Circulating sex hormone binding globulin levels are modified with intensive lifestyle intervention, but their changes did not independently predict diabetes risk in the Diabetes Prevention Program

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ABSTRACT

Introduction  Sex hormone binding globulin (SHBG) levels are reported to be inversely associated with diabetes risk. It is unknown whether diabetes prevention interventions increase SHBG and whether resultant changes in SHBG affect diabetes risk. The purpose of this analysis was to determine whether intensive lifestyle intervention (ILS) or metformin changed circulating SHBG and if resultant changes influenced diabetes risk in the Diabetes Prevention Program (DPP).

Research design and methods  This is a secondary analysis from the DPP (1996–2001), a randomized trial of ILS or metformin versus placebo on diabetes risk over a mean follow-up of 3.2 years. The DPP was conducted across 27 academic study centers in the USA. Men, premenopausal and postmenopausal women without hormone use in the DPP were evaluated. The DPP included overweight/obese persons with elevated fasting glucose and impaired glucose tolerance. Main outcomes measures were changes in SHBG levels at 1 year and risk of diabetes over 3 years.

Results  ILS resulted in significantly higher increases (postmenopausal women: p<0.01) or smaller decrements (men: p<0.05; premenopausal women: p<0.01) in SHBG compared with placebo or metformin. Changes in SHBG were primarily attributable to changes in adiposity. There were no consistent associations of change in SHBG with the risk of diabetes by treatment arm or participant group.

Conclusions  Lifestyle intervention may be associated with favorable changes in circulating SHBG, which is largely due to changes in adiposity. Changes in circulating SHBG do not independently predict reductions in diabetes incidence.

INTRODUCTION

Sex hormone binding globulin (SHBG) levels have been reported to be inversely proportional to diabetes risk in multiple studies. In longitudinal observational studies, for example, higher baseline SHBG levels have been prospectively associated with lower risk of incident type 2 diabetes, and conversely lower baseline SHBG levels have been prospectively associated with a higher risk of incident diabetes, even after adjustment of other variables such as age, body mass index, and free sex steroids.1–3 Cross-sectional studies have likewise demonstrated inverse associations of circulating SHBG levels...
with indices of insulin resistance and with measures of glycemia.9–13 Short-term randomized trials have demonstrated an effect of interventions related to glucose homeostasis on circulating SHBG concentrations, including lifestyle changes, weight loss, and medications.9 14–17 No prior publications have evaluated whether changes in circulating SHBG levels as a result of such interventions impact the long-term risk of diabetes. The Diabetes Prevention Program (DPP), a 3-year multicenter randomized controlled clinical trial, compared intensive lifestyle intervention (ILS) with a targeted goal of weight reduction of at least 7% of initial body weight, metformin, and placebo on the risk of diabetes in overweight/obese individuals with elevated fasting glucose and glucose intolerance.18 Earlier reports from the DPP showed that ILS significantly increased SHBG among postmenopausal glucose-intolerant women in the DPP not using exogenous estrogen (n=382), while metformin did not affect endogenous sex hormones or SHBG.19 Contrary to expectations, analyses using the complete evaluable DPP population showed that baseline SHBG alone or SHBG single-nucleotide polymorphisms (SNPs) did not predict 3-year incident diabetes in the DPP population.20

In the follow-up analyses reported here, we further explore the relationship between circulating SHBG and diabetes risk. Specifically, we evaluated whether intervention-associated changes in SHBG, in turn, impacted diabetes risk in the high-risk DPP population. We did so by: (1) evaluating the extent to which diabetes prevention interventions (ILS, metformin, and placebo) impacted circulating SHBG levels in the DPP population, in subgroups defined by sex and menopausal status; (2) exploring concomitant determinants of SHBG change in the DPP (eg, adiposity, sex hormones, and glucose measures); and (3) examining the effects of change in SHBG on diabetes risk over an average of 3.2-year follow-up.

