Dipeptidyl Peptidase-4 Inhibition Potentiates Stimulated Growth Hormone Secretion and Vasodilation in Women

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Background—Diminished growth hormone (GH) is associated with impaired endothelial function and fibrinolysis. GH-releasing hormone is the primary stimulus for GH secretion and a substrate of dipeptidyl peptidase-4. We tested the hypothesis that dipeptidyl peptidase-4 inhibition with sitagliptin increases stimulated GH secretion, vasodilation, and tissue plasminogen activator (tPA) activity.

Methods and Results—Healthy adults participated in a 2-part double-blind, randomized, placebo-controlled, crossover study. First, 39 patients (29 women) received sitagliptin or placebo on each of 2 days separated by a washout. One hour after study drug, blood was sampled and then arginine (30 g IV) was given to stimulate GH. Vasodilation was assessed by plethysmography and blood sampled for 150 minutes. Following a washout, 19 of the original 29 women received sitagliptin alone versus sitagliptin plus antagonist to delineate GH receptor (GHR)– (n=5), nitric oxide– (n=7), or glucagon-like peptide-1 receptor– (n=7) dependent effects. Sitagliptin enhanced stimulated GH secretion (P<0.01 versus placebo, for 30 minutes) and free insulin–like growth factor-1 (P<0.001 versus placebo, after adjustment for baseline) in women. Vasodilation and tPA increased in all patients, but sitagliptin enhanced vasodilation (P=0.01 versus placebo) and increased tPA (P<0.001) in women only. GHR blockade decreased free insulin–like growth factor-1 (P=0.04 versus sitagliptin alone) and increased stimulated GH (P<0.01), but decreased vascular resistance (P=0.01) such that nadir vascular resistance correlated inversely with GH (r = −0.90, P<0.001). GHR blockade suppressed tPA. Neither nitric oxide nor glucagon-like peptide-1 receptor blockade affected vasodilation or tPA.

Conclusions—Sitagliptin enhances stimulated GH, vasodilation, and fibrinolysis in women. During sitagliptin, increases in free insulin–like growth factor-1 and tPA occur via the GHR, whereas vasodilation correlates with GH but occurs through a GHR-independent mechanism.

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The growth hormone (GH) insulin-like growth factor-1 (IGF-1) axis regulates vascular function and fibrinolytic capacity in adults. GH is secreted in a pulsatile fashion from the pituitary gland and acts directly or indirectly through IGF-1, which is secreted by the liver in response to hepatic GH receptor (GHR) activation. GH and IGF-1 receptors are ubiquitously present on human endothelial cells and the myocardium. In healthy patients, acute systemic and intraarterial GH infusion increases GH without affecting IGF-1, and increases endothelium-dependent vasodilator function. Patients with GH deficiency demonstrate altered fibrinolytic balance, characterized by elevated plasminogen activator inhibitor-1 (PAI-1) antigen levels and lower tissue plasminogen activator (tPA) activity, which resolves with GH replacement therapy. IGF-1 also exerts many favorable endothelial effects via the IGF-1 receptor, including enhanced...
endothelium-dependent vasodilation, anti-inflammatory effects, and maintenance of vascular integrity via activated endothelial progenitor cells.2,7 Thus, the vascular effects of GH may be mediated by either direct effects of GH or indirect effects of increased IGF-1.

Adults with low GH or IGF-1 levels have elevated cardiovascular risk. Patients who have attenuated GH secretion include those with abdominal obesity, physiologic aging, HIV infection, polycystic ovarian syndrome, and pituitary and hypothalamic disease.8–10 In cardiovascular epidemiology studies, individuals in the lowest IGF-1 quartile have a higher risk of ischemic heart disease,11 heart failure,12 and all-cause mortality.13 Similarly, overweight and obese adults demonstrate a decreased response to GH secretagogues, which correlates with the presence of cardiovascular risk markers.14 For this reason, pharmacologic interventions designed to increase GH and IGF-1 in deficient populations have been developed in the past 2 decades in hopes of mitigating cardiovascular risk. Recombinant GH therapy was the first such therapy approved to increase GH and IGF-1 levels in deficient patients; however, exogenous GH is not restrained by physiologic negative feedback by IGF-1, does not restore pulsatile secretion, and is limited by the side effect of hyperglycemia.

An alternative and previously unexplored method to enhance GH and downstream IGF-1 secretion in humans is to inhibit the degradation of endogenous GH-releasing hormone (GHRH) by dipeptidyl peptidase-4 (DPP4). GHRH is the primary stimulus for pituitary GH secretion and determines GH pulsatility. Endogenous GHRH has a half-life of ≈6 minutes in humans as it is degraded and inactivated by DPP4.15 Sitagliptin was the first DPP4 inhibitor approved by the US Food and Drug Administration in 2006 for the management of hyperglycemia in patients with type 2 diabetes mellitus. Sitagliptin decreases the degradation of the incretin hormone, glucagon-like peptide-1 (GLP-1), and thereby improves postprandial hyperglycemia in patients with diabetes mellitus in a glucose-dependent manner.

