Association of Cortisol and the Metabolic Syndrome in Korean Men and Women

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INTRODUCTION

Glucocorticoids, mediated by activation of the hypothalamic-pituitary-adrenal (HPA) axis, have an impact on metabolic responses, insulin-resistance and lipolysis (1). Glucocorticoids directly impair insulin sensitivity in adipocytes (2) and promote free fatty acid release from mature adipocytes through hormone sensitive lipase (HSL)-mediated lipolysis (3). These events are mediated by Glucocorticoid Receptors (GR). The density of these receptors is higher in intraabdominal visceral than other fat depots, and the activity of cortisol will be accentuated in this particular adipose tissue (4). It is now apparent that tissue-specific changes in cortisol metabolism is primarily increased intracellular glucocorticoid reactivation, catalysed by the enzyme 11-hydroxysteroid dehydrogenase type (HSD)-1 in adipose tissue (5).

Abdominal obesity promotes insulin resistance and is considered a main part of the metabolic syndrome (6, 7). Although cortisol has been implicated as a pathophysiological contributor to idiopathic obesity, circulating cortisol has not consistently been shown to be elevated in obese as compared to normal weight individuals (8). A recent study using both cross-sectional and longitudinal analysis to determine the associations between body composition and serum cortisol concentrations in men showed that weight change drives the changes in cortisol concentrations, rather than vice versa (9).

In addition, studies have shown that cortisol may add to the effect of adiposity on metabolic risk. Ward et al. (10) reported that higher cortisol concentrations increased the risk for cardiovascular disease (CVD) risk factors in a South Asian cohort study, above what could be explained by the effect of adiposity alone. In that study, associations between cortisol concentration and CVD risk factors were also stronger than what has previously been shown in Caucasian populations, despite similar cortisol levels. Asian populations have been shown to have a different fat distribution pattern than Western populations and seem to be more prone to central obesity and metabolic dysregulation even at lower body mass index (BMI) levels (11, 12). However, increased prevalence of overweight and obesity does not fully explain the higher prevalence of the metabolic syndrome in Asian populations (12-14), and the role of glucocorticoids has not been investigated in Korean individuals. We hypothesized that cortisol would be positively and independently associated with increased CVD risk, assessed as components of the metabolic syndrome.

Obesity and the metabolic syndrome are closely related and have become increasingly prevalent in Korea. The cardiovascular disease (CVD) risk factors comprising the metabolic syndrome have previously been associated with increased hypothalamic-pituitary-adrenal axis (HPAA) activity, but the associations have not been extensively examined in non-Caucasian populations. The aim of the present study was to investigate the relationships between cortisol, adiposity and the metabolic syndrome in a Korean population. A total of 1,881 adults participated in the study between January 2001 and February 2008. Sociodemographic data were assessed by questionnaires. Body composition, clinic blood pressures as well as metabolic variables including glucose, insulin, and lipid profile were assessed and analyzed in relation to cortisol levels. Mean age of the participants was 58.7 ± 10.8 yr. Higher levels of cortisol was associated with elevated blood pressure, fasting glucose and total cholesterol in men, and between cortisol and systolic blood pressure, fasting glucose and total cholesterol in women. There was an increased risk for the metabolic syndrome associated with higher cortisol levels in both men (P < 0.001) and women (P = 0.040) adjusting for age and body mass index. Higher cortisol levels are associated with several CVD risk factors and the metabolic syndrome, independent of overall adiposity level, in Korean men and women.
MATERIALS AND METHODS

Study population
We enrolled 2,219 Korean individuals > 20 yr old who had undertaken medical evaluation at Ajou University Hospital between January 2001 and February 2008. Participants were excluded if they were taking steroid medication or did not provide information about smoking status, alcohol consumption, or physical activity (n = 338) leaving a sample of 1,881 participants (56.1% female).

