Vertical sleeve gastrectomy improves glucose and lipid metabolism and delays diabetes onset in UCD-T2DM rats.
Vertical Sleeve Gastrectomy Improves Glucose and Lipid Metabolism and Delays Diabetes Onset in UCD-T2DM Rats

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Vertical sleeve gastrectomy (VSG) has gained interest as a low morbidity bariatric surgery, which is effective in producing weight loss and causing type 2 diabetes resolution. However, the efficacy of VSG to prevent the onset of type 2 diabetes has not been previously investigated. VSG or sham surgery was performed on 2-month-old prediabetic male University of California Davis-type 2 diabetes mellitus rats. Sham-operated animals were either sham-operated *ad libitum* fed (S-AL) or were weight-matched to VSG-operated animals (S-WM). Diabetes onset was determined by weekly nonfasting blood glucose measurements. Animals underwent oral glucose tolerance tests at 1 and 4 months after surgery and indirect calorimetry at 1.5 months after surgery. VSG surgery significantly delayed diabetes onset compared with both S-AL and S-WM animals. VSG-operated animals ate 23% less and weighed 20% less than S-AL. Energy expenditure did not differ between VSG-operated animals and controls. Results from the oral glucose tolerance tests demonstrate improved glucose tolerance and islet function in VSG-operated animals compared with S-AL and S-WM. Nutrient-stimulated glucagon-like peptide (GLP)-1, GLP-2, and peptide YY excursions were greater in VSG-operated animals. VSG surgery resulted in decreased fasting plasma insulin, ghrelin and lipid concentrations, and markedly higher fasting plasma adiponectin and bile acid concentrations, independent of body weight. Increases of circulating bile acid concentrations were due to selective increases of taurine-conjugated bile acids. Thus, VSG delays type 2 diabetes onset in the University of California Davis-type 2 diabetes mellitus rat, independent of body weight. This is potentially mediated by increases of circulating bile acids, adiponectin, and nutrient-stimulated GLP-1 secretion and decreased circulating ghrelin concentrations. (*Endocrinology* 153: 3620–3632, 2012)

Bariatric surgery is currently the most effective long-term treatment for obesity and often results in resolution of type 2 diabetes (1) and may also prevent or delay the onset of diabetes (2, 3). However, the mechanisms by which this occurs remain undefined. Identification of these mechanisms may provide new therapeutic targets for the treatment and prevention of type 2 diabetes. Vertical sleeve gastrectomy (VSG) has gained interest as a low morbidity bariatric surgery that is highly effective in producing weight loss and improving glucose homeostasis. VSG involves removal of the greater curvature of the stomach and thus has been considered to be primarily a restrictive surgery. However, retrospective studies have revealed that VSG reverses diabetes in 80–84% of patients with type 2 diabetes (4–6) with improvements of circulating insulin and glucose concentrations observed as early as 3 d postoperatively and before any significant weight loss (7). Interestingly, VSG has been shown to produce diabetes re-

Abbreviations: AG, Acyl-ghrelin; AUC, area under the curve; DAG, desacyl-ghrelin; FBS, fetal bovine serum; GLP, glucagon-like peptide; IT, ileal interposition; OGTT, oral glucose tolerance test; PYY, peptide YY; PDS, polydioxanone; RYGB, Roux en Y gastric bypass; S-AL, sham-operated *ad libitum* fed; S-WM, sham-operated weight matched; TBP, TATA box-binding protein; TG, triglyceride; UCD-T2DM, University of California Davis-type 2 diabetes mellitus; VSG, vertical sleeve gastrectomy.
mission and improved glucose homeostasis to a similar degree as Roux en Y gastric bypass (RYGB) in both rodent and human clinical studies (4, 8). Furthermore, recent clinical studies in humans have shown that, similar to RYGB, metabolic improvements after VSG may be related to increases of nutrient-stimulated glucagon-like peptide (GLP)-1 and peptide YY (PYY) secretion (9). In addition, VSG results in marked decreases of circulating ghrelin concentrations, which have been postulated to contribute to greater early excess body weight loss (10).

In addition to the well-described effects of ghrelin to increase appetite, ghrelin also influences fat mass, growth hormone secretion, and glucose homeostasis. Ghrelin is primarily produced in the gastric mucosa and its receptor (GH secretagogue receptor-1a) is expressed on hypothalamic cells, pituitary, liver, adipocytes, and pancreas (11). Furthermore, GH secretagogue receptor-1a and ghrelin are both expressed in pancreatic α-, β-, and ε-cells (12–14). Recent studies suggest that excess ghrelin production could contribute to the development of type 2 diabetes, because ghrelin has been shown to impair glucose-stimulated insulin secretion and promote insulin resistance (15–17).