In this study, we tested the hypothesis that inhibition of DPP4 activity by sitagliptin would enhance stimulated GH secretion and thereby increase vasodilation and plasma tPA activity in healthy young men and women. To further elucidate the mechanism by which stimulating the GH–IGF-1 axis influences vascular function and fibrinolysis, we then investigated the contribution of nitric oxide, GHR activation, and GLP-1 receptor activation to the vascular changes observed with stimulated GH secretion during DPP4 inhibition.

### Methods

Anonymized data and analytic methods will be made publicly available (NCT01701973 at www.clinicaltrials.gov).

### Study Protocol

Healthy, lean (body mass index ≤25 kg/m²), nonsmoking adults, aged 18 to 40 years, participated in a 2-part double-blind, randomized, placebo-controlled, crossover study. (See Table 1 for patient characteristics.) The study adhered to the principles of the Declaration of Helsinki and Title 45, US Code

| Parameter         | N=39     |
|-------------------|---------|
| Age, y            | 25±5    |
| Race, No. (%)     |         |
| White             | 31 (80) |
| Black             | 2 (5)   |
| Hispanic          | 4 (10)  |
| Asian             | 2 (5)   |
| Sex, No. (%)      |         |
| Women             | 29 (74) |
| Men               | 10 (26) |
| Weight, kg        | 65.4±9.3|
| Body mass index, kg/m² | 22.9±1.8 |

Values are expressed as mean±SD unless otherwise indicated.
of Federal Regulations, Part 46, Protection of Human Subjects, and was approved by the Vanderbilt University Medical Center’s institutional review board. All patients provided written informed consent before initiation of study procedures. Patients with a history of chronic illness, including diabetes mellitus, hypertension, cardiovascular disease, and chronic renal or hepatic insufficiency, were excluded. Medication use other than a multivitamin was prohibited at the time of study; oral contraceptive use was not permitted in women. Pregnancy was excluded in women of child-bearing age by serum pregnancy testing.

In the first half of the study (Figure S1A), 39 patients underwent 2 study days separated by a washout period to determine the effect of DPP4 inhibition on stimulated GH secretion. Patients were assigned to treatment order (sitagliptin or matching placebo) using a block randomization algorithm. On each study day, patients reported to the Vanderbilt Clinical Research Center in the morning after an overnight fast. All patients were studied in the supine position in a temperature-controlled room. Participants were given oral study drug (sitagliptin or matching placebo), and a peripheral intravenous line was placed in the antecubital fossa of their nondominant arm. Sitagliptin dose was 200 mg in the first 14 patients (7 women) to achieve >80% inhibition of DPP4 activity within 1 hour. We changed to sitagliptin 100 mg daily for 4 days in the remaining 25 patients to achieve steady-state dosing and to be consistent with the current Food and Drug Administration–approved dose. Study days were separated by a 1-week washout for the first 14 participants and a 2-week washout in the remaining 25 participants. Vasodilation was assessed via strain-gauge plethysmography (Data S1) and venous blood samples were obtained 60 minutes following study drug. Patients then received arginine (30 g) intravenously over 30 minutes to stimulate endogenous GH secretion in a GHRH-dependent manner, as previously described.16 Vasodilation was assessed and venous blood samples were obtained 150 minutes after completion of arginine infusion. On the second study day, the protocol was repeated using the opposite study drug (sitagliptin or matching placebo). Blood pressure and heart rate were monitored throughout each study day.

Following at least an 8-week washout period, 19 of the 29 women from the first half of the study participated in 2 additional study visits designed to elucidate the mechanism by which sitagliptin potentiated vasodilation and tPA release (Figure S1B). We studied women only based on the results of the first part of the study. Each patient’s sitagliptin dose and washout period was the same as in the first half of the protocol. Patients were divided into 3 subgroups. In the first subgroup, 5 women were randomized to a single double-blinded subcutaneous injection of saline vehicle or pegvisomant 80 mg (Pfizer Inc) administered 72 hours before the study day to block the GHR. This dose of pegvisomant produces peak drug levels 72 hours following administration with a reduction in free IGF-1 indicative of efficient GHR blockade.17 In the second subgroup, 7 women were randomized to either double-blinded saline vehicle infusion or the nitric oxide synthase inhibitor, L-N-monomethylarginine (LNMMA [acetate]; Clinalfa; Bachem Americas, Inc), administered as a 3-mg/kg 15-minute intravenous infusion preceding arginine followed by an additional 6 mg/kg infused over 120 minutes. This dose demonstrates peak hemodynamic effects 20 minutes after the start of the infusion and does not affect arginine-stimulated GH secretion.18,19 In the third subgroup, 7 women were randomized to either double-blinded saline vehicle infusion or Exendin 9-39 (Exendin 9-39 Acetate; Clinalfa; Bachem Americas, Inc), administered as an intravenous bolus infusion of 7500 pmol/kg over 1 minute preceding arginine followed by a continuous infusion of 750 pmol/kg per minute for 150 minutes, to block the GLP-1 receptor.20

