RESEARCH ARTICLE

Associations of Serum Isoflavone, Adiponectin and Insulin Levels with Risk for Epithelial Ovarian Cancer: Results of a Case-control Study

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Abstract

Background: The aim of this study was to examine the association of serum isoflavones, adiponectin, and insulin levels with ovarian cancer risk. Materials and Methods: We gathered cases with histologically confirmed epithelial ovarian cancer at Sapporo Medical University Hospital from October 2010 to September 2012. Potential controls were recruited from female inpatients without any history of cancer or diabetes mellitus in different wards of the same hospital over the same period of time. Serum isoflavones, adiponectin, and insulin levels were measured in order to estimate associations with ovarian cancer risk in a case-control study. Data from 71 cases and 80 controls were analyzed with a logistic regression model adjusting for known risk factors. Results: A significant reduction in ovarian cancer risk was observed for the high tertile of serum daidzein level versus the low (P \text{ trend} < 0.001). A significant reduction in ovarian cancer risk was also observed for the high tertile of serum glycitein level versus the low (P \text{ trend} = 0.005). Furthermore, a significant reduction in ovarian cancer risk was observed for the high tertile of serum adiponectin level versus the low (P \text{ trend} = 0.004). Conversely, serum insulin level showed significantly elevated risk for ovarian cancer with the high tertile versus the low (P \text{ trend} < 0.001). Conclusions: Decreased serum isoflavones levels, such as those for daidzein and glycitein, decreased serum adiponectin levels, and increased serum insulin levels could be shown to be associated with elevated risk of ovarian cancer.

Keywords: Case-control study - isoflavones - insulin - adiponectin

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Materials and Methods

We collected cases with histologically confirmed epithelial ovarian cancer at Sapporo Medical University Hospital from October 2010 to September 2012. Potential controls were gathered from female inpatients in different wards of the same hospital over the same period. Patients with cancer at other sites or diabetes mellitus were excluded from the control group. Most of the control subjects were primarily diagnosed with having hypertension, cardiac insufficiency, atrial or ventricular tachycardia, or other circulatory diseases.

Information on demographic, anthropometric, and reproductive variables was obtained using a self-administrated structured questionnaire. Clinical information, including histologic types of ovarian cancer, such as serous, endometrioid, clear cell, and mucinous carcinomas, was collected following hospital protocol. Comparison of these serum levels was also performed according to histologic types of ovarian cancer.

All blood samples were obtained after fasting early in the morning. All serum isoflavones such as genistein, daidzein, glycitein, and equol, were measured using a liquid chromatography mass spectrograph; LC-MS/MS. Insulin and high molecular weight (HMW) adiponectin levels were measured using Chemiluminescent Enzymoassays (CLEIA) with sensitivities of 0.31µIU/mL and 0.20 µg/mL, respectively. All analyses were performed at a special reference laboratory (SRL Co., Tokyo Japan).

Descriptive statistics including mean (SD; standard deviation) and median values (interquartile ranges) for continuous or ordinal variables as well as proportions for categorical variables were computed for both the case and control groups. Comparisons were made between the case and control groups using a chi-square test for categorical variables, and with the Mann-Whitney test for continuous or ordinal variables. A Bonferroni correction was used for the multiple comparison.

Odds ratios (OR) and 95% confidence intervals (95% CI) for ovarian cancer risk were estimated according to the tertile of genistein, daidzein, glycitein, adiponectin and insulin levels through analysis of the unconditional logistic regression model, adjusting for potentially confounding variables such as age, body mass index (BMI), oral contraceptive usage, fertility drugs usage and amount of parity. Significance level was set at 5%. We conducted all analyses using SPSS version 18 (SPSS Inc., Chicago, IL, U.S.A.). The study was approved by the Institutional Review Board at Sapporo Medical University, and written informed consent was obtained from every participant.

Results

As shown in Table 1, the proportion of the cases with histories of ovarian cancer in a mother or sister was significantly higher than that of the controls (p=0.047). However, age, BMI, frequency of oral contraceptive usage and fertility drug usage, and the amount of parity were not different between the cases and the controls. Serum levels of daidzein (p<0.001) and glycitein (p=0.010) were significantly lower in the cases than in the controls. Although it was not significant, serum adiponectin level was lower in the cases than in the controls (p=0.090). Serum insulin levels were significantly higher in the cases than in the controls (p=0.001).

Table 2 showed ORs and 95% CIs of epithelial ovarian cancer risk

| Items                                           | Case (N=71) | Control (N=80) | p value |
|-------------------------------------------------|------------|----------------|---------|
| Mean age (SD) in years                          | 57.4 (11.7) | 56.3 (14.5)    | 0.807   |
| Mean body mass index (SD) in kg/m²              | 21.8 (3.3)  | 21.4 (4.0)     | 0.198   |
| Oral contraceptive use, no (%)                 | 66 (93.0)   | 68 (85.0)      | 0.123   |
| Never                                           | 5 (7.0)     | 12 (15.0)      |         |
| Has used                                        |             |                |         |
| Fertility drugs, no (%)                        | 67 (94.4)   | 76 (85.0)      | 0.862   |
| Never                                           | 4 (5.6)     | 4 (5.0)        |         |
| Has used                                        |             |                |         |
| Parity (%)                                      |             |                | 0.506   |
| 0                                               | 21 (29.6)   | 20 (25.0)      |         |
| 1-2                                             | 36 (50.5)   | 48 (60.0)      |         |
| >2                                              | 14 (19.7)   | 12 (15.0)      |         |
| History of ovarian cancer in a mother or sister (%)| 67(94.4)   | 80(100.0)      | 0.047   |
| Yes                                             | 4(5.6)      | 0(0.0)         |         |

