL, et al. An outpatient-based clinical program for diabetes prevention: an update. *Endocr Pract*. 2012;18:200-8. [PMID: 22068253] doi:10.4158/EP11226.OR
105. Uusitupa M, Peltonen M, Lindström J, Aunola S, Ilanne-Parikka P, Keinänen-Kiukaanniemi S, et al; Finnish Diabetes Prevention Study Group. Ten-year mortality and cardiovascular morbidity in the Finnish Diabetes Prevention Study—secondary analysis of the randomized trial. *PLoS One*. 2009;4:e5656. [PMID: 19479072] doi:10.1371/journal.pone.0005656
106. Uusitupa MI, Stancáková A, Peltonen M, Eriksson JG, Lindström J, Aunola S, et al. Impact of positive family history and genetic risk variants on the incidence of diabetes: the Finnish Diabetes Prevention Study. *Diabetes Care*. 2011;34:418-23. [PMID: 20980412] doi:10.2337/dc10-1013
107. Vanderwood KK, Hall TO, Harwell TS, Butcher MK, Helgeson SD; Montana Cardiovascular Disease and Diabetes Prevention Program Workgroup. Implementing a state-based cardiovascular disease and diabetes prevention program. *Diabetes Care*. 2010;33:2543-5. [PMID: 20805260] doi:10.2337/dc10-0862
108. Vermunt PW, Milder IE, Wielaard F, de Vries JH, Baan CA, van Oers JA, et al. A lifestyle intervention to reduce type 2 diabetes risk in Dutch primary care: 2.5-year results of a randomized controlled trial. *Diabet Med*. 2012;29:e223-31. [PMID: 22416789] doi:10.1111/j.1464-5491.2012.03648.x
109. Vlaar EM, van Valkengoed IG, Nierkens V, Nicolaou M, Middekoop BJ, Stronks K. Feasibility and effectiveness of a targeted diabetes prevention program for 18 to 60-year-old South Asian migrants: design and methods of the DHIAAN study. *BMC Public Health*. 2012;12:371. [PMID: 22621376] doi:10.1186/1471-2458-12-371
110. Vojta D, Koehler TB, Longjohn M, Lever JA, Caputo NF. A coordinated national model for diabetes prevention: linking health systems to an evidence-based community program. *Am J Prev Med*. 2013;44:S301-6. [PMID: 23498291] doi:10.1016/j.amepre.2012.12.018
111. West DS, Elaine Prewitt T, Bursac Z, Felix HC. Weight loss of black, white, and Hispanic men and women in the Diabetes Prevention Program. *Obesity (Silver Spring)*. 2008;16:1413-20. [PMID: 18421273] doi:10.1038/oby.2008.224
112. Wikström K, Peltonen M, Eriksson JG, Aunola S, Ilanne-Parikka P, Keinänen-Kiukaanniemi S, et al. Educational attainment and effectiveness of lifestyle intervention in the Finnish Diabetes Prevention Study. *Diabetes Res Clin Pract*. 2009;86:e1-5. [PMID: 19592125] doi:10.1016/j.diabres.2009.06.014
113. Wing RR, Hamman RF, Bray GA, Delahanty L, Edelstein SL, Hill JO, et al; Diabetes Prevention Program Research Group. Achieving weight and activity goals among diabetes prevention program lifestyle participants. *Obes Res*. 2004;12:1426-34. [PMID: 15483207]
114. Xiao L, Yank V, Wilson SR, Lavori PW, Ma J. Two-year weight-loss maintenance in primary care-based Diabetes Prevention Program lifestyle interventions. *Nutr Diabetes*. 2013;3:e76. [PMID: 23797383] doi:10.1038/nutd.2013.17
115. Trief PM, Cibula D, Delahanty LM, Weinstock RS. Depression, stress, and weight loss in individuals with metabolic syndrome in SHINE, a DPP translation study. *Obesity (Silver Spring)*. 2014;22:2532-8. [PMID: 25251749] doi:10.1002/oby.20916
116. Diabetes Prevention Program Research Group. HbA1c as a predictor of diabetes and as an outcome in the Diabetes Prevention Program: a randomized clinical trial. *Diabetes Care*. 2015;38:51-8. [PMID: 25336746] doi:10.2337/dc14-0886
117. Rautio N, Jokelainen J, Oksa H, Saaristo T, Moilanen L, Vanhala M, et al. Do depressive symptoms have an impact on the effectiveness of lifestyle counseling in prevention of type 2 diabetes? One-year follow-up of FIN-D2D. *Prim Care Diabetes*. 2014;8:43-7. [PMID: 24238822] doi:10.1016/j.pcd.2013.10.005
118. den Boer AT, Herraets IJ, Stegen J, Roumen C, Corpeleijn E, Schaper NC, et al. Prevention of the metabolic syndrome in IGT subjects in a lifestyle intervention: results from the SLIM study. *Nutr Metab Cardiovasc Dis*. 2013;23:1147-53. [PMID: 23462149] doi:10.1016/j.numecd.2012.12.005
119. Azar KM, Xiao L, Ma J. Baseline obesity status modifies effectiveness of adapted diabetes prevention program lifestyle interventions for weight management in primary care. *Biomed Res Int*. 2013;2013:191209. [PMID: 24369008] doi:10.1155/2013/191209
120. Alibasic E, Ramic E, Alic A. Prevention of diabetes in family medicine. *Mater Sociomed*. 2013;25:80-2. [PMID: 24082827] doi:10.5455/msm.2013.25.80-82
121. Gillett M, Royle P, Snaith A, Scotland G, Poobalan A, Imamura M, et al. Non-pharmacological interventions to reduce the risk of diabetes in people with impaired glucose regulation: a systematic