GLP-1 and PYY are secreted by L cells, located primarily in the distal small intestine, in response to direct stimulation by nutrients and by vagal stimulation (18, 19). PYY acts to decrease food intake and thus maintain weight loss (20). GLP-1 acts to decrease food intake, increase glucose-stimulated insulin secretion and improve insulin sensitivity, and may preserve islet integrity (18).

In the present study, we used the University of California Davis-type 2 diabetes mellitus (UCD-T2DM) rat model of type 2 diabetes to investigate the effects of VSG on the onset of diabetes. The UCD-T2DM rat model develops adult-onset polygenic obesity, insulin resistance, and hyperglycemia, without a defect in leptin signaling, making this model ideal for investigating the efficacy of VSG to delay type 2 diabetes onset (21). Previous rodent studies of VSG surgery have used models of obesity such as Zucker rats (22) and high-fat-fed Long-Evans rats (8). To our knowledge, this is the first study of the effects of VSG in an animal model of type 2 diabetes. Here, we report that VSG significantly delayed diabetes onset and was accompanied by significant improvements of insulin secretion and glucose and lipid metabolism. Furthermore, VSG-operated animals exhibited marked increases of nutrient-stimulated GLP-1, GLP-2, and PYY secretion, decreased fasting plasma ghrelin concentrations, and increased fasting plasma adiponectin and bile acid concentrations, which are likely contributors to the delay in diabetes onset.

Materials and Methods

Diet and animals

Male UCD-T2DM rats were individually housed in hanging wire cages in the animal facility in the Department of Nutrition at the University of California Davis and maintained on a 14-h light, 10-h dark cycle. At 2 months of age rats underwent sham or VSG surgery. Sham-operated animals were then divided into sham-operated ad libitum-fed (S-AL) or sham-operated weight-matched (S-WM) groups. Animals were followed until at least 8 months of age to determine the time to diabetes onset. Baseline body weights, measured on the day of surgery, were 385 ± 10, 385 ± 6, and 385 ± 7 g in S-AL (n = 14), S-WM (n = 16), and VSG-operated animals (n = 13), respectively. All animals received ground chow (no. 5012; Ralston Purina, Belmont, CA). Food intake and body weight were measured three times a week in S-AL and VSG-operated animals, and S-WM animals received daily rations of food to match their body weights with the VSG-operated animals. Nonfasting blood glucose was measured weekly with a glucometer (One-Touch Ultra; LifeScan, Milpitas, CA) at 1400–1600 h. Diabetes onset was defined as a nonfasted blood glucose value above 11.1 mmol/liter (200 mg/dl) for two consecutive weeks. Indirect calorimetry was performed at 1.5 months after surgery using an Integra ME system (AccuScan, Columbus, OH). The experimental protocols were approved by the University of California Davis Institutional Animal Care and Use Committee.

Bariatric procedures, including VSG surgery, can result in malabsorption of certain vitamins and minerals, including vitamin B12, often necessitating supplementation (23). In the rat, vitamin B12 deficiency can result from removal of the glandular portion of the stomach at 1–3 months after surgery, leading to the subsequent development of anemia. The glandular portion of the stomach is responsible for producing intrinsic factor, which is necessary for vitamin B12 absorption (24). To control for this, hematocrit was monitored, and when mild anemia was noted, vitamin B12 supplementation was initiated in VSG-operated animals. At 5 months after surgery, mild anemia was first noted in the VSG-operated animals (hematocrit, 34 ± 2%; rat reference range, 35–45%) (25). VSG-operated animals received 50 μg of cyanocobalamin sc once weekly thereafter (25). After one month of treatment, hematocrit values in VSG-operated animals were normalized to 40 ± 2% and remained stable thereafter.

