Comparison of serum androgens and endometrial thickness in obese and non-obese postmenopausal women

Obeye obez olmayan postmenopozal hastalarda serum androjen seviyeleri ve endometrial kalınığın karşılaştırılması

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Abstract

Objective: In this study, we investigated whether serum androgen levels and endometrial thickness differed in obese and non-obese women.

Material and Methods: Thirty-two non-obese (BMI <30) and 48 obese (BMI ≥30) women were enrolled. Blood samples were analyzed for testosterone, free testosterone, androstenedione, DHEAS, and SHBG, and transvaginal ultrasonography was performed.

Results: Obese women had significantly higher free testosterone and endometrial thickness and significantly lower SHBG. Eight of 17 women with endometrial thickness >5 mm had significant pathology.

Conclusion: These results suggest that obesity may be a risk factor for endometrial carcinoma and other pathologies in post-menopausal women through an action on androgen concentrations.

Methods: (J Turkish-German Gynecol Assoc 2010; 11: 149-51)

Key words: Serum androgene, endometrial thickness, obesity, post-menopause

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Introduction

The major hormonal changes associated with the occurrence of the final menstrual period (FMP) in normal subjects include a profound fall in circulating estradiol (E2), primarily of ovarian origin, beginning up to about one year earlier, and an accompanying large increase in the circulating gonadotropins, FSH and LH (1, 2). Changes in circulating androgens are more complex. In cross-sectional studies, levels of testosterone [T] have been reported to be lower in postmenopausal than premenopausal women (3, 4), whereas in the few longitudinal studies, either no change (5) or a small fall, of the order of 15% (2), in total T has been reported. The major adrenal androgen, dehydroepiandrosterone sulfate (DHEAS), reaches its peak levels in young women in their twenties and declines progressively thereafter, with no obvious relationship to the menopause (6). Serum sex hormone binding globulin (SHBG) levels fall to a small degree postmenopausally (1).

SHBG is responsible for the transport of sex hormones in the plasma, especially testosterone, dehydrotestosterone and estradiol (higher affinity for testosterone and lower for estradiol). It has a negative relationship with free testosterone and is used as an indirect indicator of androgenity (7). Transvaginal sonography (TVS) is a non-invasive and reliable method for evaluating the uterus and endometrium (8). In this study we investigated whether serum androgen levels and endometrial thickness differed in obese and non-obese postmenopausal women.

Material and Methods

Eighty postmenopausal women who applied to the menopause clinic of Istanbul Education and Research Hospital, Department of Obstetrics and Gynecology, between July 2004, were prospectively included in the study. These women had experienced natural menopause, had an FSH

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level greater than 40 mIU/ml, were non-smokers; and at least one year had passed since their last menstrual period. Seven women who had previously used hormone treatment or a medication that could affect body mass index, and three women who had genital or systemic pathology, were excluded from the study. PAP smears and physical examinations were performed after obtaining patient histories. Complete blood count, liver and renal functional tests, hormone profile (FSH, LH, E2, progesterone, prolactin, testosterone, free testosterone, androstenedione, DHEAS, SHBG, and fasting insulin levels) were carried out for each woman. After excluding any systemic disease, age, height and weight of each woman were noted at the first visit. Blood samples were taken after 12 hours fasting for the evaluation of sex steroids. Samples were centrifuged at 3000 rpm and sera were stored under -20ºC until evaluation of SHBG. A chemiluminescent enzyme immunometric assay technique was used for the determination of hormone concentrations, except SHBG, for which an enzyme linked immunosorbant assay (ELISA) was used. Transvaginal ultrasonography was performed with a Toshiba 2700 SA ultrasonography machine, and the criteria given by the American College of Obstetricians and Gynecology in 1995 was used for the measurement of endometrial thickness. The evaluation was performed in the three planes (anterior-posterior, long axis, semi-axial). Fluid collection in the cavity was not added to the thickness when making the measurement. We performed fractional curettage when we detected an endometrial thickness value of 5 mm or more. We added to the thickness when making the measurement. We performed fractional curettage when we detected an endometrial thickness value of 5 mm or more. We added to the thickness when making the measurement.

The women were allocated into two groups according to Body Mass Index (BMI), which was calculated as weight (kg) / height² (m²), and a cut-off value for obesity of 30 kg/m² used. Those with BMI > 30 kg/m² were considered as the study group.