Data collection
Sociodemographic characteristics, smoking status, alcohol habits, and activity level were assessed by questionnaire. Socioeconomic status (SES) meant financial affordability; and exercise was defined by moderate or severe vs none or sedentary. Height and weight were measured using bioelectrical impedance analysis (Inbody 3.0, Biospace, Seoul, Korea) following overnight fast. BMI was calculated as weight divided by height squared (kg/m²) (15). Waist circumference was measured between the lower rib and iliac crest by a trained nurse. Blood pressure was measured by a semi-automated blood pressure monitor (TM-2650A, PMS Instruments, Tokyo, Japan) after rest of at least 15 min. Venous blood was drawn following an 8-hr overnight fast and 24-hr abstinence from vigorous activity. Fasting glucose, total cholesterol and triglyceride level were assessed with an enzymatic colorimetric method (TBA-200FR, Toshiba, Tokyo, Japan); Serum insulin was determined with an immunoradiometric assay kit (Dnabot, Tokyo, Japan) and cortisol with a RIA method (Diagnostic Products Cooperation, Los Angeles, CA, USA). We calculated the insulin resistance index using the homeostasis model assessment (HOMA) of insulin resistance (16).

Statistical analysis
We followed the NCEP-ATP III Asian guideline components to define metabolic syndrome, comprised of central obesity (waist circumference ≥ 90 cm for men and ≥ 85 cm for women), blood pressure ≥ 130/80 mmHg, triglyceride ≥ 150 mg/dL, fasting glucose ≥ 110 mg/dL, and high-density lipoprotein cholesterol (men < 40 mg/dL and women < 50 mg/dL). In line with the guidelines, subjects who met three or more of the criteria mentioned above were defined as having the metabolic syndrome (17, 18).

Chi-square test and Student’s t-test were used to calculate the difference in mean values of general characteristics and laboratory tests between men and women. Regression analysis was conducted to assess the association between cortisol and CVD risk factors while controlling for age and BMI. Further regression analysis was assessed to evaluate the relation of cortisol concentration to the separate CVD risk variables while simultaneously controlling for confounders such as SES, smoking status, alcohol intake and exercise. Finally, logistic regression analysis was used to evaluate the association of cortisol levels to the metabolic syndrome, controlling for age and BMI. A P < 0.05 was used as cutoff for significance. SPSS in version 12.0 was used for all statistical analysis.

RESULTS

Demographic characteristics
A total of 1,881 individuals were assessed, 826 men (43.9%) and 1,055 women (56.1%) (Table 1). The mean age of the subjects was 58.7 ± 10.8 yr and the mean BMI for all subjects was 24.3 ± 3.3 kg/m². The mean cortisol level for all subjects was 12.7 ± 4.8 µg/dL. Mean levels (± SD) of study variables are presented by gender in Table 1. There were gender differences on several demographic and clinical variables, including age, waist circumference (WC), BMI, cortisol levels, HDL cholesterol, total cholesterol, and systolic blood pressure. In addition, men had higher alcohol consumption and were more often smokers in comparison to women. Almost 32% of the women and 21% of the men met criteria for the metabolic syndrome.

Table 1. General characteristics of the study subjects

| Parameters          | Male (n = 826) | Female (n = 1,055) | P values |
|---------------------|---------------|-------------------|----------|
| Age (yr)            | 57.9 ± 11.8   | 59.4 ± 10.0       | 0.004    |
| WC (cm)             | 84.8 ± 9.0    | 83.6 ± 8.4        | 0.003    |
| BMI (kg/m²)         | 23.3 ± 3.2    | 24.7 ± 3.3        | < 0.001  |
| SES (%)             |               |                   | 0.215    |
| Low                 | 20.6          | 22.3              |          |
| Middle              | 72.9          | 72.6              |          |
| High                | 5.3           | 3.4               |          |
| Current smoking (%) | 36.4          | 4.2               | < 0.001  |
| Alcohol intake (%)  | 62.6          | 14.7              | < 0.001  |
| Exercise (%)        | 32.8          | 27.7              |          |
| MS ( IFNα (%), %)   | 176/650 (21.3/78.7) | 338/717 (81.9/68.0) | < 0.001  |
| SBP (mmHg)          | 127.2 ± 17.4  | 130.1 ± 18.8      | < 0.001  |
| DBP (mmHg)          | 78.8 ± 11.6   | 78.1 ± 10.7       | 0.148    |
| FBS (mg/dL)         | 104.8 ± 28.1  | 103.9 ± 25.4      | 0.487    |
| T-Chol (mg/dL)      | 192.9 ± 36.6  | 205.8 ± 38.3      | < 0.001  |
| TG (mg/dL)          | 145.1 ± 97.2  | 139.7 ± 94.9      | 0.216    |
| HDL-C (mg/dL)       | 51.9 ± 13.9   | 55.0 ± 13.8       | < 0.001  |
| Cortisol (µg/dL)    | 13.8 ± 4.6    | 11.8 ± 4.8        | < 0.001  |
| Insulin (µIU/mL)    | 7.0 ± 4.2     | 7.3 ± 4.8         | 0.460    |
| HOMA-IR             | 1.89 ± 1.57   | 1.91 ± 1.34       | 0.287    |