VSG surgery

Rats were placed on a liquid diet (Boost, Novartis, Minneapolis, MN) 4 d before surgery and for 14 d after surgery and received enrofloxacin (20 mg/kg, sc once daily) and meloxicam (2 mg/kg sc once daily). Anesthesia was induced and maintained with isoflurane (2–5%). A midline abdominal incision was made, and connective tissue attachments to the liver and spleen were transected to allow isolation of the stomach outside the abdominal cavity. The stomach was packed-off with gauze sponges and 4–0 polydioxanone (PDS) suture (Ethicon, Corne- lia, GA) was used to create a line of sutures encompassing both gastric walls just below the intended incision. Hemostats were placed just above the intended incision, and approximately 70% of the stomach (including the entire fundus) was removed by cutting along the greater curvature between the suture line and the hemostats to create a tubular remnant connecting the esoph-
agus and pylorus. A second line of sutures was placed using 6–0 PDS (Ethicon) to reinforce apposition of the gastric mucosa. The gastric remnant was lavaged and placed back into the abdominal cavity. The abdominal cavity was closed in two layers using 4–0 PDS (Ethicon).

Sham-operated animals were treated in similar manner as the VSG group. Sham surgeries were performed by making a laparotomy incision and isolating the stomach outside of the abdominal cavity. A simple continuous pattern of suture was placed through one layer of the greater curvature of the stomach in the same location as the VSG-operated animals using 6–0 PDS (Ethicon) along both gastric walls. The stomach was lavaged and returned to the abdominal cavity.

**Oral glucose tolerance tests (OGTT)**

At 1 and 4 months after surgery, OGTT were performed. Animals were fasted overnight (12 h) and then received a 50% dextrose solution (1 g/kg body weight) by oral gavage. Blood was collected from the tail for measurement of glucose and insulin concentrations. A second aliquot of blood was placed in tubes containing EDTA, aprotinin, and a dipeptidyl peptidase-IV inhibitor and analyzed for GLP-1 and PYY. Serum glucose was measured using an enzymatic colorimetric assay for glucose (Thermo DMA, Louisville, CO). Serum insulin was measured by ELISA (Millipore, St. Charles, MO). Total GLP-1 was measured by sandwich electrochemiluminescence immunoassay (Meso Scale Discovery, Gaithersburg, MA). Plasma PYY was measured by RIA (Millipore).

**Monthly fasted hormone and metabolite profiles**

Baseline and monthly blood samples were collected after an overnight (12 h) fast from the tail. Plasma was assayed for glucose, insulin, triglycerides (TG), cholesterol, leptin, adiponectin, and ghrelin. Plasma glucose, cholesterol, and TG were measured using enzymatic colorimetric assays (Thermo DMA; L-type TG H kit, Wako Chemicals USA, Inc., Richmond, VA). Leptin and adiponectin were measured with rodent/rat-specific RIA (rat leptin, mouse adiponectin; Millipore). Insulin was measured by ELISA (Millipore). Ghrelin secretion was measured by sandwich electrochemiluminescence immunoassay (Meso Scale Discovery).

**Monthly fasting plasma bile acid profiles**

Monthly bile acid profiles were analyzed as previously described (26, 27). An internal standard mixture of D4-cholate, D4-β-muricholate, D4-α-muricholate, D4-chenoxycholate, D4-deoxycholate, D4-hydoxycholate, D4-ursodeoxycholate, and D4-lithocholate, and their tauroconjugated and glycoconjugated counterparts, was prepared in acetonitrile. Eluted bile salts were detected by negative ion, electrospray mass spectrometry on a linear trap linear ion trap mass spectrometer (Thermo Finnigan, San Jose, CA) using a data-dependent stage of detection triggering a qualitative 0 m/z neutral loss scan at 27% normalized collision energy. Peak areas from initial scans of individual bile salts at 314.3, 498.3, 464.3, 447.3, 411.3, 407.3, 395.3, 393.3, and 391.3 m/z were integrated, and response factors were defined by peak area ratios of analytes to that of internal, deuterated standards. The response factors were read against those obtained from standard curves in surrogate matrix, and molar levels of serum or plasma levels were interpolated from standard curves. Response factors for all samples were comprised of peak area ratios of nonlabeled salts normalized to the stable-labeled counterparts. Concentrations were interpolated by linear regression from curves of known standards.