Results

The mean height was 156.17±6.26 cm, weight 89.02±19.73 kg, BMI 32.85±6.79 kg/m², and endometrial thickness 5.56±0.06 mm. The demographic characteristics of the women according to BMI are listed in Table 1. The hormonal concentrations and endometrial thicknesses of the two groups are compared in Table 2. SHBG levels were significantly lower in the group in which BMI was greater than 30 kg/m², whereas free testosterone and endometrial thicknesses were significantly higher (Table 2). Only one of the 17 women whose endometrial thickness was higher than 5 mm had a BMI less than 30 kg/m². Fractional curettage was performed in these 17 women, and endometrial carcinoma was found in two, complex atypical hyperplasia in one, simple endometrial hyperplasia in one, and endometrial polyp in four. No pathology was detected in nine of these women.

Discussion

The relation between obesity and the increased endometrial cancer risk is well established in a review by Kaaks et al. and explained by the increasing amount of bioavailable estrogens in the circulation and the endometrial tissue (9). After the menopause, the major source of estrogen is via peripheral aromatization of androgens, especially in the adipose tissue (9). Furthermore, hyperinsulinemia in obese women inhibits the synthesis of SHBG (9). As a result, obesity is associated with increasing levels of estradiol unbound to SHBG, which can freely diffuse to target tissues. Many articles have reported a decrease in SHBG as the BMI increases (10-12), and we also found lower SHBG concentrations in the group which had the higher BMI. Douchi et al. reported a prospective study in which a significant relationship was demonstrated between endometrial thickness and obesity (13). Warming et al. demonstrated a positive correlation between BMI and endometrial thickness in a study that included healthy post-menopausal women (14). Serin et al. reported that increased endometrial thickness was associated with obesity but not with hypertension (15). We also observed increased endometrial thickness among women with high BMI and furthermore, independently of these surveys, we evaluated the role of androgens for this relation and detected high levels of free testosterone in obese women with increased endometrial thickness. Decreased SHBG concentrations as a result of increased BMI in post-menopausal women increase the free androgen and estrogen levels in the circulation, with a resultant increased risk of endometrial carcinoma. An increased endometrial thickness with increased BMI could be an indicator of this probable risk. The question then arises whether this increased

| Table 1. Demographic characteristics of the two groups according to BMI |
|-----------------|-------|-------|--------|-------|
|                 | BMI < 30 | BMI ≥ 30 |  |  |
|                 | Mean ± SD | Mean ± SD | p  |  |
| Age (years)     | 53.06 ± 6.58 | 52.54 ± 6.87 | 0.813  |  |
| Parity (n)      | 3.31 ± 1.66 | 3.08 ± 1.50 | 0.653  |  |
| Years post-menopause | 6.56 ± 8.35 | 5.43 ± 5.62 | 0.613  |  |
| Height (cm)     | 155.25 ± 5.30 | 156.17 ± 6.26 | 0.633  |  |
| Weight (kg)     | 66.56 ± 7.22 | 89.02 ± 19.73 | 0.0001 |  |
endometrial carcinoma risk is solely due to the increased estrogen concentration or not. Allen et al. have evaluated the endogenous sex hormones and endometrial cancer risk in women in a prospective case control study (16). They concluded that high levels of free testosterone and estrogens are associated with an increased endometrial cancer risk in postmenopausal women (16). Furthermore, they suggested that the association of endometrial cancer risk with free testosterone levels is a result of peripheral conversion of androgens into estradiol. Also, SHBG was significantly inversely and BMI positively associated with risk.

In another prospective case control study by Lukanova et al., it was concluded that increased endometrial cancer risk in postmenopausal women with high blood concentrations of androgens seems to be primarily due to their role as precursor hormones for estrogen synthesis (17). Interestingly, we have detected endometrial adenocarcinoma in two subjects, complex atypical hyperplasia in one, and simple endometrial hyperplasia in three. Free testosterone levels of these cases were high and SHBG levels were low. Although our study did not definitively demonstrate a relationship between elevation of androgens and endometrial pathologies, it indicates that the probability of such a relationship should be kept in mind. Considering free testosterone and SHBG concentrations together with the endometrial thickness may help to detect endometrial pathologies.

Future studies are needed to demonstrate conclusively that elevated androgen levels, as well as increased estrogen, can give rise to endometrial pathologies in postmenopausal women.

**Conflicts of interest**

No conflict of interest is declared by authors.

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