review and economic evaluation. *Health Technol Assess.* 2012;16:1-236, iii-iv. [PMID: 22935084] doi:10.3310/hta16330

122. Schellenberg ES, Dryden DM, Vandermeer B, Ha C, Korownyk C. Lifestyle interventions for patients with and at risk for type 2 diabetes: a systematic review and meta-analysis. *Ann Intern Med.* 2013;159:543-51. [PMID: 24126648] doi:10.7326/0003-4819-159-8-201310150-00007

123. Dunkley AJ, Bodicoat DH, Greaves CJ, Russell C, Yates T, Davies MJ, et al. Diabetes prevention in the real world: effectiveness of

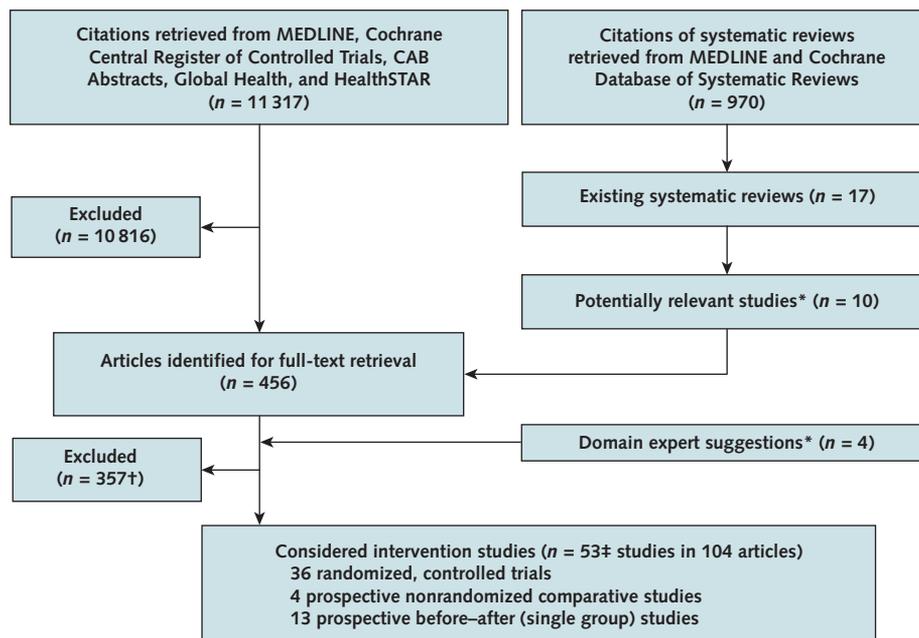
pragmatic lifestyle interventions for the prevention of type 2 diabetes and of the impact of adherence to guideline recommendations: a systematic review and meta-analysis. *Diabetes Care.* 2014;37:922-33. [PMID: 24652723] doi:10.2337/dc13-2195

124. Aguiar EJ, Morgan PJ, Collins CE, Plotnikoff RC, Callister R. Efficacy of interventions that include diet, aerobic and resistance training components for type 2 diabetes prevention: a systematic review with meta-analysis. *Int J Behav Nutr Phys Act.* 2014;11:2. [PMID: 24423095] doi:10.1186/1479-5868-11-2

**Current Author Addresses:** Dr. Balk: Brown University School of Public Health, Box G-S121-8, Providence, RI 02912.  
 Ms. Earley: Mapi USA, 180 Canal Street, Suite 503, Boston, MA 02114.  
 Dr. Raman and Ms. Avendano: Center for Clinical Evidence Synthesis, Tufts Medical Center, 800 Washington Street, Box 063, Boston, MA 02111.  
 Dr. Pittas: Division of Endocrinology, Diabetes and Metabolism, Tufts Medical Center, 800 Washington Street, Box 268, Boston, MA 02111.  
 Dr. Remington: Health Science Learning Center, Room 4263, 750 Highland Avenue, Madison, WI 53705.

