Dose-Dependent Cortisol-Induced Increases in Plasma Leptin Concentration in Healthy Humans

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Background: Leptin is a hormone that regulates fat metabolism and appetite. The secretion of leptin is regulated by adiposity and, in the rodent, by factors such as insulin, β-adrenergic agonists, and glucocorticoids (GCs). Increased secretion of the endogenous human GC, cortisol, occurs during stress and in disorders such as major depression. Pharmacological GCs can robustly increase plasma leptin concentrations in humans, leading us to hypothesize that cortisol may serve as a physiological regulator of human leptin secretion.

Methods: A randomized double-blind placebo-controlled comparison of 2 fixed oral dosages of cortisol (40 mg/d and 160 mg/d), given for 4 days to matched groups of healthy subjects (n = 47). Low-dose treatment approximated GC output during mild stress, while high-dose treatment approximated GC output during maximal stress, spanning a range of GC secretion relevant to physiological stress.

Results: Cortisol produced dose-dependent and time-dependent increases in plasma leptin concentrations (time x treatment condition x body mass index; F6,123 = 10.73; P < .001). Initial treatment-induced increases in plasma leptin concentration returned toward baseline values during 4 treatment days, suggesting tolerance to this GC effect in these healthy subjects.

Conclusions: The results indicate an important role for GCs in the short-term regulation of human leptin secretion. Glucocorticoid-induced increases in leptin secretion suggest a mechanism that may contribute to anorexia and weight loss during stress and disease states such as major depression, if these conditions are associated with sustained increases in plasma leptin concentrations.

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SUBJECTS AND METHODS

SUBJECTS

All subjects (N = 47) gave written informed consent for participation in a protocol approved by the Human Studies Committee at Washington University School of Medicine, St Louis, Mo. Subjects were recruited using local advertisement and screened for general medical health and psychiatric disorders using the Diagnostic Interview for Genetic Studies. The inclusion criteria were subjects aged between 18 and 30 years and able to give informed consent. The exclusion criteria were as follows: (1) non–insulin-dependent or insulin-dependent diabetes mellitus; hypertension (treated or untreated); any major surgery in the preceding 6 months; any cardiac condition causing documented hemodynamic compromise; any respiratory condition causing documented or clinically recognized hypoxia; fever; dehydration; nausea; epilepsy; other endocrine disease; body weight less than 80% ideal body weight; any other medical condition requiring more than 7 days of hospitalization in the preceding 4 weeks; pregnancy or high-dose estrogen therapy; narcotic therapy; corticosteroid or spironolactone therapy; or psychotropic therapy; (2) any Axis I psychiatric disorder, including any substance use disorders; and (3) mental retardation as defined by DSM-IV criteria. Clinical characteristics of the sample were as follows (±SD): mean age = 22.4 years (2.8 years); BMI = 23.19 kg/m² (3.95 kg/m²); 23 men; 24 women. Thirteen women were taking oral birth control, and, of the remaining 11, 6 were in the luteal phase and 5 were in the follicular phase of their menstrual cycle at study baseline (as defined in the analysis section below). No significant differences in age (F₂,₄₄ = 0.034; P = .9), BMI (F₂,₄₄ = 1.59; P = .2), baseline plasma leptin levels (F₂,₄₄ = 0.67; P = .5), cortisol levels (F₂,₄₄ = 1.59; P = .2), insulin levels (F₂,₄₄ = 0.009; P = .9), or glucose levels (F₂,₄₄ = 0.01; P = .99) were detected across the high-dose CORT (7 men and 7 women), low-dose CORT (6 men and 8 women), or placebo-treated groups (10 men and 9 women). In addition, no significant main effect of sex or sex–by–treatment-condition interaction was detected for any of these variables.

PROCEDURE

The present study was a randomized double-blind placebo-controlled comparison of 2 fixed oral dosages of CORT (hydrocortisone), 160 mg/d and 40 mg/d, given for 4 days. All subjects received 1 of the 2 oral CORT doses or placebo using matched capsules given in divided doses (at 7 AM and 7 PM), using an approximately 3:2 dose ratio (high dose, 100 mg:60 mg; low dose, 25 mg:15 mg) to approximate circadian changes in endogenous CORT. Low-dose treatment approximated CORT output during mild physiological stress, while the high-dose treatment approximated CORT output during maximal physiological stress. Plasma sampling and behavioral and adverse effect measurements were obtained at baseline (study day 0), after 1 day and 4 days of treatment (study days 1 and 4), and after a 6-day

