Influence of cigarette smoking on hormone and lipid metabolism in women in late reproductive stage

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Background: The aim of the study was to analyze lipid and hormone metabolism, body mass index (BMI), and age parameters in late reproductive stage women in relation to cigarette smoking.

Methods: The study enrolled 345 healthy late reproductive stage women living in Poland; 13.33% were smokers. The first part of the study assessed lipid metabolism (total cholesterol, high-density lipoprotein [HDL], low-density lipoprotein [LDL], and triglycerides) and hormone metabolism (estradiol [E2], follicle-stimulating hormone [FSH], and anti-Müllerian hormone [AMH] levels) in women in the early phase of the follicular menstrual cycle. The second part of the study was carried out using the diagnostic survey method, with a standardized questionnaire (Primary Care Evaluation of Mental Disorders [PRIME-MD]) and the authors’ own research tools.

Results: The women were aged 42.3±4.5 years (mean ± SD). The BMI (24.8±4.04 kg/m²) did not differ significantly between the groups. The women who smoked cigarettes had a statistically significantly lower level of HDL as well as higher LDL and triglyceride levels (p<0.05). Differences were also shown in hormone levels: non-smoking participants had statistically significantly higher levels of E2 and FSH (p<0.05). In the group of non-smoking women, age was a predictor exerting a significant positive impact on the levels of total cholesterol, LDL, triglycerides, and AMH (p<0.05). BMI contributed to a decline in HDL and triglyceride levels. In the group of smoking women, age significantly positively influenced the level of E2, and negatively influenced AMH. BMI was associated with a significant decrease in the HDL level.

Conclusion: Smoking cigarettes affects the physical health of women in late reproductive stage through negative influences on lipid and hormone metabolism, among other factors. Age is an unmodifiable factor adversely affecting both lipids and hormones. Higher BMI has a negative influence on lipid metabolism in both groups of women in this study.

Keywords: smoking, cholesterol profile, gonadal steroid hormones

Introduction

Smoking cigarettes is a well-documented risk factor for the development of cardiovascular disease, atherosclerosis, and type 2 diabetes. Several studies have looked at the relationships between cigarette smoking and triglyceride concentration in blood serum, fasting blood glucose, and high-density lipoprotein (HDL) cholesterol levels.¹⁻⁵ The outcomes did not always demonstrate the expected relationship. The study conducted by Freeman et al on a group of men proved only a slight influence of smoking on the level of triglycerides.² Other reports did not show statistically significant differences...
in fasting glycemia level and insulin resistance between smokers and non-smokers. An important factor influencing these relationships may be body mass, as an increase in body mass is strongly associated with an increase in triglyceride levels in serum and a decrease in HDL concentration. Some studies report that people who smoke tend to maintain a lower body weight than non-smokers.

Moreover, epidemiological studies reveal a negative impact of cigarette smoking on ovarian function and fertility, as well as faster onset of menopause. The effect of environmental factors, such as cigarette smoking and obesity, on ovarian reserve may be important for determining the time of menopause. There is more and more scientific evidence of the usefulness of anti-Müllerian hormone (AMH) testing, together with follicle-stimulating hormone (FSH), in measuring ovarian reserve and evaluating the impact of environmental factors on fertility.

AMH belongs to the transforming growth factor-β family. Initiation of AMH expression in granular layer cells of Graafian follicles takes place during folliculogenesis. AMH expression is higher in precaervenous granular cells and in small cavernous follicles, and gradually undergoes changes in further stages of follicular development. Studies conducted among women have mainly focused on analyzing the role of AMH in serum as a determinant of ovarian function. van Rooij et al found a strong relationship between the number of cavernous follicles, evaluated with the use of ultrasonography, which reflects the size of the primary follicle pool, and the level of AMH in serum. This suggests that AMH is an excellent marker of ovarian reserve. Studies by Biggs et al revealed that this hormone has strong predictive properties of ovarian reserve, after taking into account age as well as FSH, luteinizing hormone (LH), and estradiol (E2) hormone levels.