**RESEARCH DESIGN AND METHODS**

**Study design**

The DPP design, eligibility, and baseline characteristics have been reported previously.18 21 The DPP enrolled 3234 participants at least 25 years of age, body mass index (BMI) 24 kg/m² or higher (≥22 kg/m² in Asian-Americans), with fasting blood glucose 95–125 mg/dL (≤125 mg/dL in American Indians), and impaired glucose tolerance. Participants were randomly assigned to one of three treatments: an intensive lifestyle program (ILS) with a goal of at least 7% of weight loss through dietary modification and 150 min/week of moderate intensity exercise, metformin 850 mg twice daily; or matching placebo. Mean follow-up at the end of DPP was 3.2 years. All participants provided written informed consent. The current analyses are from participants who had baseline and 1-year SHBG blood samples and provided approval for the use of their blood samples in secondary analyses. In order to avoid potentially confounding effects of hormone therapy (ie, hormonal contraception or hormone replacement) on SHBG concentrations, users of these medications were excluded from the current analyses.

The final analysis included 2142 men and women with no self-reported hormone use in the DPP who had baseline and year 1 SHBG levels. Participants were classified prior to analysis as: men (n=886), premenopausal women with no self-reported hormone use (n=801), henceforth referred to as ‘premenopausal women’, and postmenopausal women with no self-reported hormone use (n=455), referred to as ‘postmenopausal women’. Postmenopausal status was defined as age at enrollment of 55 years, or self-reported natural or surgical menopause. Hormone use was assessed by self-reported medications.

**Diabetes ascertainment and glucose measures**

Diabetes status was determined by fasting glucose assessment every 6 months and/or by annual 75 g oral glucose tolerance test. Diabetes was defined as fasting glucose ≥126 mg/dL and/or 2-hour postchallenge glucose ≥200 mg/dL, confirmed. 1/fasting insulin was used as a surrogate measure of insulin sensitivity.22

**SHBG and sex hormone measurements**

SHBG and sex hormone measurements (dehydroepiandrosterone, dehydroepiandrosterone sulfate, estrone; estrone-S, estradiol, testosterone and dihydrotestosterone (DHT)) were conducted on morning fasting plasma samples, without regard to endogenous hormonal cycle. Sex hormones were measured by gas chromatography/mass spectrometry (Endoceutics, Quebec City, Canada), and SHBG was measured using ELISA (Bioline), as previously described.20 23 24

**Statistical analysis**

Three analysis groups (males, premenopausal women, and postmenopausal women) were considered. Quantitative characteristics with normal distributions were presented as means±SD and compared among groups using analysis of variance. Characteristics that were not normally distributed were presented as median (25th percentile and 75th percentile) and compared across groups using the Kruskal-Wallis test. Qualitative characteristics were compared among groups using the Pearson χ² test. 95% confidence intervals (CI) were calculated by bootstrapping. The statistical significance level was set at α=0.05 in all analyses. All P values were two-sided. All missing data were excluded from analyses. All analyses were performed using Stata (StataCorp, College Station, TX).
characteristics were presented as frequency (%), and the χ² test of association was used to compare these characteristics among groups.

The relationship of various characteristics with the change in SHBG from baseline to year 1 was evaluated with multivariable linear regression models. These models were tested separately within each of the different groups of interest, adjusted for treatment and baseline demographics (age, Caucasian race), lifestyle factors (smoking, alcohol intake, and leisure activity), glucose metabolism measures (fasting plasma glucose, 2-hour postchallenge glucose, and 1/fasting insulin), and adiposity measures (waist circumference and BMI), and sex hormone measures (estradiol, estrone, and testosterone). We identified a number of individually significant factors, which were then evaluated in a set of hierarchical models that were sequentially tested, as: model 1: baseline SHBG, treatment group, demographics, and lifestyle factors; model 2: additionally adjusted for baseline and change in sex hormones; model 3: further adjusted for baseline and change in adiposity; and then model 4: additionally adjusted for baseline and change in glucose parameters.