**Cell culture**

3T3-L1 cells were maintained in high glucose (25 mM) DMEM supplemented with 10% newborn calf serum and penicillin-streptomycin. When confluent (designated d 0), cells were switched to high insulin (1.7 μM), high glucose DMEM supplemented 10% fetal bovine serum (FBS). Differentiation of 3T3-L1 preadipocytes was induced at d 2 by switching to high insulin, high glucose DMEM supplemented with 10% FBS, 1 μM dexamethasone, and 0.5 mM 3-isobutyl-1-methylxanthine. After 48 h, the medium was switched back to high insulin, high glucose DMEM supplemented 10% FBS. Differentiated adipocytes were exposed to one of three doses of acyl-ghrelin (AG) or desacyl-ghrelin (DAG) (Bachem; Torrance, CA) calculated to correspond with the concentrations seen in the VSG-operated animals (AG low, 22 pmol/liter; and DAG low, 89 pmol/liter) and the S-WM animals (AG medium, 65 pmol/liter; and DAG medium, 260 pmol/liter). A high dose (AG high, 130 pmol/liter; and DAG high, 520 pmol/liter) corresponding to a 2-fold higher dose than the medium dose was also included. The concentration of AG present in the circulation was estimated using the 1-month total ghrelin concentration data set and previously published data demonstrating a 1:5 ratio of acyl to total ghrelin in the circulation in rats (28). Cells were incubated with the appropriate dose of ghrelin for 24 h, and then media were collected for assessment of adiponectin secretion by RIA, and mRNA was extracted from cells using TRIzol reagent (Invitrogen, Carlsbad, CA) for assessment of adiponectin mRNA expression. CDNA was generated using high-capacity cDNA RT kit (Applied Biosystems, Foster City, CA). Adiponectin mRNA levels were determined by real-time PCR (iCycler; Bio-Rad, Hercules, CA) and normalized to TATA box-binding protein (TBP). Primers used were adiponectin forward, GTTGCAAGCTCTCCTGTTCC and adiponectin reverse, CTCCAGGAGTGCCATCTCT; TBP forward, GAAG CTGGGCTACATTTCCAG and TBP reverse, CCCCTTGTA CCCCCAATAAT.

**Statistics and data analysis**

Data are presented as mean ± SEM. Statistical analyses were performed using GraphPad Prism 4.00 for Windows (GraphPad Software, San Diego, CA). The incidence of diabetes was analyzed by log-rank testing of Kaplan-Meier survival curves. All other data were analyzed by two-factor (time × treatment) repeated measures ANOVA followed by post hoc analysis with Bonferroni’s multiple comparison test or Student’s t test as indicated. Differences were considered significant at P < 0.05. Values are expressed as mean ± SEM.

**Results**

**VSG surgery delays the onset of diabetes**

Compared with S-AL animals, the onset of diabetes was delayed in S-WM and VSG-operated animals by approximately 1 and 3.5 months, respectively. Furthermore, the age of diabetes onset was delayed in VSG-operated ani-
mals compared with S-WM animals by approximately 2.5 months. The mean ages of diabetes onset were 6.1 ± 0.5, 7.0 ± 0.2, and 9.6 ± 0.2 months in S-AL, S-WM, and VSG-operated animals, respectively (P < 0.05 VSG compared with S-WM and S-AL) (Table 1). By 8 months of age, none of the VSG-operated animals had become diabetic, whereas 3/16 S-WM and 11/14 S-AL animals were diabetic. Thus, within the first 8 months of age, VSG significantly delayed diabetes onset compared with both the S-AL and S-WM groups (P < 0.05) (Fig. 1A). At 1 yr of age, S-WM and VSG-operated animals both exhibited significantly delayed diabetes onset compared with S-AL animals (P < 0.001), but at this small sample size and longer time course, the time of onset was not significantly different between S-WM and VSG groups by Kaplan-Meier analysis. Similarly, diabetes-free days up to 1 yr of age were significantly greater in S-WM and VSG-operated animals compared with S-AL animals (P < 0.001) but did not differ between S-WM and VSG-operated animals (Table 1). The delay in the onset of diabetes corresponded with lower fasting serum glucose concentrations in S-WM and VSG-operated animals compared with S-AL animals (Fig. 1B).

As expected, energy intake and body weight were higher in S-AL animals compared with S-WM and VSG-operated animals (P < 0.001) (Fig. 2, A and B). Energy intake was reduced by approximately 23% in VSG-operated animals compared with S-AL. Energy intake and body weight did not differ between S-WM and VSG-operated animals. Energy expenditure in VSG-operated animals did not differ compared with S-AL or S-WM groups (Fig. 2, C and D). However, energy expenditure was approximately 5% lower in S-WM animals compared with S-AL (P < 0.05). The observed decrease of energy expenditure with restriction of energy intake is similar to that reported in previous rodent and primate studies examining the effect of chronic energy intake restriction on energy expenditure (29, 30). It is interesting that surgically induced energy restriction did not produce a similar effect and suggests a mechanism by which bariatric surgery patients are better able to defend body weight loss after bariatric surgery compared with dieting alone (31).