Laboratory Analyses

All samples were obtained after the first 3 mL of blood were discarded. Blood samples were collected on ice, centrifuged immediately, and plasma-stored at −80°C in prespecified aliquots until time of assay. Venous DPP4 activity was determined by ELISA (eBioscience). Venous DPP4 activity was assayed by incubating 20 µL of serum sample in 80 µL assay buffer (0.1 mol/L Tris at a pH of 8.0; Bachem) for 30 minutes at 37°C with colorimetric substrate (2 mmol/L L-glycyl-L-prolyl p-nitroanilide hydrochloride [Sigma Aldrich]) for a total reaction volume of 200 µL, as previously described.21 The enzyme activity was assessed by measuring the increase in specific absorbance at 405 nm at 0, 15, and 30 minutes and was expressed as nmol/mL per minute. GH levels were determined using the Access Ultralitative hGH Assay (Beckman Coulter), while GH levels after pegvisomant administration were analyzed using the IDS-iSYS hGH assay; both assays were calibrated against National Institute for Biological Standards and Control World Health Organization International Standard 98/574. Free IGF-1 was determined using a commercially available ELISA (R&D systems). IGF-1 was analyzed by Luminex assay (EMD Millipore), which is calibrated against the National Institute for Biological Standards and Control World Health Organization International Standard 02/254. tPA activity and PAI-1 antigen levels were measured in blood collected in acidified citrate anticoagulant (TrinILIZE Stabilyte tubes, Tcoag; Bray Co). tPA activity was analyzed using a biofunctional immunoassay calibrated against National Institute for Biological Standards and Control World Health Organization International Standard 86/670 (TriniLIZE tPA Activity, Tcoag, Co) in the first 14 patients.
Following discontinuation of this assay, the remaining samples were analyzed using an ELISA calibrated against National Institute for Biological Standards and Control World Health Organization International Standard 98/714 (Oxford Biomedical Research). Samples from each patient were assayed using the same method. PAI-1 antigen was analyzed using a TintElize PAI-1 antigen assay (Toag, Co). Plasma cGMP was determined using a competitive enzyme immunoassay (GE Healthcare Bio-Sciences Corp). Samples for analysis of active GLP-1 were collected in aprotinin and analyzed using the MILLPLEX MAP Human Metabolic Hormone Magnetic Bead Panel (EMD Millipore Corporation). Insulin and estradiol were analyzed by double-antibody radioimmunoassay. Bedside blood glucose was determined by YSI bedside glucose analyzer (YSI Life Sciences).

Statistical Analysis

Data are presented as mean±SD, unless otherwise noted. We tested for carryover effect using the t test approach proposed by Jones and Kenward. Wilcoxon signed-rank test was used to compare baseline variables between treatment conditions as well as GH levels (untransformed) between treatment conditions at each time point. Wilcoxon rank-sum test was used to compare percent DPP4 inhibition before GH stimulation, peak GH during placebo, and peak GH during sitagliptin between men and women. Percent DPP4 inhibition was determined by the equation: \[1 - \text{DPP4 activity during sitagliptin} / \text{DPP4 activity during placebo}\] × 100. Spearman correlation was used to evaluate the association between continuous variables. Mixed effect models were used to analyze the data with a random subject effect and with interaction terms removed from the final model when the \(P\) value from the corresponding overall test for interaction was >0.2. Results from mixed effect models are presented as the mean difference between treatments with 95% confidence interval. The end points GLP-1, insulin, and GH were log transformed to satisfy model assumptions. Statistical analyses were performed using IBM SPSS software version 23.0, GraphPad Prism 5 and R 2.15.0 (www.r-project.org). Sample size calculations are included in Data S1.

Results

Effect of Sitagliptin on DPP4 Activity and GLP-1

Sitagliptin significantly decreased DPP4 activity (\(P<0.001\) versus placebo) and increased GLP-1 levels both at baseline and throughout stimulated GH secretion (\(P<0.0001\) versus placebo) (Table 2). Sitagliptin did not affect insulin levels (\(P=0.45\) versus placebo) or blood glucose levels (\(P=0.58\) versus placebo) during stimulated GH secretion or at baseline (Table 2). Sitagliptin (200 mg) reduced DPP4 activity similarly in men and women (percent DPP4 inhibition 78±13% in 7 women versus 80±6% in 7 men, \(P>0.999\)). Safety data are summarized in Data S1.