**Author Contributions:** Conception and design: E.M. Balk, A. Earley, A.G. Pittas, P.L. Remington.  
 Analysis and interpretation of the data: E.M. Balk, A. Earley, G. Raman, A.G. Pittas, P.L. Remington.  
 Drafting of the article: E.M. Balk, A. Earley, G. Raman.  
 Critical revision for important intellectual content: A. Earley, G. Raman, A.G. Pittas.  
 Final approval of the article: E.M. Balk, A. Earley, G. Raman, E.E. Avendano, A.G. Pittas, P.L. Remington.  
 Provision of study materials or patients: A. Earley, G. Raman.  
 Statistical expertise: E.M. Balk, G. Raman.  
 Administrative, technical, or logistic support: A. Earley, E.E. Avendano, P.L. Remington.  
 Collection and assembly of data: E.M. Balk, A. Earley, G. Raman, E.E. Avendano.

**Appendix Figure 1.** Summary of evidence search and selection.

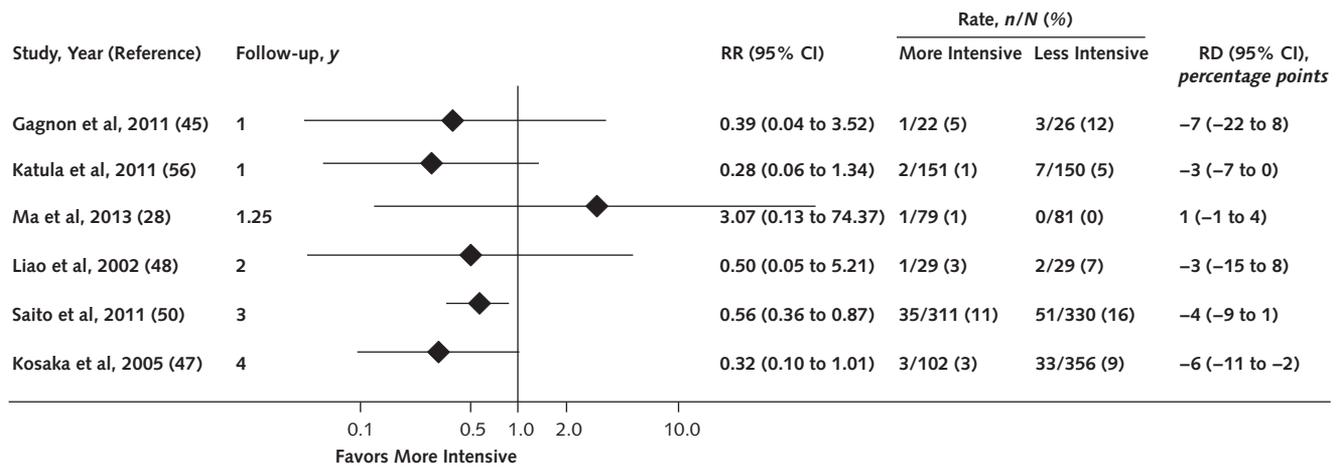


\* Not already screened.

† Not a population of interest ( $n = 70$ ), diet or physical activity alone ( $n = 47$ ), no outcome of interest reported ( $n = 36$ ), not intervention of interest ( $n = 31$ ), single-group study with  $<100$  participants ( $n = 25$ ), protocol or baseline data only ( $n = 21$ ), not a primary study ( $n = 18$ ), no additional data compared with included article ( $n = 18$ ), cost-effectiveness analysis only ( $n = 15$ ),  $<30$  participants per group ( $n = 15$ ),  $>10\%$  of participants did not meet eligibility criteria ( $n = 15$ ), intervention lasted  $<3$  mo or involved only 1 session ( $n = 13$ ),  $<6$  mo of follow-up ( $n = 13$ ), no analyses of interest ( $n = 10$ ), abstract only ( $n = 6$ ), retrospective study or retracted or unavailable article ( $n = 4$ ).