The association between the change in SHBG and the risk of diabetes was evaluated using Cox proportional hazards regression models, tested separately by treatment group in each of the three participant groups, and adjusted for demographic measures. Correlation analyses were conducted to evaluate the relationship between changes in SHBG and changes in circulating sex hormones and glucose measures. To further evaluate whether changes in SHBG contributed to centrally mediated downstream effects, mediation analyses were performed with change in SHBG as the predictor, change in Follicle-Stimulating Hormone (FSH) as the mediator, and change in circulating sex hormones or change in glucose measures (fasting glucose, hemoglobin A1c (HbA1c), 2-hour postchallenge glucose, homeostatic model assessment of insulin resistance (HOMA-IR), and 1/fasting insulin) as the outcome, after adjusting for treatment, age, and Caucasian race. Statistical analysis was performed with SAS V.9.4, and all tests were two sided done at a significance level of 0.05.

RESULTS
Study population
As described previously, the original DPP cohort consisted of two-thirds women and 45% ethnic minorities, with an average age of 51±11 years (mean±SD) at baseline. Baseline descriptive characteristics by groups (men: n=886; premenopausal women: n=801; postmenopausal women: n=455) are presented in table 1. Baseline sex hormones differed as expected between the three groups, with postmenopausal women having lower levels of circulating sex hormones relative to premenopausal women.Baseline SHBG was lowest in postmenopausal women (median (Q1, Q3) of 34.9 (25.9, 46.2) nmol/L), with a median value of 45.2 (30.9, 68.6) nmol/L in premenopausal women and 39.8 (26.9, 56.3) nmol/L in men (table 1).

Effect of DPP interventions (ILS, metformin, and placebo) on change in SHBG at 1 year
Figure 1 and online supplemental table 1 show the unadjusted mean of SHBG at baseline and year 1 by treatment arm. In men, SHBG decreased from baseline to year 1 in all three arms, but ILS was associated with the smallest decrement (−9.91±22.27 nmol/L) (mean±SD) compared with metformin (−12.40±20.12 nmol/L) or placebo (−11.97±17.30 nmol/L) (p<0.01 ILS vs metformin and p<0.01 ILS vs placebo) (figure 1, online supplemental table 1). Overall, changes in SHBG in the ILS participants were significantly different from changes in SHBG in the placebo or metformin groups, with either increases or smaller decrements comparatively.

In a model adjusting for treatment assignment, analysis group (ie, men, premenopausal women, and postmenopausal women), change in SHBG, age at randomization, and race (Caucasian vs not), the interaction of treatment with analysis group was statistically significant (p=0.047), suggesting the effect of treatment on SHBG was different in the three analysis groups. When tested with in each analysis group, in models adjusting for treatment, change in SHBG, age at randomization, and race (Caucasian vs not), the interaction of change in SHBG with treatment arm was statistically significant only in postmenopausal women (p=0.023), suggesting that in this group the effect of change in SHBG on the risk of diabetes is different among the three treatment arms; in men and premenopausal women, the interaction of change in SHBG with treatment arm was not significant.

Determinants of change in SHBG
The observed differences between treatment arms suggest a number of hypothesized mediators that might drive the changes in SHBG. Therefore, hierarchical multivariable linear regression models of changes in SHBG from baseline to year 1 were conducted evaluating potential determinants of this change (figure 2, online supplemental table 2). The ILS group continued to have less decrease (men and premenopausal women) or greater increase (postmenopausal women) in SHBG in all groups after adjustment for demographics and lifestyle factors (model 1), which were slightly attenuated following additional adjustment for sex hormones (model 2) and completely eliminated following adjustment for adiposity measures.
Table 1  Descriptive characteristics of participants for current analyses at DPP baseline