Dose-dependent CORT-induced increases in plasma leptin concentrations were detected in both male and female subjects, with greater increases in plasma leptin in women vs men and in individuals with higher BMI. The initial analysis-of-variance model testing the effect of treatment condition and BMI on serial plasma leptin concentrations in the total subject sample detected an interaction between plasma sample time and treatment condition (F₆,₁₂₃ = 8.18; P < .001; G-G adjustment) and a 3-way interaction among sample time, treatment condition, and BMI (F₆,₁₂₃ = 10.73; P < .001; G-G adjustment). The strength of these results is explained in part by the observation that 100% of CORT-treated subjects demonstrated a numerical increase in leptin concentrations on day 1, compared with baseline. The
washout period (study day 10). All assessments were performed at approximately 4 PM, with no food permitted after 1 PM. Previous investigations have established that plasma leptin concentrations do not undergo significant diurnal variation in the absence of prolonged starvation or massive caloric loading.

PLASMA MEASURES

Plasma leptin concentration was determined by radioimmunoassay (Linco Research, St Louis, Mo), using purified recombinant human leptin as a standard and as a radiolabel and a rabbit polyclonal antibody to human leptin, with a second antibody separation method. The assay limits of detection and linearity were 0.5 ng/mL and 100 ng/mL in plasma, with leptin values greater than 100 ng/mL assayed again on dilution. Intra-assay and interassay coefficients of variation were less than 7%, with no cross-reactivity with common peptide hormones. Plasma glucose was measured with a glucose oxidase method (Beckman Instruments, Fullerton, Calif). Plasma insulin and total CORT were measured by radioimmunoassay.

ANALYSIS

The comparability of the different treatment group characteristics (eg, age and BMI) was tested using analysis of variance. The main hypothesis concerning treatment effects on plasma leptin concentration was first tested using analysis of variance to evaluate leptin variance as a function of plasma sample time, treatment condition, and BMI. Body mass index was included even in the initial model based on previous work from this and other laboratories indicating a critical effect of adiposity on leptin levels. The hypothesis was further tested using more inclusive analysis of variance and analysis of covariance models, that included subject sex and plasma covariates (CORT, glucose, and insulin). Significant effects were further evaluated using additional models and simple regressions within separate groups to better understand the contributions of individual continuous and factorial variables. To assess the validity of unadjusted F tests on the within-subject “time” factor and its interactions, the sphericity assumption for this data set was assessed using the test for sphericity applied to orthogonal components. Because the test for sphericity was significant (Mauchly criterion, 0.288; χ² = 53.2; P < .001), it is appropriate to use P values for the univariate tests of hypotheses for within-subject effects that have the df adjusted. Two tests were applied that use adjusted P values, the Greenhouse-Geisser (G-G) and Huynh-Feldt adjustments. The former provided the most conservative estimate of significance in this analysis, and this test is presented below as appropriate. The effects of phase of menstrual cycle and oral birth control on plasma leptin concentrations were explored using unpaired t tests, comparing plasma leptin values across female subjects receiving oral birth control, female subjects in the luteal phase of their menstrual cycle (defined as day of cycle at baseline ≤ [cycle length in days/2]), and female subjects in the follicular phase of their menstrual cycle (defined as day of cycle ≥ [cycle length in days/2]).

Results can be further understood by examining male vs female subjects (Figure 1). A more inclusive model for predicting leptin levels detected a significant interaction among time, treatment condition, sex, plasma CORT concentrations, and BMI (F [3,98] = 3.47; P < .001). This was explained by larger CORT-induced leptin increases in women (Figure 1), subjects receiving higher dose CORT treatment, and those with higher BMI (Figure 2). Importantly, CORT-induced increases in plasma leptin levels in these healthy subjects were detected after 1 treatment day, but were not sustained after 4 days of treatment, despite sustained CORT elevation.

Dividing subjects into obese (defined as men with BMI of ≥ 27.3 kg/m² and women with BMI of ≥ 27.8 kg/m²) vs lean individuals, regressions for plasma leptin levels vs CORT levels on day 1 were significant for low (F [1,42] = 5.78; P = .02), but not high BMI, possibly because of the few obese subjects in our sample. Notably, the slopes of the regression lines for both lean and obese subjects were nearly identical (0.367 and 0.383, respectively), suggesting that data from additional obese subjects might fall along the same line. Higher mean plasma leptin levels in the male subjects receiving low-dose vs high-dose CORT treatment (Figure 1, bottom) were accounted for by a single obese male subject in the low-dose treatment group with higher baseline leptin and a robust CORT-induced leptin response. Exploring the effects of varying ovarian endocrine status in the female subjects using unpaired t tests, no differences in plasma leptin concentrations were detected among women in the luteal vs follicular phase of their menstrual cycle (as defined above) vs those receiving oral birth control at either baseline or at study day 1, consistent with most but not all, previous reports.

