Short communication

Frequency of poly cystic ovary syndrome in patients with premenopausal breast cancer

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Abstract

Background: Polycystic ovary syndrome (PCOS) is a heterogeneous, complex genetic disorder characterized by hyperandrogenemia, hyperinsulinemia, insulin resistance, and chronic anovulation. It is the most common endocrine disorder in women of reproductive age with an enigmatic pathophysiologic and molecular basis. Obesity, hyperandrogenism, and infertility occur frequently in PCOS, which mostly have a genetic predisposition, and are features known to be associated with the development of breast cancer risk.

Objective: In present study, frequency of PCOS in patients with premenopausal breast cancer was compared with the frequency in women without breast cancer.

Materials and Methods: This is a case-control study, which compared PCOS frequency in 166 patients with premenopausal breast cancer and 166 healthy controls with normal mammography in last 6 months.

Results: Eleven patients (6.62%) in case group and 16 patients (9.63%) in control group had polycystic ovary syndrome according to their questionnaire. The difference was not significant (p=0.645).

Conclusion: There was no relationship between frequency of polycystic ovary syndrome and breast cancer in this study. This might be due to the age of patients with breast cancer in this study, which was mostly over 40. It could be significant if the patients were chosen in lower age for showing more effect of genetic than environment. The adjustment or matching of other risk factors could help to find the better results.

Key words: Polycystic ovary syndrome, Premenopausal breast cancer, Endocrine disorder.

Introduction

Polycystic ovary syndrome (PCOS) was seen in 15% of women at reproductive age (1). It is one of the most important endocrine problems in reproduction, which in it chronic anovulation and

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Nasrin Ghasemi, Department of Genetics, Research and Clinical Centre for Infertility, Shahid Sadoughi University of Medical Sciences, Yazd, Iran. **Email:** n479g@ssu.ac.ir hyperandrogenism happens (2). The risk of a number of gynecological neoplasias, including endometrial, breast, and ovarian cancer increase (3). women with PCOS Obesity, in hyperandrogenism, and infertility occur frequently in PCOS, and are features known to be associated with the development of breast cancer. However, studies examining the relationship between PCOS and breast carcinoma have not always identified a significantly increased risk (4). The endocrine abnormalities in PCOS including prolonged exposure to estrogen, progesterone deficiency, and androgen excess, could contribute to an increased risk for gynecological cancers. Previous studies have revealed that the etiology of breast cancer is linked to long exposure of breast epithelium to estrogens, especially in conjunction with progesterone (5). Long reproductive period (early age at menarche and/or late age at menopause) represents risk factors for this malignancy (6). As for the effect of pregnancy on breast cancer risk, there is a transient increase risk in the first 3-4 years after delivery of a baby, but lifetime risk seems lower than that of women who remain nulliparous (5). The elevated androgen levels and the increased levels of insulin and IGF-I detected PCOS and obesity could enhance the in development of breast cancer. PCOS usually accompanies with obesity, which have elevated androgen levels and increased levels of insulin and IGF-I. These conditions could enhance the development of breast cancer, which could happens to direct stimulation of AR-positive cancer cell in binding with androgen; stimulation **ER**-positive cells of by intratissular changes of testosterone to estradiol, mitogenic stimulation of cancer cells by insulin and IGF-I; and decreased levels of SHBG and increased levels of free estrogens in hyperandrogenic women (7-8). Because of the etiologic implications of the previously reported positive association between polycystic ovaries and breast cancer (9-11), present study attempted to confirm the relation. The study used data from a case-control study that compare frequency of PCOS among breast cancer patients with healthy controls.

Materials and methods

To investigate the relation between polycystic ovaries and breast cancer, frequency of PCOS was evaluated between 166 patients with breast cancer. Cases included premenopausal women with histologically confirmed primary breast cancer. This frequency was compared with frequency of PCOS among 166 women without breast cancer, which was confirmed by mammography in last six months. The age of cases and controls was matched. All cases and controls were fertile and the frequency of the use of contraceptive pill in both groups was not significantly different. PCOS patients in women with breast cancer and healthy controls were detected by face to face interview and filling the questionnaire, then it was confirmed by sonography in both groups. Cases and controls filled the informed consent form to be part of this study. The number of cases and

controls were calculated according to approximately 10% prevalence of PCOS in 2003 population. The Rotterdam consensus workshop was used to diagnose PCOS, which is a syndrome of ovarian dysfunction along with the cardinal features hyperandrogenism and polycystic ovary PCO. The results were analyzed by SPSS software (version 15) using chi-square test and odd ratio with CI.