|                       | Men (n=886) | Premenopausal women (n=801) | Postmenopausal women (n=455) | P value |
|-----------------------|------------|----------------------------|-----------------------------|---------|
| Age (years)           | 54.0±10.9  | 42.3±5.9                   | 56.7±9.6                    | <0.001* |
| Race                  |            |                            |                             |         |
| Caucasian             | 521 (59)   | 400 (50)                   | 226 (50)                    | <0.001† |
| Other                 | 365 (41)   | 401 (50)                   | 229 (50)                    |         |
| Alcohol (g/day)       | 0.94 (0.00, 5.32) | 0.00 (0.00, 0.94) | 0.00 (0.00, 0.94) | <0.001‡ |
| Leisure activity (Met-hour/week) | 14.9 (5.7, 27.6) | 7.7 (3.0, 16.0) | 7.8 (2.7, 16.5) | <0.001‡ |
| Weight (kg)           | 98.4±19.3  | 95.2±21.1                  | 91.9±19.5                   | <0.001* |
| BMI (kg/m²)           | 32.0±5.5   | 35.8±7.0                   | 34.9±6.7                    | <0.001* |
| Waist circumference (cm) | 108.0±13.3 | 104.5±15.1                 | 104.6±14.3                  | <0.001* |
| FPG (mg/dL)           | 108.4±8.2  | 105.7±8.0                  | 106.6±7.7                   | <0.001* |
| 2-hour postchallenge glucose (mg/dL) | 163.9±17.0 | 164.0±17.0                 | 163.7±17.1                  | <0.001* |
| 1/fasting insulin (mL/µU) | 0.0435 (0.0313, 0.0625) | 0.0385 (0.0278, 0.0556) | 0.0417 (0.0303, 0.0626) | <0.001‡ |
| DHEA (pg/mL)          | 3042 (2461, 4077) | 2117 (1412, 3120) | 1600 (1030, 2400) | <0.001‡ |
| DHEA-S (ng/mL)        | 1000 (602, 1545) | 870 (556, 1231) | 588 (359, 909) | <0.001‡ |
| Estrone (pg/mL)       | 38.2 (31.4, 47.0) | 85.2 (54.9, 128.3) | 70.9 (39.0, 118.7) | <0.001‡ |
| Estrone-S (pg/mL)     | 626 (426, 910) | 1186 (604, 2204) | 413 (204, 740) | <0.001‡ |
| Estradiol (pg/mL)     | 22.4 (18.2, 27.3) | 53.6 (23.1, 104.7) | 9.5 (5.9, 17.2) | <0.001‡ |
| Testosterone (pg/mL)  | 3165 (2476, 3960) | 176 (134, 254) | 140 (100, 207) | <0.001‡ |
| DHT (pg/mL)           | 236 (172, 321) | – | – |         |
| SHBG (nmol/L)         | 39.8 (26.9, 56.3) | 45.2 (30.9, 68.6) | 34.9 (25.9, 46.2) | <0.001‡ |

Values are expressed as n (%), mean±SD, or median (25th percentile, 75th percentile).
Comparisons across the three groups are presented.
*P value from analysis of variance.
†P value from χ² test.
‡P value from Kruskal-Wallis non-parametric test.
BMI, body mass index; DHEA, dehydroepiandrosterone; DHEA-S, dehydroepiandrosterone-sulfate; DHT, dihydrotestosterone; DPP, Diabetes Prevention Program; FPG, fasting plasma glucose; SHBG, sex hormone binding globulin.
Clinical care/Education/Nutrition

Effects of change in SHBG on diabetes risk
The relationship between 1-year change in SHBG and diabetes risk was inconsistent across treatment arms and participant subgroups (table 2). A significant effect of change in SHBG on diabetes risk was seen in men in the placebo group, HR 0.77, 95% CI 0.63 to 0.95, p=0.01, for each SD relative increase in SHBG. In addition, a significant effect of change in SHBG on diabetes risk was seen in postmenopausal women in the metformin group (HR 1.61, 95% CI 1.05 to 2.49, p=0.03). However, these were not consistent findings across treatment arms and participant groups (table 2).