Effect of Sitagliptin on Stimulated GH Secretion and Free IGF-1

Arginine infusion stimulated GH secretion to a greater extent in women (\(n=29\)) than in men (\(n=10\)), as previously

Table 2. Initial Biochemical Parameters Before GH Stimulation

| Variable                        | Placebo                          | Sitagliptin                     | No.* | \(P\) Value |
|---------------------------------|----------------------------------|---------------------------------|------|------------|
| DPP4 activity, nmol/mL per minute | 25.4±6.5 (24.8, 9.7)             | 8.1±4.5 (7.0, 5.6)              | 39   | <0.001     |
| DPP4 antigen, ng/mL             | 458.1±158.1 (458.4, 191.7)       | 431.2±149.4 (423.8, 250.7)      | 38   | 0.22       |
| Blood glucose, mg/dL            | 86.6±5.8 (86.0, 8.4)             | 85.4±7.8 (85.1, 7.9)            | 39   | 0.24       |
| Insulin, µU/mL                  | 6.8±3.2 (6.6, 4.1)               | 7.3±4.0 (7.0, 6.0)              | 39   | 0.30       |
| GLP-1, pg/mL                    | 2.7±2.8 (2.0, 0.8)               | 19.1±12.5 (19.1, 16.6)          | 39   | <0.001     |
| Estradiol, pg/mL                | 252.9±256.6 (177.6, 145.7)       | 250.9±173.9 (238.4, 195.8)      | 29   | 0.67       |
| tPA activity, IU/mL             | 0.19±0.18 (0.14, 0.18)           | 0.24±0.26 (0.14, 0.31)          | 36   | 0.13       |
| PAI-1 antigen, ng/mL            | 4.4±4.1 (3.0, 4.6)               | 3.5±2.7 (2.9, 4.0)              | 36   | 0.26       |
| Total IGF-1, ng/mL              | 108.3±30.4 (102.3, 46.6)         | 104.6±30.5 (106.8, 38.6)        | 39   | 0.11       |
| Free IGF-1, ng/mL               | 0.65±0.31 (0.61, 0.45)           | 0.65±0.24 (0.66, 0.39)          | 37   | 0.67       |

Results are presented as mean±SD (median, interquartile range). DPP4 indicates dipeptidyl peptidase-4; GLP-1, glucagon-like peptide-1; IGF-1, insulin-like growth factor-1; PAI-1, plasminogen activator inhibitor-1; tPA, tissue plasminogen activator.

*Results were analyzed in patients with data available on both days.

†Analyzed in women only.
DPP4 Inhibition Increases Growth Hormone and Vasodilation

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Effect of Sitagliptin on Vasodilation and cGMP Levels During Stimulated GH Secretion

DPP4 inhibition did not significantly affect blood pressure, pulse rate, or vasodilation before arginine infusion, as compared with placebo (Table 3). Vasodilator response is presented as the change in both forearm blood flow (FBF) and forearm vascular resistance (FVR) following arginine infusion. FBF increased (P<0.001 effect of time) and FVR decreased (P<0.001 effect of time) following stimulated GH secretion. Sitagliptin enhanced the increase in FBF (P=0.01 versus placebo) and decrease in FVR (P=0.003 versus placebo) in women only (Figure 2). Sitagliptin also increased pulse rate (P=0.03 versus placebo) following stimulated GH secretion in women. Sitagliptin increased cGMP levels during stimulated GH secretion in both women (increase of 98.57 fmol [95% confidence interval, 33.18–163.96], P=0.003 versus placebo) and men (increase of 116.39 fmol [95% confidence interval, 43.33–189.33], P=0.002 versus placebo).

Effect of Sitagliptin on tPA Activity During Stimulated GH Secretion

tPA activity increased following stimulated GH secretion (P<0.001). Acute inhibition of DPP4 activity with 200 mg of sitagliptin increased tPA activity levels in women (P<0.001 versus placebo, n=7) but decreased tPA activity in men (P=0.02 versus placebo, n=7) (Figure 3). This effect of DPP4 inhibition on tPA activity levels was not observed following 100 mg of sitagliptin. PAI-1 antigen levels during stimulated GH secretion were unaffected by sitagliptin in men and women (P=0.33 versus placebo).

Effect of GHR Blockade on Vasodilation and tPA Activity During Stimulated GH Secretion in Women

Pegvisomant significantly decreased free IGF-1 during sitagliptin (P=0.04 versus sitagliptin alone, n=5) and increased GH levels (P<0.01 versus sitagliptin alone), consistent with effective GHR blockade (Figure 4A). The addition of GHR blockade significantly increased vasodilation (P<0.01 versus sitagliptin alone for change in FVR) throughout stimulated GH secretion. Moreover, at the nadir in vascular resistance, GH levels correlated inversely with vascular resistance (r=−0.90, P<0.001) (Figure 4B). Pegvisomant suppressed tPA activity before (0.24±0.12 after sitagliptin alone versus 0.10±0.08 IU/mL after addition of pegvisomant, P=0.04) and during stimulated GH secretion (P<0.001 versus sitagliptin alone) (Figure 4C).

Effect of NO Synthase Inhibition on Vasodilation and tPA Activity During Stimulated GH Secretion in Women

LNMMA significantly decreased cGMP levels during sitagliptin and stimulated GH secretion (decrease of 432.39 fmol [95% confidence interval, −792.77 to −72.02], P=0.02 versus sitagliptin alone, n=7). LNMMA did not affect the vasodilator response to stimulated GH secretion during sitagliptin (P=0.43 versus sitagliptin alone for change in FVR and P=0.94 versus sitagliptin alone for change in FVR) (Figure 5A). The addition of LNMMA to sitagliptin also had no effect on tPA activity (P=0.64 versus sitagliptin alone) (data not shown).