Statistical analysis

The frequency of PCOS in women with breast cancer compare with healthy controls using chi-square test.

Results

Frequency of PCOS in 166 women with breast cancer was compared with normal controls. The mean age of the cases was 44.3 ± 3.80 (31 to 51 years old) and in controls was 44.2 ± 3.84 (30 to 50 years old). In this study, the frequency of PCOS in breast cancer patients was 6.62% compare to 9.63% in control group. The risk of breast cancer was lower among women with history of physician-diagnosed polycystic ovaries than women without such a history (Table I). The odd ratio was 0.66 (CI 95%: 0.299-1.48). Current data show that of women with breast cancer, 15.1% have hyperanderogenism, 6.6% have anovulation, 1.8% have obesity, and 1.8% have infertility, which were not significantly different with control group. There was not a relation between the frequency of PCOS signs and breast cancer (Table II).

Table I. PCOS frequency in women with breast cancer in comparison with women without breast cancer.

Groups	Without PCOS		With PCOS		Sum	
	No	%	No	%	No	%
With BC	155	93.38	11	6.62	166	100
Without BC	150	90.37	16	9.63	166	100

Chi square test

There was no significant relation between PCOS and breast cancer (p-value =0.645).

Odd ratio was 0.66 (CI 95%: 0.299-1.48). (BC = breast cancer)

Table II. Frequency of major signs of PCOS in women withbreast cancer compare to women without breast cancer.

Groups	Acnes/Hirsutism		Oligo/Amenorrhea		Sum	
	No	%	No	%	No	%
With BC	25	15.06	11	6.62	36	21.68
Without BC	29	17.46	16	9.63	45	27.09

Chi-square test

There is no significant relation between major signs of PCOS and breast cancer (p-value =0.773). (BC=breast cancer)

Discussion

Obesity, hyperandrogenism, and infertility are features of PCOS, which were known to be associated with the development of breast cancer. However, the epidemiological data in the literature about PCOS and breast cancer risk are contrasting (4, 12-17). The results of present study did not show increased risk of PCOS in breast cancer patients. In an English cohort study, women with PCOS had no significantly increased risk of mortality or morbidity from breast cancer (18). In a US cohort study on 34 835 women, of whom 833 developed a breast cancer during the follow-up, subjects with PCOS were not more likely to have a breast cancer, and adjustment for age at menarche, age at menopause, parity, oral contraceptive use, BMI, waist/hip ratio and family history of breast cancer lowered the RR to 1 (4). Recently, a pentanucleotide repeat [(TAAAA)n] polymorphism in the promoter region of the CYP11A gene, that encodes the cholesterol sidechain cleavage enzyme, has been found to be associated with the risk of PCOS (19-22). Zheng et al investigated the association of this polymorphism with breast cancer risk in a Chinese population-based case-control study including 1015 breast cancers and 1082 community controls pentanucleotide (23).Therefore, а repeat polymorphism in the CYP11A gene could represent a linkage between PCOS and breast carcinogenesis. Gammon and Thompson, 1991 reported a reduced risk of breast cancer in women with PCOS finding an odds ratio of 0.52 (95% CI 0.32-0.87), this study however is difficult to interpret as the prevalence of PCOS as identified by a self-assessed questionnaire (13). Conversely, the study of Atiomo et al detected a statistically significant positive family history of breast cancer among women with PCOS (16). In an English study designed to test whether cardiovascular mortality was increased in 786 women with PCOS followed for an average of 30 years, the standardized mortality ratio (SMR) for all causes was 0.90, based on 59 deaths, the SRM for circulatory disease was 0.83, based on 15 deaths, and the SRM for breast cancer was 1.48, based on 13 deaths (17). In conclusion, present study did not find significant relation between PCOS and risk of breast cancer. However, most of previous study suggested the relation between PCOS and risk of breast cancer in postmenopausal women by adjusting age. This statement could be confirmed by the other study for the frequency of the PCOS in postmenopausal women with breast cancer. The

age of menarche, the age at the first pregnancy and the duration of breast feeding should be considered in future study.