**VSG surgery improves circulating lipids and hormones**

Monthly fasting plasma insulin remained significantly lower in VSG-operated animals compared with both S-AL and S-WM animals up to 3 months after surgery (P < 0.001), at which point, insulin values began to decline in S-AL animals due to the onset of diabetes and progressive β-cell decompensation (Fig. 3A). Circulating insulin concentrations continued to be lower in VSG-operated animals compared with S-WM up to 5 months after surgery, suggesting an improvement of insulin sensitivity, independent of body weight (P < 0.01). Fasting plasma leptin concentrations remained lower in VSG-operated animals compared with S-WM up to 5 months after surgery (P < 0.01) (Fig. 3B). We have previously reported that fasting plasma leptin concentrations increase in the UCD-T2DM rat before diabetes onset due to increasing adiposity and increasing circulating insulin concentrations. Conversely, fasting plasma leptin concentrations decrease after diabetes onset in the UCD-T2DM rat due to decreasing adipose mass and decreasing circulating insulin concentrations (21). Leptin concentrations in S-AL animals began to decline at 3 months after surgery, likely due to the increased prevalence of diabetes and corresponding decreases of circulating insulin concentrations and adipose mass.

Circulating lipids were markedly improved in VSG-operated animals compared with both S-AL and S-WM animals. Fasting plasma TG concentrations were 50% lower in VSG-operated animals compared with S-AL and S-WM up to 5 months after surgery (P < 0.001) (Fig. 3C). Furthermore, fasting plasma cholesterol concentrations were lower in VSG-operated animals compared with S-AL and S-WM up to 5 months after surgery (P < 0.001) (Fig. 3D).

Similar to previous studies in both rodents and humans, fasting plasma ghrelin concentrations were reduced by approximately 75% in VSG-operated animals compared with both S-AL and S-WM (P < 0.001) (Fig. 3E). Fasting plasma adiponectin concentrations were elevated in S-WM and VSG-operated animals compared with S-AL (P < 0.001) (Fig. 3F). However, fasting plasma adiponectin concentrations were further elevated in VSG-operated animals compared with S-WM, demonstrating a weight-independent increase of circulating adiponectin concentrations after VSG surgery (P < 0.05) (Fig. 3F).

A previous study reported that incubation of brown adipocytes with ghrelin impairs adiponectin production (32). Thus, we hypothesized that decreases of circulating ghrelin in VSG-operated animals may have contributed to the body weight-independent augmentation of circulating lipids.

| TABLE 1. Age of diabetes onset and incidence in animals followed to 1 yr of age |
|-----------------|-----------------|-----------------|
| Age of onset (d) | S-AL             | S-WM             | VSG             |
| Incidence (%)    | 182 ± 16         | 210 ± 6          | 288 ± 23^a      |
| Diabetes-free days | 120 ± 6          | 19 ± 6           | 23              |
| n                | 14               | 16               | 13              |

Values are mean ± SEM.

^a P < 0.05 compared with S-AL and S-WM.

^b P < 0.001 compared with S-AL by Student’s t test.
adiponectin concentrations. To test this hypothesis, we exposed differentiated 3T3-L1 adipocytes to concentrations of acylated ghrelin calculated from the total ghrelin concentrations (28) measured in the VSG-operated animals (AG low) and from the ghrelin concentrations measured in the S-WM animals (AG medium) and a dose 2-fold higher than the medium dose of ghrelin (AG high). Adiponectin secretion and mRNA expression were not

**FIG. 1.** Kaplan-Meier analysis of diabetes incidence in S-AL (n = 14), S-WM (n = 16), and VSG-operated animals up to 8 months of age (n = 13) (A). *, P < 0.05 compared with S-AL and S-WM by log-rank test. Fasting serum glucose concentrations (B). +++, P < 0.001 by two-factor repeated measures ANOVA. ***, P < 0.001; **, P < 0.01 compared with S-AL by Student’s t test.