Values are mean ± SD unless otherwise indicated. BMI, body mass index; SES, socioeconomic status; MS, metabolic syndrome; SBP, systolic blood pressure; DBP, diastolic blood pressure; FBS, fasting blood sugar; T-Chol, total cholesterol; TG, triglyceride; HDL-C, high density lipoprotein cholesterol; HOMA-IR, homeostasis model assessment of insulin resistance. Gender difference of continuous variables were compared using t-test, categorical variables were compared by chi-squared test.
Results from the present study suggest that higher levels of fasting cortisol are associated with several CVD risk factors in Korean men and women, independent of overall adiposity level as measured by BMI. Our study also showed that the risk for the metabolic syndrome was greater in individuals with high cortisol levels after adjusting for age and BMI. This finding is consistent with previous research showing that individuals with met-
abolic syndrome have increased cortisol secretion (19, 20) and cortisol metabolite levels (21). Further adjustment for SES status, smoking, alcohol drinking and daily activity reduced this effect in women, but not in the men.

We found a negative association of cortisol to BMI and WC in the women, and no association between cortisol and BMI or WC in the men. These findings were consistent with previous studies showing that cortisol levels were not elevated in obese as compared to normal weight individuals (8, 9), and also in agreement with the study suggesting a diminished stimulability of the HPA axis in obese individuals (22). Cortisol is released from the subcutaneous adipose tissue by 11beta-HSD1 in humans, and increased enzyme expression in obesity is likely to increase local glucocorticoid signaling and contribute to whole-body cortisol regeneration (23). However, because the previous findings were not entirely consistent, it is still not clear whether the enzymatic overactivity is a cause or a result of obesity. In addition, our study is using a cross sectional design, and can therefore not assess whether cortisol levels actually contribute to weight gain over time. Hence, the present study can not test whether hypercortisolemia favors accumulation of visceral fat as initially suggested by Bjorntorp (4) and a recent review (24).

The present study showed that cortisol levels were associated to fasting glucose levels in both men and women, even after controlling for age and BMI. This is in line with the notion that glucocorticoids stimulate hepatic gluconeogenesis, leading to increased fasting glucose (25). In addition, glucocorticoids inhibit insulin actions on skeletal muscle and potentiate actions on adipose tissue, promoting the development of the metabolic syndrome (26). Cortisol counteracts the insulin activation of glycogen synthesis in muscle, the insulin inhibition of hepatic glucose production and the insulin inhibition of lipolysis in adipose tissue, leading to systemic insulin resistance (27). Unfortunately, insulin was only available in a subsample of the present study participants. Our observation is somewhat consistent with in vivo and in vitro experiments showing that glucocorticoids regulate insulin secretion (28).

This differs from previous observations showing a synergistic effect of cortisol and adiposity on CVD risk factors. For example, studies have shown that the correlation between cortisol and systolic blood pressure is most marked in obese subjects (29). Although the mechanism of this interaction is unclear, a recent study suggests that glucocorticoid action may be increased in skeletal muscle, an important insulin target tissue, in obese individuals (30).

Our study population consisted of patients who visited a University Hospital, which may limit the generalizability of our findings. Second, because we employed a cross-sectional design, it is not possible to determine whether cortisol was causally related to the development of the metabolic syndrome. Longitudinal studies could assess whether circulating cortisol concentra-

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AUTHOR SUMMARY

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The cardiovascular disease (CVD) risk factors comprising the metabolic syndrome have previously been associated with increased hypothalamic-pituitary adrenal axis activity. The aim of the present study was to investigate the relationships between cortisol, adiposity and the metabolic syndrome in a Korean population (1, 881 adults). Higher cortisol levels are associated with several CVD risk factors, independent of overall of adiposity level. In addition, there is an increased risk for the metabolic syndrome associated with higher cortisol levels after adjusting for age and BMI.