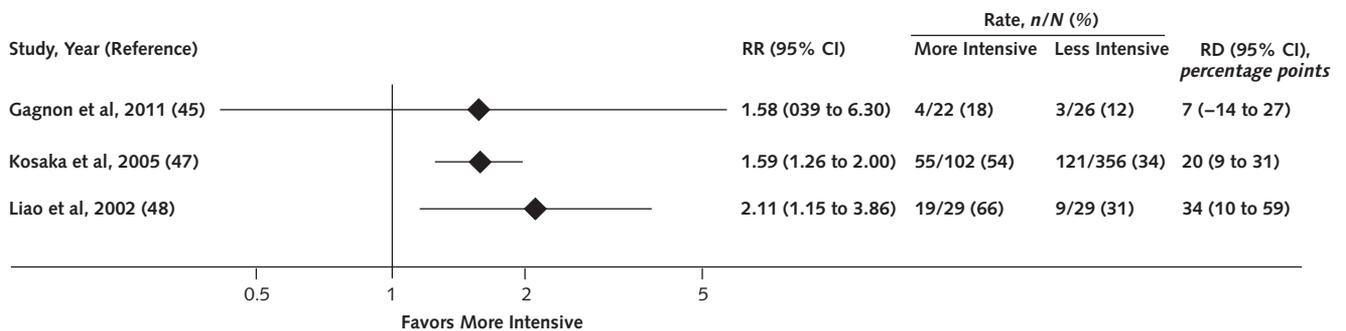
‡ Excludes 1 prospective nonrandomized comparative study that was not analyzed because of limited quality of execution.

**Appendix Figure 2.** Forest plot of RR of incident diabetes in at-risk participants in more intensive versus less intensive combined diet and physical activity promotion programs.



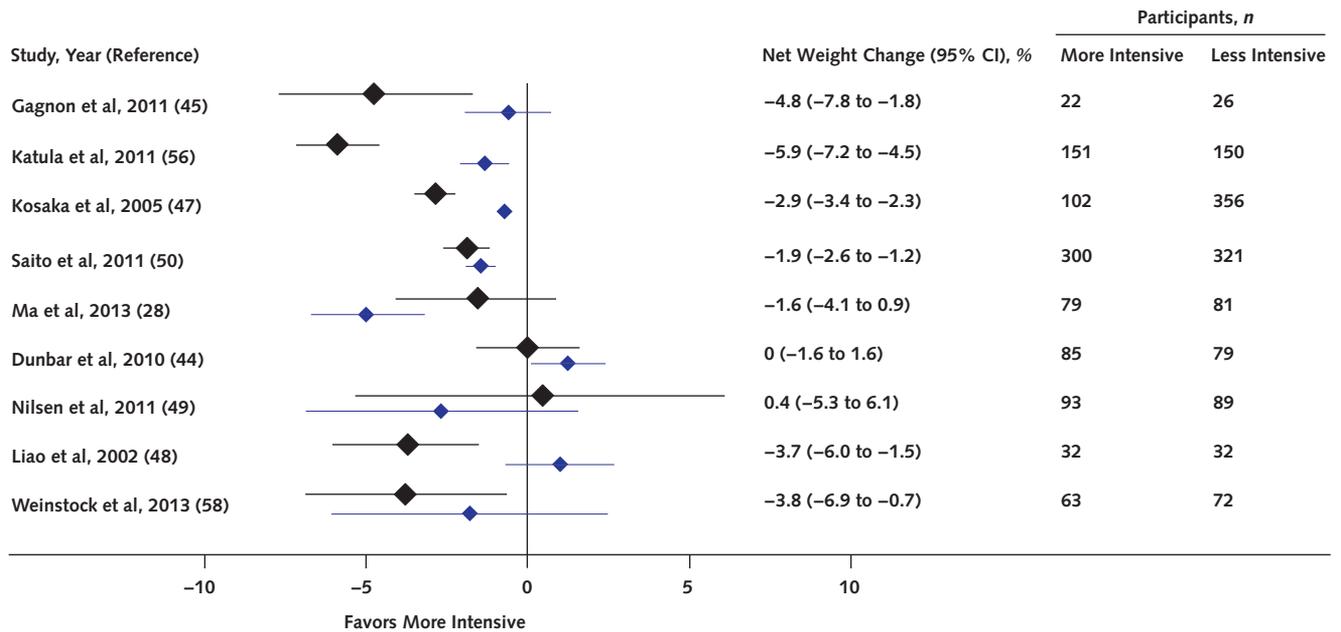
RD = risk difference; RR = risk ratio.

**Appendix Figure 3.** Forest plot of RR of reversion to normoglycemia in at-risk participants in more intensive versus less intensive combined diet and physical activity promotion programs.



RD = risk difference; RR = risk ratio.

**Appendix Figure 4.** Forest plot of net percentage change in weight (from baseline) in at-risk participants in more intensive versus less intensive combined diet and physical activity promotion programs.



Blue lines show percentage weight change in less intensive groups. The study by Ackermann and colleagues (35) was not included because it reported only that there was no significant difference between the more intensive and less intensive interventions at 12 mo (overall mean weight loss, 3.3% [CI, 2.7% to 3.9%];  $P = 0.26$  between interventions).