Correlation of SHBG changes with sex hormones and glucose measures
Changes in SHBG were positively correlated with changes in peripheral sex hormones (change in estradiol and change in testosterone), and inversely correlated with glucose measures (fasting glucose, HbA1c, 2-hour postchallenge glucose), negatively correlated with HOMA-IR and positively correlated with 1/fasting insulin (table 3). These effects were consistent across treatment arms (table 3) and were not centrally mediated by FSH (data not shown).

DISCUSSION
In summary, in the DPP, lifestyle intervention exerted effects to increase or attenuate the decline in SHBG levels, manifested differently within each subgroup. In postmenopausal women, for example, ILS resulted in an increase in SHBG, consistent with prior analyses within the DPP, whereas in men and in premenopausal women in the ILS arm, the decrement in SHBG was less than that seen in the other treatment arms. Metformin had no impact compared with placebo on SHBG in any of the groups. Through multivariable regression models, we found that a large part of the effect of ILS on SHBG levels was attributable to reductions in adiposity, with modest effects of sex hormones, glucose measures, and their changes. Given that low SHBG levels have been reported to be associated with diabetes risk and that low SHBG levels are seen in conditions associated with obesity and insulin resistance (eg, diabetes, polycystic ovary syndrome, and fatty liver disease), we explored whether the changes in SHBG contributed to reduced diabetes incidence. However, despite these apparently beneficial changes in SHBG, we did not see consistent evidence for SHBG mediating the treatment effects on diabetes prevention. Furthermore, while changes in SHBG were positively correlated with changes in circulating hormones and 1/fasting insulin, and inversely associated with changes in glucose (fasting and 2-hour postchallenge) and HOMA-IR, mediation analyses did...
not support a centrally mediated effect of SHBG on these outcomes. Taken together, these observations suggest that changes in SHBG reflect the overall metabolic and hormonal milieu, in particular the effects of weight loss, but are not directly influencing risk of development of diabetes.

Our findings are also consistent with our earlier analyses of baseline SHBG and SHBG SNPs associated with diabetes outcomes and risk of diabetes in the DPP. As earlier reported, while baseline SHBG was cross-sectionally associated with some indicators of insulin resistance and diabetes risk (inverse fasting insulin, insulinogenic index, and waist circumference), SHBG concentration at baseline was not associated with diabetes risk in any of the participant groups evaluated. Furthermore, there was no evident association of the SHBG SNPs and diabetes risk in the DPP population.20

Table 2  Association between SD change in SHBG and 3-year diabetes risk, adjusted for age at randomization and Caucasian race

|                | HR  | Lower 95% CI | Upper 95% CI | P value |
|----------------|-----|--------------|--------------|---------|
| Placebo        |     |              |              |         |
| Males (SD=17.3) | 0.771 | 0.629        | 0.946        | 0.013   |
| Premenopausal  | 1.163 | 0.888        | 1.523        | 0.272   |
| Postmenopausal | 0.882 | 0.676        | 1.151        | 0.356   |
| Metformin      |     |              |              |         |
| Males (SD=20.1) | 0.836 | 0.677        | 1.034        | 0.099   |
| Premenopausal  | 0.949 | 0.710        | 1.269        | 0.725   |
| Postmenopausal | 1.614 | 1.046        | 2.491        | 0.030   |
| Intensive lifestyle intervention |     |              |              |         |
| Males (SD=22.3) | 0.787 | 0.576        | 1.074        | 0.131   |
| Premenopausal  | 0.876 | 0.661        | 1.162        | 0.359   |
| Postmenopausal | 0.597 | 0.303        | 1.177        | 0.137   |