The process of ovarian aging is also characterized by a progressive increase in FSH concentration in serum. This increase is associated with progressive loss of ovarian function. In murine studies, it was noted that a lack of AMH causes follicles to be recruited at a significantly faster pace. Moreover, it was shown that follicles display an increased sensitivity to FSH.

The 2001 Stages of Reproductive Aging Workshop (STRAW) standardized the classification of menopause status. A decade later, a new, updated STRAW+10 version was developed. The study presented here included women who, according to this classification, were in the late reproductive stage (the –3 stage), ie, the stage typified by the onset of a decline in fecundability and first changes in the menstrual cycle. In accordance with the STRAW+10 criteria, this phase is divided into two subphases: –3b and –3a. The length of the menstrual cycle in the –3b subphase remains unchanged. What can be observed, however, is that the level of AMH and the antral follicle count decrease. The –3a subphase is characterized by a shortening of the menstrual cycle. Moreover, in the early follicular phase, the level of FSH increases, while the levels of other biomarkers of ovarian aging decrease. The level of FSH >30 mIU/mL is regarded as critical and is marked by a lack of ovarian follicles in the ovaries. In this period, the level of E2 decreases dramatically (from 40–400 pg/mL before menopause, depending on the phase of the menstrual cycle, to 20–30 pg/mL after menopause).

The aim of this study was to analyze the influence of smoking on:

1) the lipid and hormonal profiles of women in the late reproductive stage;
2) the relationship between hormonal and lipid profiles in the study sample;
3) parameters of hormonal and lipid metabolism depending on the participants’ age and body mass index (BMI).

Materials and methods

The study enrolled healthy women in the late reproductive stage, living in north-west Poland. All participants came from the general population of the West Pomerania Province (Poland). A quota sampling method was applied. Recruitment was performed based on advertisements in public places and local newspapers. All the women were classified into the late reproductive stage group based on the results of their FSH and AMH levels according to STRAW+10 criteria. The youngest participant was 35 years old, and the oldest one was 53 years old. Women with endocrine disorders, thyroid diseases, diabetes, neoplastic disease, abnormal smear test results, or mental disorders were excluded. Women with Axis I mental disorders according to the ICD-10 classification were excluded from the analysis by means of the Primary Care Evaluation of Mental Disorders (PRIME-MD) questionnaire and a psychiatric examination. The study involved 345 women; 26 women were excluded, either because they did not meet the inclusion criteria or because of incomplete data.

All subjects gave their written informed consent for inclusion in the research. The study was conducted in accordance with the Declaration of Helsinki, and the protocol was approved by the Bioethical Commission of the Pomeranian Medical University in Szczecin, Poland (permission number KB-0012/12/12). Those responsible for the whole research...
process were qualified and properly trained academic workers (Masters of Nursing) of the Pomeranian Medical University in Szczecin.

Assessments
The first part of the study comprised laboratories tests including lipid metabolism (total cholesterol, HDL cholesterol, low-density lipoprotein [LDL] cholesterol, and triglycerides) and hormone metabolism (E2, FSH, and AMH levels).

Women in the early phase of the follicular menstrual cycle had their venous blood sample taken using the closed blood collection system BD Vacutainer®, following the existing guidelines. Blood was collected in the treatment room and delivered to the laboratory in accordance with the relevant rules and procedures.

The levels of AMH, FSH, E2, total cholesterol, HDL cholesterol, LDL cholesterol, and triglycerides were measured in a certified laboratory (ISO 9001:2008 certificate).

The following norms of “Laboratoria Medis” were followed: total cholesterol 115–190 mg/dL, HDL cholesterol ≥45 mg/dL, LDL cholesterol <115 mg/dL, and triglycerides ≤150 mg/dL. For the follicular phase, the norms were E2 12.5–166 pg/mL and FSH 3.5–12.5 mIU/mL.

In the second part of the study, we carried out a survey with a standardized questionnaire (PRIME-MD) and our own research tools, concerning sociodemographic data and information about smoking. Based on the ICD-10 criteria, we established whether the women were addicted to nicotine. The group of smokers comprised only those women who had at least three of the following symptoms during the previous year: a strong need for nicotine; an increasing problem with control of smoking-related behaviors; withdrawal syndrome while making attempts to limit or give up smoking; a growing need to smoke more cigarettes to achieve the same effect that was previously attainable with a lower number; neglecting interests and activities for smoking; and smoking despite the awareness of its numerous negative consequences. Other women were included in the non-smoking group after confirming that they did not smoke at all.