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References

- 1. Legro RS. Polycystic ovary syndrome and cardiovascular disease: a premature association? *Endocr Rev* 2003; 24: 302-312.
- Chang RJ. A practical approach to the diagnosis of polycystic ovary syndrome. *Am J Obstet Gynecol* 2004; 191: 713- 717.
- Spritzer PM, Morsch DM, Wiltgen D. Polycystic ovary syndrome associated neoplasms. Arq Bras Endocrinol Metabol 2005; 49: 805-810.
- 4. Anderson KE, Sellers TA, Chen PL, Rich SS, Hong CP, Folsom AR. Association of Stein-Leventhal syndrome with the incidence of postmenopausal breast carcinoma in a large prospective study of women in Iowa. *Cancer* 1997; 79: 494-499.
- 5. Helewa M, Levesque P, Provencher D, Lea RH, Rosolowich V, Shapiro HM. Breast Disease Committee and Executive Committee and Council, Society of Obstetricians and Gynaecologists of Canada. Breast cancer, pregnancy, and breastfeeding. *J Obstet Gynaecol Can* 2002; 24:164–180.
- Ross RK. Breast cancer: epidemiology, pathology, and natural history. In: Genazzani AR, editor. Hormone replacement therapy and cancer. The current status of research and practice. New York: Parthenon Publishing; 2002; 31–37.
- 7. Secreto G, Zumoff B. Abnormal production of androgens in women with breast cancer. *Anticancer Res* 1994; 14: 2113–2117.
- Kaaks R. Nutrition, hormones, and breast cancer: Is insulin the missing link? *Cancer Causes Control* 1996;7: 605–625
- 9. Daniilidis A, Dinas K. Long term health consequences of polycystic ovarian syndrome: a review analysis. *Hippokratia* 2009; 13:90-92.
- Moini A, Eslami B. Familial associations between polycystic ovarian syndrome and common diseases. J Assist Reprod Genet 2009; 26: 123-127.
- Thiboutot DM, Harper JC, O'Connell K, Rich P, Sondheimer SJ. Improving outcomes through collaboration. *Cutis* 2008; 81 (Suppl. 1): 26-31
- Chittenden BG, Fullerton G, Maheshwari A, Bhattacharya S. Polycystic ovary syndrome and the risk of gynaecological cancer: a systematic review. *Reprod Biomed Online* 2009;19: 398-405.
- 13. Gammon MD, Thompson WD. Polycystic ovaries and the risk of breast cancer. *Am J Epidemiol* 1991; 134: 818- 824.
- 14. Reshef E, Lei ZM, Rao CV, Pridham DD, Chegini N, Luborsky JL. The presence of gonadotropin receptors in nonpregnant human uterus, human placenta, fetal

membranes, and decidua. J Clin Endocrinol Metab 1990; 70: 421-430.

- 15. Balen A. Polycystic ovary syndrome and cancer. *Hum Reprod Update* 2001; 7: 522–525.
- Atiomo WU, El-Mahdi E, Hardiman P. Familial associations in women with polycystic ovary syndrome. *Fertil Steril* 2003; 80: 143–145.
- Pierpoint T, McKeigue PM, Isaacs AJ, Wild SH, Jacobs HS. Mortality of women with polycystic ovary syndrome at longterm follow-up. *J Clin Epidemiol* 1998; 51: 581– 586.
- Wild S, Pierpoint T, Jacobs H, McKeigue P. Long-term consequences of polycystic ovary syndrome: results of a 31 year follow-up study. *Hum Fertil* 2000; 3: 101–105.
- Gharani N, Waterworth DM, Batty S, White D, Gilling-Smith C, Conway GS, et al. Association of the steroid synthesis gene CYP11a with polycystic ovary syndrome and hyperandrogenism. *Hum Mol Genet* 1997; 6: 397– 402.

- 20. Diamanti-Kandarakis E, Bartzis MI, Bergiele AT, Tsianateli TC, Kouli CR. Microsatellite polymorphism (tttta) (n) at 7528 base pairs of gene CYP11aa influences hyperandrogenemia in patients with polycystic ovary syndrome. *Fertil Steril* 2000; 73: 735–741.
- Franks S, Gharani N, McCarthy M. Candidate genes in polycystic ovary syndrome. *Hum Reprod Update* 2001; 7: 405–410.
- 22. Daneshmand S, Weitsman SR, Navab A, Jakimiuk AJ, Magoffin DA. Overexpression of theca-cell messenger RNA in polycystic ovary syndrome does not correlate with polymorphisms in the cholesterol side-chain cleavage and 17ahydroxylase/ C (17-20) lyase promoters. *Fertil Steril* 2002; 77: 274–280.
- 23. Zheng W, Gao YT, Shu XO, Wen W, Cai Q, Dai Q, et al. Population-based case–control study of CYP11A gene polymorphism and breast cancer risk. *Cancer Epidemiol Biomarkers Prev* 2004; 13: 709–714.