Effect of GLP-1 Receptor Blockade on Vasodilation and tPA Activity During Stimulated GH Secretion in Women

GLP-1 receptor blockade with Exendin 9-39 increased fasting GLP-1 (P<0.01), glucagon (P<0.09), and blood glucose levels (P<0.001), as previously described. Exendin 9-39 briefly caused vasoconstriction immediately after arginine infusion (P=0.02 versus sitagliptin alone for FBF and P=0.02 versus sitagliptin alone for FVR at 60 minutes, n=7) (Figure 5B). Following stimulated GH secretion, FBF increased (P<0.001 effect of time) and FVR decreased (P<0.001 effect of time). The addition of Exendin 9-39 to sitagliptin did not prevent vasodilation following stimulated GH secretion (P=0.88 versus sitagliptin alone for change in FBF and P=0.57 versus sitagliptin alone for change in FVR). The
Figure 1. Dipeptidyl peptidase-4 (DPP4) inhibition with sitagliptin enhances early stimulated growth hormone (GH) secretion (A) and free insulin–like growth factor-1 (IGF-1) levels (B) and shortens the time to peak GH (C) in women (n=29) but not men (n=10). Change in free IGF-1 levels from baseline to 90 minutes following arginine. (Free IGF-1 data available in 28 women and 9 men.) Data are presented as mean±SEM. *P≤0.05; †P<0.10 vs placebo at same time point by Wilcoxon signed-rank test. The effect of treatment on free IGF-1, in the linear model, was significant in women (P<0.001 vs placebo, after adjustment for baseline free IGF-1). The effect of treatment on ln(GH), in the linear model, was significant in women for 30 minutes following arginine (P<0.01 vs placebo, after adjustment for baseline GH).
addition of Exendin 9-39 to sitagliptin also had no effect on tPA activity ($P=0.58$ versus sitagliptin alone) (data not shown).

Reproducibility of Stimulated GH Secretion During DPP4 Inhibition

The reproducibility of the effect of DPP4 inhibition on stimulated GH secretion was assessed by comparing GH levels during sitagliptin alone with GH levels obtained during sitagliptin plus saline vehicle infusion in the 19 women who completed both crossover studies (Figure 6). There was a significant correlation between stimulated GH secretion following sitagliptin and stimulated GH secretion following sitagliptin plus saline infusion (peak GH response: $r_s=0.65$, $P=0.003$; GH 30 minutes after arginine: $r_s=0.51$, $P=0.02$).

Discussion

This study tested the hypothesis that DPP4 inhibition potentiates arginine-stimulated GH secretion in humans. We found that sitagliptin significantly enhanced stimulated GH secretion and shortened the time to peak GH in healthy

Table 3. Initial Hemodynamic Parameters Before GH Stimulation

| Variable                  | Placebo | Sitagliptin | $P$ Value |
|---------------------------|---------|-------------|-----------|
| Systolic BP, mm Hg        | 108.8±10.8 (110.0, 18.0) | 109.9±10.7 (108.0, 12.0) | 0.40 |
| Diastolic BP, mm Hg       | 65.6±6.4 (65.0, 9.0) | 65.6±6.5 (66.0, 6.0) | 0.97 |
| Mean arterial pressure, mm Hg | 82.9±6.2 (84.0, 10.0) | 83.2±5.9 (83.0, 6.0) | 0.59 |
| Pulse rate, beats per min | 58.3±8.3 (57.0, 13.0) | 59.3±8.1 (60.0, 11.0) | 0.29 |
| FVR, mm Hg/(mL/min/100mL) | 36.9±11.4 (36.2, 19.4) | 37.4±10.2 (35.6, 12.3) | 0.58 |
| FBF, mL/min per 100 mL    | 2.5±0.9 (2.2, 1.3) | 2.4±0.6 (2.4, 0.8) | 0.65 |

BP indicates blood pressure; FBF, forearm blood flow; FVR, forearm vascular resistance; GH, growth hormone. Results are presented as mean±SD (median, interquartile range).

Figure 2. Dipeptidyl peptidase-4 inhibition with sitagliptin enhances vasodilation during arginine (Arg) stimulated growth hormone secretion in women ($n=29$) but not men ($n=10$). The overall effect of treatment, as determined by linear model, was significant in women ($P=0.013$ effect of treatment on percent change [Δ] in forearm blood flow and $P=0.003$ effect of treatment on percent change in forearm vascular resistance). Data are presented as mean±SEM. *$P<0.05$ vs placebo at specified time point in the linear model.
women but not men. Similarly, sitagliptin increased free IGF-1 levels in women. Forearm vasodilation after peak GH was potentiated by sitagliptin only in women. GHR blockade further increased vasodilation during DPP4 inhibition in association with increased GH levels. The latter indicates that GH induces endothelium-independent vasodilation through a GHR-independent mechanism.