**FIG. 2.** Energy intake (A), body weight (B), energy expenditure (C), and AUC of energy expenditure (D) in S-AL (n = 14), S-WM (n = 16), and VSG-operated animals (n = 13). +++, P < 0.001 for S-WM and VSG compared with S-AL by two-factor repeated measures ANOVA with Bonferroni’s post hoc test compared with S-AL.
affected by exposure to AG concentrations similar to the concentrations found in VSG-operated animals. However, adiponectin secretion and mRNA expression were reduced in a step-wise manner with exposure to AG concentrations similar to the concentrations found in S-WM animals as well as by 2-fold higher AG concentrations. Incubation of differentiated 3T3-L1 adipocytes with similar concentrations of inactive DAG did not affect adiponectin secretion or adiponectin mRNA expression (Fig. 4).

**VSG surgery increases circulating bile acid concentrations**

Circulating bile acid profiles did not differ between groups at baseline (Fig. 5A). At 3 months after surgery, there was a 2-fold increase of circulating deoxycholic acid concentrations ($P < 0.05$), but total circulating bile acid concentrations did not differ between groups (Fig. 5B). However, at 5 months after surgery, total bile acid concentrations and all bile acid subtype concentrations were elevated in VSG-operated animals compared with S-AL and S-WM ($P < 0.05$) (Fig. 5C). The elevation of circulating total bile acid concentrations in VSG-operated animals was primarily due to an increase of conjugated bile acids as both an absolute value and as a percentage of the total bile acid pool ($P < 0.05$) (Fig. 6, A and B). Specifically, taurine-conjugated bile acids were elevated approximately 3-fold in VSG-operated animals compared with...
S-AL and S-WM ($P < 0.05$) (Fig. 6, A and B). Glycine-conjugated bile acids did not differ between groups as an absolute value (Fig. 6C), but glycine-conjugated bile acids as a proportion of total conjugated bile acids were lower in VSG-operated animals compared with S-AL and S-WM ($P < 0.05$) (Fig. 6D). Fasting plasma-unconjugated bile acids did not differ between groups when expressed as an absolute value (Fig. 6E). However, unconjugated bile acids were proportionally lower in VSG-operated animals compared with S-AL and S-WM ($P < 0.05$) (Fig. 6F). Thus, VSG surgery results in a preferential increase of circulating taurine-conjugated bile acid concentrations.

**VSG surgery improves insulin secretion and increases nutrient-stimulated GLP-1, GLP-2, and PYY secretion**

OGTT were performed at 1 and 4 months after surgery to assess changes of glucose tolerance and nutrient-stimulated gastrointestinal hormone secretion over time. Glucose excursions were significantly lower in S-WM and VSG-operated animals at both 1 and 4 months after surgery ($P < 0.01$) (Fig. 7, A and B). VSG resulted in significantly lower insulin excursions, as indexed by area under the curve (AUC), compared with S-WM at both 1 and 4 months after surgery, suggesting an improvement of insulin sensitivity ($P < 0.05$) (Fig. 7, C and D). At 4 months after surgery, insulin excursions were markedly blunted in S-AL animals due to the greater prevalence of diabetes in this group and the concomitant impairment of $\beta$-cell function (Fig. 7D). Importantly, the percent increase of plasma insulin concentrations from fasting to peak values was significantly higher in VSG-operated animals compared with both S-AL and S-WM during both the 1 and 4 month OGTT, demonstrating superior islet function in VSG-operated animals ($P < 0.01$) (Fig. 7, C and D).
Peak plasma GLP-1 concentrations were 3-fold higher in VSG-operated animals compared with S-AL and S-WM animals at both 1 and 4 months after surgery \((P < 0.001)\) (Fig. 7, E and F). Furthermore, GLP-1 excursions were significantly higher at 4 months after surgery compared with at 1 month after surgery in VSG-operated animals, demonstrating preservation of postoperative increases of postprandial GLP-1 secretion \((P < 0.05)\). The GLP-1 AUC was significantly lower in S-WM compared with S-AL animals during both the 1 and 4 month OGTT \((P < 0.05)\). Similarly, nutrient-stimulated GLP-2 secretion was significantly elevated in VSG-operated animals compared with S-AL and S-WM \((P < 0.05)\) (Fig. 7, G and H). This effect was preserved up to 4 months after surgery.