As expected, a significant plasma sample time × treatment condition interaction was detected on circulating plasma CORT concentrations (F [1,123] = 9.54; P < .001; G-G adjustment), with low-dose CORT treatment producing little change in plasma CORT concentrations measured at 4 PM compared with placebo, while the higher dose produced stable stress-physiological CORT concentrations at 4 PM throughout the 4 days of treatment (Table). A main effect of sex on plasma CORT levels (F [1,43] = 8.09; P = .007) was explained by higher overall CORT levels in women vs men. In contrast to these differences in overall levels, treatment conditions produced a similar pattern of effects on plasma CORT levels across men and women (time × treatment condition × sex interaction: F [1,123] = 1.64; P = .17; G-G adjustment). This result indicates that sex differences in leptin response are probably not explained by sex differences in the plasma CORT response to fixed CORT doses. A significant treatment effect on plasma glucose levels (time × treatment condition: F [1,123] = 5.19; P < .001; G-G adjustment) was explained by a treatment-induced rise in plasma glucose during the high-dose CORT condition only (Table).
Pharmacological doses of GCs can increase circulating leptin levels by establishing the significance of this effect for humans using CORT doses relevant to physiological stress. This study demonstrates dose-dependent and time-dependent CORT-induced increases in circulating leptin levels in young healthy humans, with greater dose-dependent CORT-induced increases in leptin levels in women and subjects with higher BMI. Body mass index remained a strong predictor of plasma leptin levels in this study as in our previous report, interacting with CORT remaining a strong predictor of plasma leptin levels in this study as in our previous report,22 interacting with CORT and leptin levels in obesity. Interestingly, plasma leptin levels escaped from the stimulating effects of sustained CORT exposure, suggesting tolerance to GC-induced stimulation or a homeostatic down-regulation of GC effects on leptin during the 4-day treatment interval in these healthy subjects.

In contrast, leptin levels remain increased over time in patients with Cushing disease even after controlling for increased BMI, suggesting that some disease processes may interfere with the down-regulation of GC effects on leptin secretion. In addition, the failure of leptin levels to decrease after correction of hypercortisolism in patients with pituitary Cushing disease suggests the possibility of long-term complex changes in mechanisms coupling GCs to leptin secretion. However, the interpretation of studies in patients with Cushing disease is complicated, with mechanisms coupling GCs to leptin secretion and other disease factors confounded. Future studies might address longer periods of cortisol treatment in healthy subjects and measure leptin activity in patients with major depression and other disease states associated with hypercortisolism and weight loss. Physiological CORT elevations in such diverse clinical states as depression, dementia, severe illness (eg, acquired immunodeficiency syndrome wasting syndrome), and stress might reduce appetite and weight, at least in part, through increases in circulating leptin levels. However, more basic studies confirming leptin effects on appetite and weight in humans are still needed.

Pharmacological doses of GCs have clinically recognized orexigenic effects, and GC-induced leptin secretion might be conceptualized as a counterregulatory mechanism limiting GC-related hyperphagia in all but pharmacological GC dosing conditions. Alternatively, some investigators have proposed that chronic exposure to excess GCs might result in a down-regulation of leptin activity even to the point of leptin insensitivity and the development of obesity. The escape of plasma leptin levels from the stimulating effects of sustained CORT exposure in the healthy subjects we studied is consistent with this proposal and might be understood as an adapt-
Significance is given for treatment condition elimination half-life of 1.5 hours may explain the observation, allowing plasma CORT levels to decrease toward the end of each dosing interval. The relatively brief CORT plasma reduction, allowing plasma CORT levels to decrease toward the end of each dosing interval. This distinction between plasma levels of 4 PM was no longer increased for the 40-mg/d group compared with the placebo group (Table). However, the biological (vs plasma elimination) half-life of CORT is 8 to 12 hours, suggesting that CORT effects on leptin secretion could persist throughout the dosing interval. This distinction between the plasma elimination time and biological effects of CORT, as well as the continuous CORT infusion data from humans noted above, may explain why increases in plasma leptin concentrations were detected in female subjects even in the lower 40-mg/d CORT treatment group despite no increase in plasma CORT concentration at 4 PM.

While leptin effects on appetite, physical activity levels, and calorigenesis are well characterized in the rodent, the physiological significance of leptin activity in humans remains the subject of intensive ongoing investigation. In the future, it will be critical to define the short-term and extended physiological effects of increased plasma leptin concentrations in humans. In summary, the results of this study indicate an important physiological role for GCs in the short-term regulation of leptin secretion in humans. These GC-induced increases in leptin secretion suggest a mechanism that may contribute to anorexia and weight loss during stress and disease states such as major depression, if these conditions are associated with sustained increases in plasma leptin concentrations.

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