SHBG, sex hormone binding globulin.
To date, it has not been clear whether low SHBG levels represent a biomarker of metabolic abnormality and diabetes risk or are perhaps somehow contributory and causative of disease. Our data would support that dynamic changes in SHBG reflect the changes in the surrounding metabolic environment. Several studies support this line of thought. First, multiple studies have demonstrated that levels of SHBG, insulin sensitivity, and metformin responsiveness are strongly linked. This may be because metformin primarily suppresses hepatic glucose production and is not generally regarded as a potent insulin sensitizer. In contrast, intervention studies with insulin sensitizers (thiazolidinediones), have demonstrated responsive increases in SHBG. In a prospective randomized controlled trial of women with polycystic ovary syndrome (PCOS), for example, serum SHBG levels were correlated with glucose disposal rate (insulin sensitivity), as assessed by hyperinsulinemic euglycemic clamp, and increased significantly during treatment with pioglitazone, a treatment known to directly modulate insulin sensitivity. Furthermore, the improvement in glucose disposal and insulin sensitivity was directly associated with increases in circulating SHBG. In another study evaluating rosiglitazone on metabolic and ovarian effects in polycystic ovary syndrome (PCOS), an increase in SHBG was demonstrated in response to rosiglitazone treatment. These changes in SHBG were also shown to be associated with improvements in insulin sensitivity.

### Table 3: Correlation of SHBG changes with changes in hormone and glucose measures, overall and by treatment arm

| Treatment Arm | Change in SHBG (overall) – correlation coefficient, p(n) | Change in SHBG (placebo) – correlation coefficient, p(n) | Change in SHBG (metformin) – correlation coefficient, p(n) | Change in SHBG (intensive lifestyle intervention) – correlation coefficient, p(n) |
|---------------|----------------------------------------------------------|----------------------------------------------------------|-------------------------------------------------------------|---------------------------------------------------------------------------------|
|               | Change in FSH | Change in estradiol | Change in testosterone | Change in fasting glucose | Change in HbA1c | Change in 2-hour glucose | Change in HOMA-IR | Change in 1/fasting insulin |
|               | -0.19 | 0.08 | 0.12 | -0.20 | -0.12 | -0.11 | -0.13 | 0.15 |
|               | p<0.001 (990) | p<0.001 (1924) | p<0.001 (1996) | p<0.001 (2138) | p<0.001 (2122) | p<0.001 (2097) | p<0.001 (2097) | p<0.001 (2097) |
|               | -0.30 | 0.12 | 0.08 | -0.19 | -0.12 | -0.06 | -0.10 | 0.10 |
|               | p<0.001 (351) | p<0.001 (649) | p<0.001 (674) | p<0.001 (725) | p<0.001 (716) | p=0.097 (724) | p=0.006 (708) | p=0.010 (708) |
|               | -0.18 | 0.02 | 0.06 | -0.11 | -0.05 | -0.03 | -0.06 | 0.09 |
|               | p=0.002 (310) | p=0.003 (636) | p=0.001 (656) | p=0.003 (698) | p=0.159 (696) | p=0.356 (698) | p=0.129 (684) | p=0.024 (684) |
|               | -0.12 | 0.10 | 0.07 | -0.24 | -0.07 | -0.13 | -0.16 | 0.18 |
|               | p=0.027 (329) | p=0.016 (639) | p=0.063 (666) | p=0.001 (715) | p=0.057 (710) | p<0.001 (715) | p<0.001 (715) | p<0.001 (715) |

FSH, Follicle-Stimulating Hormone; HbA1c, hemoglobin A1c; HOMA-IR, homeostatic model assessment of insulin resistance; SHBG, sex hormone binding globulin.
Consistent with our findings that SHBG may reflect the metabolic milieu and changes as such, it is of interest to note that the hepatic environment has also been directly implicated in regulating SHBG. Selva et al. previously showed that monosaccharide (glucose and fructose) induced hepatic lipogenesis reduced SHBG production by downregulating hepatocyte nuclear factor 4α (HNF-4α) levels, a key transcription factor regulating hepatic expression of SHBG. Supporting this, Winters et al. evaluated SHBG gene expression in human liver samples and found that SHBG mRNA was a strong predictor of circulating SHBG levels. They described an inverse association between hepatic triglyceride content and SHBG mRNA and serum SHBG, with a suggestion that the low level of SHBG mRNA was largely due to a low level of HNF-4α mRNA expression in the liver, which is also reduced in insulin resistance. Thus, it is plausible that improvements in insulin sensitivity, as seen with weight loss and therapies that directly modify insulin sensitivity, may have an impact on the liver, which then affects expression of SHBG mRNA and production of SHBG.