PYY excursions were markedly elevated in VSG-operated animals compared with S-AL and S-WM animals at both 1 and 4 months after surgery \((P < 0.001)\) (Fig. 7, I and J). Although the PYY AUC in VSG-operated animals tended to be higher at 4 months after surgery compared

![Graphs showing plasma bile acid concentrations](image-url)
with 1 month after surgery, this difference did not reach statistical significance. However, these data also demonstrate the durability of postoperative increases of postprandial PYY secretion in VSG-operated animals.

**Discussion**

In this study, we demonstrate that VSG performed in prediabetic male UCD-T2DM rats delays the onset of type 2 diabetes and results in long-lasting improvements of glucose and lipid metabolism. Postoperative decreases of food intake and body weight contributed to the delay in diabetes onset, as there was approximately a 5-month increase in the number of diabetes-free months in S-WM and VSG-operated animals compared with S-AL. However, metabolic and endocrine changes after VSG surgery likely contributed to the delay in diabetes onset, as VSG surgery resulted in a significantly higher age of diabetes onset and a significant delay in diabetes onset up to 8 months of age by Kaplan-Meier analysis compared with weight-matched controls. The weight-independent metabolic and endo-
crine changes that we observed after VSG surgery that likely contributed to the delay in diabetes onset include increased postprandial GLP-1 secretion, increased circulating bile acid and adiponectin concentrations, and decreased circulating ghrelin concentrations.

Glucose-stimulated GLP-1 and GLP-2 secretion were elevated at both 1 and 4 months after surgery in VSG-operated animals. Postprandial GLP-1 secretion in VSG-operated animals was further elevated at 4 months after surgery compared with 1 month after surgery, demonstrating that this response does not taper over time. The “hind-gut” hypothesis postulates that increases of postprandial GLP-1 secretion after bariatric surgery are due to increased delivery of incompletely absorbed nutrients to the distal small intestine, resulting in direct stimulation of L cells to release GLP-1. Although VSG surgery increases gastric emptying, the ability of VSG surgery to increase the flux of incompletely absorbed nutrients in the distal small intestine is questionable, because it does not involve bypass of the proximal small intestine. Furthermore, human clinical studies have demonstrated similar postprandial GLP-1 secretion profiles after VSG compared with RYGB. This suggests that mechanisms other than that proposed in the hind-gut hypothesis may contribute to postoperative increases of postprandial GLP-1 secretion. One potential contributor is activation of a vagal reflex by nutrients in the proximal small intestine.

Increases of GLP-1 secretion likely contributed to the improvement of insulin sensitivity and islet function. Improvement of glucose-stimulated insulin secretion was observed during the OGTT at 1 and 4 months after surgery, in which VSG-operated animals exhibited approximately 2-fold greater glucose-stimulated insulin secretion compared with both S-AL and S-WM animals. As an incretin hormone, GLP-1 potentiates glucose-stimulated insulin secretion. Furthermore, GLP-1 has been shown to have beneficial effects on the islet by increasing insulin synthesis, stimulating β-cell proliferation, and preventing β-cell apoptosis. Increases of GLP-2 secretion may have contributed to the increases of nutrient-stimulated GLP-1 secretion, because GLP-2 is known to stimulate proliferation of intestinal cells and thus may contribute to an increase in L-cell number. Nutrient-stimulated GLP-2 secretion has been shown to be elevated after ileal interposition (IT) surgery in rats and early after RYGB and VSG in humans. However, this study demonstrates that the increased GLP-2 response is maintained over time.

The decreases of fasting circulating insulin concentrations and decreased insulin AUC during both the 1 and 4 month OGTT compared with weight-matched animals suggest an improvement of insulin sensitivity that is independent of body weight. Increases of nutrient-stimulated GLP-1 secretion may have improved insulin sensitivity by reducing glucotoxicity and lipotoxicity. GLP-1 reduces glucotoxicity by improving islet function and reducing hepatic gluconeogenesis. GLP-1 may contribute to reductions in lipotoxicity by stimulating fat oxidation. VSG-operated animals exhibited improvements of lipid metabolism with markedly lower fasting plasma lipids.

Bile acids have been shown to be elevated after RYGB and IT surgery and may represent another mechanism involved in the metabolic improvements after bariatric surgery. Here, we found marked increases of total circulating bile acid concentrations at 5 months after surgery. These increases were similar in magnitude to those reported in obese rats after IT surgery. However, bile acid profiles were different in VSG-operated animals compared with what has been previously reported after IT surgery. The increase of circulating bile acid concentrations after VSG surgery was primarily due to increased taurine-conjugated bile acids with proportional decreases of unconjugated bile acid concentrations. In contrast, Kohli et al. report marked preferential increases of unconjugated bile acids along with increases of glycine and taurine-conjugated bile acid concentrations. These differences in circulating bile acid profiles between IT and VSG surgery suggest that the mechanisms responsible for postoperative increases of circulating bile acid concentrations likely differ based on the type of bariatric surgery performed.