Table 1. Baseline Characteristics and Serum Levels of Isoflavones, Adiponectin, Insulin in the Epithelial Ovarian Cancer Cases and the Controls
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Genistein: low (<24.1 ng/mL), middle (24.1-106.4 ng/mL), high (≥106.5 ng/mL); Daidzein: low (<10.9 ng/mL), middle (10.9-43.8 ng/mL), high (≥43.9 ng/mL); Glycitein: low (<0.5 ng/mL), middle (0.5-2.2 ng/mL), high (≥2.3 ng/mL); Equol; low (<0.6 ng/mL), high (≥0.6 ng/mL); Adiponectin: low (<4.7 µg/mL), middle (4.7-9.6 µg/mL), high (≥9.7 µg/mL); Insulin: low (<4.1 µIU/mL), middle (4.1-8.9 µIU/mL), high (≥9 µIU/mL); Model 1: adjusted for age, BMI, oral contraceptive use, use of fertility drugs, and parity; Model 2: adjusted for variables in Model 1 plus daidzein; Model 3: adjusted for variables in Model 1 plus genistein; Model 4: adjusted for variables in Model 1 plus daidzein and insulin

Table 2. Adjusted Odds Ratio (OR) and 95% Confidence Interval (95% CI) for the Epithelial Ovarian Cancer Cases Compared with the Controls

Table 3. Comparison of Age, Serum Isoflavones, Daidzein and Insulin Levels among the Cases with Histologic Types of Ovarian Cancer and the Controls

Values are expressed as means (SD) or medians [interquartile range]. *p<0.05 after Bonferroni correction; P1 and P2 presented comparisons between the cases with clear cell carcinoma and the controls; P3 presented comparison between the cases with mucinous carcinoma and the controls

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as well as mucinous carcinoma were significantly higher than those in the controls (both, p<0.01 after Bonferroni correction). Other comparisons did not reveal differences among the cases with different histological types, or differences between the cases and the controls.

Discussion
Consistent with previous studies showing the preventive effect of isoflavones against ovarian cancer (Lee et al., 2014; Rossi et al., 2008; Chang et al., 2007; Zhang et al., 2004), our study has indicated that low serum daidzein and glycitein levels were associated with increased risk for ovarian cancer. Especially, we observed significantly decreased daidzein levels in ovarian cancer cases with clear cell carcinoma as the specific histologic type. As a biological explanation for the protective effect of isoflavones against ovarian cancer, isoflavones have been shown to induce apoptosis and inhibit the growth and proliferation of ovarian cancer cells (Gossner, 2007). Another plausible mechanism is through antiangiostatic effects on ovarian cancer cell lines, since it is well known that ovarian cancer is an estrogen-dependent cancer.

Our study has also shown that low serum adiponectin levels are associated with an elevated risk of ovarian cancer. Although epidemiological studies on the association between serum adiponectin levels and ovarian cancer risk have been scarcely reported, as far as our knowledge, adipokines signaling pathways were suggested to be involved in ovarian follicle development and ovarian cancer (Dupont, et al., 2012). Furthermore, low adiponectin antagonizes many effects of TNF-α, and have been recognized as a key regulator of insulin sensitivity and tissue inflammation (Lihn et al., 2005, Fantuzzi, 2005). TNF-α, a major inflammatory cytokine, is abundant in the ovarian cancer microenvironment, and chronic inflammation may be implicated in the pathophysiology of ovarian cancer.

In this study, the high tertile of insulin levels had an approximately fivefold risk of ovarian cancer through mechanisms other than obesity, because a significant relationship was still observed even after adjustment of BMI. IGF-1 has been implicated in the regulation of ovarian epithelial cell proliferation (Lukanova, et al., 2002), and this may provide an etiological explanation for increased risk of insulin for ovarian cancer risk. As another possible mechanism of IGF-1 involvement in ovarian carcinogenesis, an experimental study has shown that the malignant transformation of ovarian cells can be induced by overexpression of IGF-1 receptors (Coppola et al., 1999).

We conducted histologic comparisons of ovarian carcinoma in terms of serum levels of measured substances, because of different risk factors (Risch et al., 1996), and because different morphologic properties (Soslow, 2008) have been shown among various types of histopathology in ovarian cancer. Significantly lower serum daidzein levels and significantly higher serum insulin levels were observed in the cases with clear cell carcinoma. Furthermore, significantly higher serum insulin levels were noted in cases with mucinous cell carcinoma. However, the sample size of each histopathology was not enough to conclude these characteristics. Further study is necessary to confirm them.

As a major limitation of this study, the study design was a hospital-based case-control study, where controls were recruited only from inpatients who might have characteristics different from the general population. This may have resulted in a biased estimate of the association between exposure and diseases. Secondly, because the sample size was limited, type II errors might exist in the study. In addition, because only Japanese women were studied, our results may not be applicable to other racial/ethnic groups.

In conclusion, decreased serum isoflavones levels, such as those for daidzein and glycitein, decreased serum adiponectin levels, and increased serum insulin levels are shown to be associated with an elevated risk of ovarian cancer.

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