In contrast, there is emerging evidence that SHBG may play a causative role in disease and may not merely be an ‘innocent bystander’. Sáez-López et al. recently described a significant inverse relation between SHBG mRNA expression and hepatic triglyceride content, as well as levels of acetyl-coenzyme A carboxylase, a key lipogenic enzyme, in liver samples obtained from obese humans with non-alcoholic fatty liver disease undergoing bariatric surgery. Furthermore, the authors found that SHBG overexpression in cultures of human hepatic (HepG2) cells was able to abrogate the increase in multiple hepatic lipogenic enzymes in the liver when triggered under high-glucose culture conditions. Although this was studied in fatty liver, it is possible that SHBG is not only a biomarker, but it may independently contribute to the pathogenesis of metabolic disease.

In order to estimate a causal role of circulating SHBG for type 2 diabetes, Wang et al. applied quantitative nuclear magnetic resonance metabolomics in three Finnish population-based cohorts to profile circulating lipids and metabolites and their association with SHBG. Higher SHBG levels were associated with a more favorable cardiometabolic risk profile, and SHBG was predictive of future insulin resistance and type 2 diabetes. The observed association of SHBG with type 2 diabetes (OR=0.83 per 1 SD) was less than that seen in prospective observational associations by meta-analysis (HR 0.47). These results suggest that circulating SHBG may have a minor direct contributory role in the development of type 2 diabetes but is more likely largely reflective of other factors.

There are several strengths and limitations to these analyses. First, a prospective, controlled evaluation of changes in SHBG by sex and menopausal status in the very well characterized DPP population provided the opportunity to directly assess whether the study interventions affected SHBG and what other factors contributed to those changes. In addition, our sample size is commensurate with other studies in the literature for the relevant constituent grouping to have provided meaningful analyses and comparative results. However, given the predefined criteria of the DPP to identify those individuals already at high risk of development of diabetes based on glucose and weight measures, we may have been limited in seeing additional impact of SHBG on risk. Furthermore, SHBG and sex hormone measurements were conducted on samples that were not timed to the endogenous hormonal cycle, and hormone use was assessed by self-report. Randomization may help to minimize potential influence of this and other unanticipated factors that may regulate SHBG.

In summary, in the DPP, ILS was consistently associated with changes in circulating SHBG levels compared with placebo or metformin, specifically increased levels in postmenopausal women, and attenuation of decrease in men and premenopausal women. The effects of ILS on SHBG seemed largely due to changes in adiposity but may also be influenced by other changes (e.g., glucose measures and sex steroids). The observed changes in SHBG related to the interventions did not consistently translate to changes in diabetes risk.

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Contributors The manuscript was conceived by VRA and KJM, with manuscript questions and analytic plan designed by VRA, CAC, CK, SLE and KJM. VRA wrote the manuscript, interpreted the data, critically reviewed and revised the manuscript. CAC contributed to writing, data analysis, data interpretation, critical review and revision, SLE contributed to data analysis, data interpretation, critical review and revision. LP, CK, SHG, EH and KJM contributed to data interpretation, critical review and revision. All authors had access to the data and all authors agreed to submit the final manuscript. At the time of publication KJM was a full-time employee of Eli Lilly and Company. However, prior to employment at Eli Lilly and Company, KJM served as Principal Investigator for this NIH funded study. As such, data collection occurred prior to and independent of this employment and therefore, data analysis and preparation of the manuscript were independent of Eli Lilly and Company.

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