Our study is the first to define an off-target effect of the antidiabetic medication sitagliptin on GH and the first study of the effect of DPP4 inhibition on the GH axis to include women. An understanding of the effect of DPP4 inhibition on GH can only be achieved by studying humans because of significant interspecies variation in the neuroregulation of GH secretion. Bergman et al examined the effect of 10-day treatment with sitagliptin, in doses ranging from 25 mg daily to 300 mg twice daily, on IGF-1 levels in 8 healthy young men. Although IGF-1 increased in nearly every treatment group after 10 days, the change was not statistically significant. Schopman et al reported that sitagliptin decreased GH levels after insulin-induced hypoglycemia in insulin-dependent men with diabetes mellitus and attributed this to an inhibitory effect of GLP-1 stimulation. GLP-1 is unlikely to contribute to enhanced GH secretion in women in the present study, as GLP-1 receptor blockade did not influence GH secretion.

We found an effect of sitagliptin in women but not in men, consistent with the mechanism of known sexual dimorphism in GH secretion. The observation that sitagliptin affects GH secretion in women only may be explained by work of others, which demonstrates that GHRH drives GH pulsatility in women and that interpulse GH secretion in women is more GHRH dependent and less susceptible to feedback inhibition by free IGF-1. We also observed that arginine did not stimulate GH secretion as effectively in men. This dose of arginine may have been insufficient for us to detect a significant effect of sitagliptin on stimulated GH secretion in men. Given these known differences in GH secretion between men and women, a separate adequately powered study is needed to investigate what effect sitagliptin has on GH secretion in men.

Our study is unique in that it investigates for the first time in humans the specific contribution of GHR activation and IGF-1 to the vasodilation observed after an increase in GH. The vasodilator effects of GH and IGF-1 in humans are well described. Napoli et al demonstrated that intra-arterial GH resulting in a 10-fold increase in GH levels in the human forearm with no effect on IGF-1 doubles forearm blood flow several hours later. IGF-1 causes relaxation in harvested human internal mammary artery that is unaffected by removal of the endothelium and NO inhibition but abolished by potassium chloride, suggesting that IGF-1–induced vasorelaxation in humans involves the potassium channels in vascular smooth muscle cells. We are not aware of any studies evaluating the effect of IGF-1 on the vasculature in the human forearm.

We found that GHR blockade paradoxically increased vasodilation following arginine-stimulated GH release in sitagliptin-treated women, even though IGF-1 concentrations were decreased. Increased vasodilation correlated with GH concentrations, which were increased as a result of loss of feedback inhibition, consistent with a GHR- and IGF-1–independent vasodilator effect of GH. Consistent with this mechanism, chronic pegvisomant therapy in patients with acromegaly improves flow-mediated dilation. GH treatment
in hypophysectomized rats upregulates vascular smooth muscle ATP-sensitive potassium channel mRNA, which could lead to depolarization and vasorelaxation. Lastly, adults with primary GH resistance, characterized by elevated GH but defective GHR signaling and IGF-1 deficiency, demonstrate normal conduit artery vascular function. 

Figure 4. The addition of growth hormone (GH) receptor blockade (pegvisomant 80 mg administered SC 72 hours prior) to sitagliptin increases GH levels as a result of reduced negative feedback (n=5 women) (A) and decreases free insulin–like growth factor-1 (IGF-1) (B). The addition of GH receptor blockade to sitagliptin further decreases forearm vascular resistance (FVR) during stimulated GH secretion (C). At the nadir in vascular resistance (arrow), a significant correlation between vascular resistance and GH levels was found (D). Pegvisomant decreases tissue plasminogen activator (tPA) activity levels before and throughout arginine (Arg)-stimulated GH secretion (E). Data are presented as mean±SEM. *P≤0.05 vs placebo at specified time point in the linear model. Linear model–based P values are: P<0.01 effect of treatment on GH, P<0.01 effect of treatment on FVR percent change (Δ), and P<0.001 effect of treatment on tPA activity.
Late vasodilation after arginine is believed to be mediated by GH, as octreotide infusion blocks arginine-stimulated GH secretion and prevents vasodilation. \(^{36}\) GH has also been reported to cause vasodilation via NO-dependent mechanisms. Prior studies indicate that coinfusion of LNMMA decreases the vasodilator response to GH. \(^{4}\) Li et al\(^3\) corroborated these findings using an acute systemic GH infusion and further demonstrated that GH increases phosphorylation and activity of endothelial NO synthase in human aortic endothelial cells in vitro. Others have found that chronic GH therapy, which also increases IGF-1 levels, increases markers of NO bioavailability, decreases peripheral resistance, and improves conduit artery vascular function. \(^{37,38}\) Our results do not support a contribution by NO to enhanced vasodilation following arginine-stimulated GH secretion during DPP4 inhibition, as vasodilation was not blocked by LNMMA and thus was NO synthase independent.