The increases of circulating bile acid concentrations after VSG surgery may have contributed to the observed improvements of insulin sensitivity and circulating lipids. Bile acids have been shown to decrease hepatic gluconeogenesis and lipogenesis and increase insulin-mediated glucose disposal in adipocytes by signaling through farnesoid X receptor. Furthermore, administration of taurine-conjugated bile acids to rats has been shown to improve insulin signaling and decrease the expression of gluconeogenic genes, likely by signaling through farnesoid X receptor. Bile acids have also been shown to signal through TGR5 receptors located in the distal small intestine to increase GLP-1 secretion and through TGR5 receptors on skeletal muscle and brown adipose tissue to increase energy expenditure. Energy expenditure did not differ between VSG-operated animals and controls. However, energy expenditure was measured at 1.5 months after surgery, before a significant increase of circulating bile acid concentrations was observed.

Similar to previous studies of RYGB and VSG, circulating adiponectin concentrations were significantly elevated after VSG surgery compared with ad libitum-fed
controls (45). However, to our knowledge, this is the first study to demonstrate postoperative increases of circulating adiponectin concentrations compared with a weight-matched control, indicating that postoperative increases of circulating adiponectin are likely due to additional mechanisms beyond the reduction of body weight. We hypothesized that decreases of circulating ghrelin concentrations after VSG surgery contributed to the body weight-independent effects of VSG surgery to increase circulating adiponectin concentrations. This is supported by our in vitro results demonstrating significant inhibition of both adiponectin mRNA expression and secretion in 3T3-L1 adipocytes exposed to active acylated ghrelin concentrations similar to those of sham-operated animals. Increases of circulating adiponectin likely contributed to the improvement of insulin sensitivity and circulating lipids. Adiponectin is an adipokine that has been demonstrated to improve insulin sensitivity and lipid metabolism (46). Furthermore, circulating adiponectin concentrations are reduced in obesity and type 2 diabetes (47). Adiponectin has been proposed to improve insulin sensitivity by signaling in liver and skeletal muscle through activation of AMP kinase to promote decreased gluconeogenesis, increased insulin-mediated glucose uptake, and increased fatty acid oxidation resulting in decreased ectopic lipid deposition (46).

Similar to previous studies in rodents and humans, we demonstrated a marked decrease of circulating ghrelin concentrations after VSG surgery (10, 48). Decreased circulating ghrelin concentrations likely contributed to reduced food intake, improved glucose-stimulated insulin secretion, and improved insulin sensitivity. Ghrelin has been shown to promote food intake and increase adiposity (49). Reductions of circulating ghrelin likely contributed to improved islet function, because previous studies have demonstrated that both physiologic and pharmacologic doses of exogenous ghrelin impair glucose-stimulated insulin secretion in cell culture studies and in humans (15, 17). This has been suggested to be mediated by activation of voltage-dependent K+ channels resulting in decreased calcium-mediated insulin release (15). Ghrelin has also been shown to promote insulin resistance, because ablation of ghrelin in both the ob/ob mouse and high-fat-fed mice has been shown to improve insulin sensitivity (16). Furthermore, inhibition of ghrelin activation by ghrelin o-acyltransferase using a ghrelin o-acyltransferase antagonist results in improvement of glucose tolerance and body weight, making ghrelin an attractive pharmaceutical target for the treatment of type 2 diabetes (50).

In conclusion, we have shown for the first time that VSG surgery delays diabetes onset in the UCD-T2DM rat model of type 2 diabetes, an effect that is partially independent of reduced body weight. Furthermore, VSG surgery results in marked improvements of glucose and lipid metabolism with improved islet function and decreased circulating lipids. Increases of circulating adiponectin, bile acids, and GLP-1 concentrations and decreases of circulating ghrelin all likely contributed to the effect of VSG surgery to delay type 2 diabetes onset. Further studies on the effect of VSG surgery and other bariatric surgeries to delay type 2 diabetes onset in the UCD-T2DM rat will help to identify new pharmaceutical targets for the treatment and prevention of type 2 diabetes.

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