Sitagliptin also increases levels of intact GLP-1 and insulin, and intact GLP-1 has previously been reported to enhance the vasodilator response to intra-arterial insulin in adults with metabolic syndrome. \(^{39}\) It is unlikely that increased insulin and GLP-1 receptor activation contributed to the enhanced vasodilation during sitagliptin. Insulin levels were identical across study days and the mechanism of vasodilation was NO synthase independent. While sitagliptin increased fasting GLP-1 levels, the addition of GLP-1 receptor blockade did not prevent vasodilation. Furthermore, our group previously demonstrated that intra-arterial infusion of GLP-1 in the setting of sitagliptin has no effect on vasodilation in the forearm of healthy adults. \(^{40}\) Ban et al\(^{41}\) demonstrated that GLP-1 (9-36) increases endothelium-dependent vasodilation in mice lacking a GLP-1 receptor. A GLP-1 receptor-independent mechanism involving GLP-1 (9-36) is unlikely to explain the

**Figure 5.** L-N-monomethylarginine (LNMMA) did not affect the vasodilator response to arginine (Arg)-stimulated growth hormone (GH) secretion during sitagliptin (A) \((n=7\) women). Glucagon-like peptidase-1 (GLP-1) receptor blockade (Exendin 9-39) did not affect the vasodilator response to stimulated GH secretion during sitagliptin (B) \((n=7\) women). Data are presented as mean±SEM. \(^*P<0.05\) vs placebo at specified time point in the linear model. The linear model–based \(P\) values for overall effect of treatment were not significant.

**Figure 6.** The increase in arginine (Arg)-stimulated growth hormone (GH) secretion during dipeptidyl peptidase-4 inhibition with sitagliptin is reproducible \((n=19\) women). Data are presented as mean±SEM unless otherwise noted. There was a significant correlation between stimulated GH secretion following sitagliptin and stimulated GH secretion following sitagliptin plus saline infusion (peak GH response: \(r_s=0.65, P=0.003;\) GH 30 minutes after arginine: \(r_s=0.51, P=0.02\)).
observed vasodilation in this study as sitagliptin prevents the formation of GLP-1 (9-36). It is also possible that the enhanced vasodilation which followed GHR blockade was mediated in part by peripheral GHRH or GH activation of the prolactin receptor. Prolactin and GHRH receptor antagonists are not presently available for use in humans.

We observed an increase in tPA activity during stimulated GH secretion that was potentiated by sitagliptin in women. We previously demonstrated that adults with GH deficiency have decreased tPA activity, along with a defective fibrinolytic response to venous occlusion.5 Miljic et al6 also reported an improvement in stimulated endothelial tPA release following venous occlusion after 1 year of GH replacement aimed to normalize IGF-1 in GH-deficient adults. We demonstrated that GH mediates tPA activity levels through the GHR in women, as the addition of GHR blockade suppressed tPA activity before and during stimulated GH secretion. Muller et al42 found no effect of pegvisomant on tPA activity in men, whereas we detected a decrease in tPA activity.

Study Limitations and Strengths
Our findings are limited by a few study design considerations. We studied healthy, lean individuals to avoid medications and diseases that may affect stimulated GH secretion, endothelial function, and fibrinolysis. We studied patients who fasted, while sitagliptin exerts its incretin-specific effects in the postprandial state. Because sitagliptin may influence GH secretion by lowering postprandial blood sugar and free fatty acids, sitagliptin could have a greater effect on GH secretion in the postprandial state. We administered sitagliptin for a limited duration to decrease the likelihood of an increase in IGF-1 influencing GH secretion through feedback inhibition. The effect of chronic sitagliptin on GH secretion may be less pronounced. Measurement of plasma GHRH levels was not performed, given the several 1000-fold dilution of pituitary-portal GHRH and the inability of assays to distinguish between pituitary and peripheral GHRH. We used arginine as a stimulus for endogenous GH secretion. Other currently available GH secretagogues, insulin and glucagon, may be unsafe to administer during sitagliptin and would cause hormonal changes that would confound results. While arginine-stimulated GH secretion is unaffected by phase of the menstrual cycle,43 vasodilation is highest during the late follicular phase when estradiol levels are highest.44 We were limited in our ability to coordinate all assessments to the same phase of the menstrual cycle in our participants; however, we did not observe a difference in estradiol levels across study treatments. Similar to other researchers, we found low intraindividual variability in the GH response to arginine.45 In fact, a strength of our study is the highly reproducible stimulated GH secretion profile during sitagliptin in women. We were not able to confirm complete blockade of the GHR. The resulting increased GH levels may have been sufficient to stimulate an incompletely blocked GHR albeit to a lesser degree. Our study is also limited by our inability to define the mechanism by which increased GH causes sex-specific vasodilation. Further study using glibenclamide in an appropriate study population may elucidate a contribution by the vascular smooth muscle ATP-sensitive potassium channel. Lastly, our limited sample size in the second half of the study limits our conclusions on the impact of NO synthase inhibition and GLP-1 receptor blockade.

Conclusions
In this study, we tested a previously unexplored method to enhance endogenous pulsatile GH secretion in humans using the oral antidiabetic DPP4 inhibitor sitagliptin. Our data are the first to support an off-target effect of DPP4 inhibition on GH secretion in women. This is clinically relevant for 2 reasons. First, the cardiovascular effects of DPP4 inhibition and how these differ from the effects of GLP-1 analogues is an active area of investigation. Our findings suggest that DPP4 inhibition influences vasodilation by influencing GH secretion in women. Second, while current strategies to increase GH cause hyperglycemia, oral DPP4 inhibitor therapy offers an attractive, novel mechanism to enhance endogenous GH secretion while also improving glucose metabolism. It was previously well established that increases in GH and IGF-1 improve fibrinolytic capacity, vasodilator function, and lower inflammation. Medications that potentiate GH secretion may thus prevent atherosclerosis through these mechanisms. Last, an enhanced understanding of the specific effects of GH, GHR activation, and IGF-1 on cardiovascular risk and how they are modified by sex is critical to how we incorporate somatotropic medications in the management algorithms for patients with impaired GH secretion and elevated cardiovascular risk.

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Disclosures

None.

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SUPPLEMENTAL MATERIAL
Data S1.

Supplemental Methods

Forearm Blood Flow Measurements

Forearm blood flow (FBF) was measured using mercury-in-silastic strain-gauge plethysmography. The wrist was supported in a sling to raise the level of the forearm to above the level of the atrium, and a strain gauge was placed around the widest part of the forearm of the non-dominant hand. The strain gauge was connected to a plethysmograph (model EC-6, D.E. Hokanson; Issaquah, WA) connected to a chart recorder to record flow measurements. For each measurement, a cuff placed around the upper arm was inflated to 45 mmHg with a rapid cuff inflator (model E-20 rapid cuff inflator and AG 101 cuff inflator air source, Hokanson; Issaquah, WA) to occlude venous outflow from the extremity. The hand was excluded from the measurement of blood flow by inflation of a pediatric sphygmomanometer cuff to 200 mmHg around the wrist. Flow measurements were recorded for approximately seven seconds, and a minimum of 6 readings were analyzed using Non-Invasive Vascular Program (Hokanson NIVP3 version 5.40; Bellevue, WA) software to obtain each mean. Forearm vascular resistance (FVR) was calculated as mean arterial pressure (MAP) divided by FBF.

Sample Size Calculation

Our preliminary data in women demonstrated a difference in mean peak GH following arginine stimulation of 6.79 ng/mL. Sample size was calculated with PS Software using the design for a paired t-test with a 0.05 two-sided significance level.¹ A sample size of 28 women provided 80% power to detect a difference in means of 6.79 ng/mL, assuming a 12.25 ng/mL SD of the differences. Our preliminary data in men demonstrated a difference in mean peak GH in the opposite direction. Pre-menopausal women are more responsive to GHRH and secrete more
GH per pulse, a GHRH-mediated phenomenon, than men. Given the known sexual dimorphism in pulsatile GH secretion, we chose to study only enough men to permit us to estimate the difference in males. A sample size of 11 men provided a half-width of the 95% confidence for the difference to be 7.6 ng/mL. Accounting for 10% drop-out, we enrolled 31 women and 12 men.

Supplemental Results

Safety

Thirty-nine subjects (29 women) completed all study procedures in the first half of the study protocol; four subjects completed only one study day and their data was excluded from the analyses. Non-serious adverse events reported during the first half of the study included bruising related to placement of the intravenous catheter (3 subjects), palpitations after sitagliptin (1 subject), nausea after sitagliptin (3 subjects), nasal congestion after sitagliptin (1 subject) and reflux after sitagliptin (1 subject). Nineteen women completed all study procedures in the second half of the study protocol. One subject completed only one study day and her data was excluded from the analysis. Non-serious adverse events reported during the second half of the study included dizziness and paresthesias in lips during arginine infusion (1 subject) and 24 hours of abdominal cramping and diarrhea following subcutaneous injection of pegvisomant (1 subject). There were no symptoms observed during LNMMA or Exendin 9-39 infusions. There were no instances of hypoglycemia. There were no serious adverse events.
Figure S1. Subjects participated in a two-part double-blind, randomized, placebo-controlled, crossover study.

In the first half of the study (A), thirty-nine subjects underwent two study days in which they were randomized to sitagliptin or placebo. Sitagliptin was given as a single 200 mg dose one hour prior to arginine in the first 14 subjects (7 women), in order to achieve greater than 80 percent inhibition of DPP4 activity within one hour. We changed to sitagliptin 100 mg daily for four days in the remaining 25 subjects to achieve steady-state dosing and to be consistent with the current FDA-approved dose. The washout period was one week in the first 14 subjects and two weeks thereafter. In the second half of the study (B), 19 of the 29 women from the first half of the study were given sitagliptin on each of two identical study days. The dose of sitagliptin given was identical to that in the first half of the study. Subjects were divided into three subgroups, in which they were randomized to treatment with placebo or one of three antagonists in a crossover fashion.
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