In addition, the patients were weighed and measured in order to calculate their BMI values according to the formula BMI = weight (kg)/height² (m²).

Statistical analysis
Statistical analysis was conducted with the use of the PASW Statistics 18 software. The lack of normal distribution of variables analyzed was confirmed with the Shapiro–Wilk test (p<0.05). The Mann–Whitney U-test was applied to verify the null hypothesis regarding the independence of the analyzed variables. Multiple regression analysis was applied to identify predictors having a significant impact on response (or dependent) variables. In the process of statistical inference, statistically significant values were taken as p≤0.05.

Results
There were 345 women in the study. Of those, 13.3% were smokers and 86.7% were non-smokers. They were aged 42.3±4.5 years (mean ± SD), and the median age of smoking and non-smoking women was similar. Three-quarters of the participants (75.1%) had a university education and 22.6% had a secondary education. The majority of the women (72.5%) lived in cities with a population greater than 100,000 residents. Most of the participants (75.1%) lived with partners and the vast majority (96.5%) were working.

The BMI of the studied women was 24.8±4.04 kg/m² (mean ± SD), which is within the normal range. The BMI was 24.6±3.7 kg/m² in the non-smoking group and 25.4±5.4 kg/m² in the smoking group (non-significant difference).

The mean values of lipid metabolism indicators for the studied women were within the normal range for triglycerides (93.1±6±50.00 mg/dL) and HDL cholesterol (55.40±13.30 mg/dL) and slightly above the existing norms for total cholesterol (193.05±33.93 mg/dL) and LDL cholesterol (119.04±30.16 mg/dL).

A comparative analysis of lipid metabolism in the smoking and non-smoking women showed statistically significant differences in mean concentrations of HDL, LDL, and triglycerides (Table 1). The women who smoked had lower HDL, and higher LDL and triglyceride levels compared with non-smoking women in the late reproductive stage. Statistically significant differences were also shown in hormone levels, with non-smoking participants having higher levels of E2 (85.9 vs 64.4 pg/mL) and FSH (6.2 vs 7.4 mIU/mL) compared with smokers (Table 1).

Table 2 presents the results of multiple regression analysis of lipid and hormone levels in both smokers and non-smokers. Estimated models described the actual data in an unsatisfactory way (R<0.5); the differences were not statistically significant (p>0.05).

Next, we analyzed how age and BMI affected lipid and hormone levels in the groups of smokers and non-smokers. In the case of non-smoking women, age was a significant positive contributor to the levels of total cholesterol, LDL, triglycerides, and AMH (p<0.05). BMI was associated with a decrease in HDL and triglyceride levels. In the group of
smoking women, age had a significant, positive impact on the level of E2, and a negative impact on AMH. BMI was associated with a significant decline in the HDL level (Table 3).

Discussion

The outcomes of the research described in this article show adverse effects of smoking cigarettes on lipid metabolism (lower HDL, and higher LDL and triglyceride levels in smokers) and hormonal functioning (lower E2 and higher FSH levels). Factors such as age and a higher BMI make this relationship even stronger.

Dyslipidemia is considered one of the most important modifiable factors in the development of type 2 diabetes,20,21 atherosclerosis,22,23 cardiovascular disease,24,25 and stroke.26,27 A well-documented effect of smoking is hypercholesterolemia, which is associated with a higher risk of cardiovascular disease. This factor is believed to influence the increase in total cholesterol level, LDL, and triglycerides, simultaneously inducing the decrease in HDL.25 In our study, significant differences between groups of smoking and non-smoking women were shown. Significantly higher concentrations of LDL cholesterol and triglycerides were observed in smoking compared with non-smoking women, while the level of HDL cholesterol was significantly lower in smokers than in non-smokers. These results show that the risk of developing the above-mentioned diseases is considerably higher in smoking women than in their non-smoking counterparts.

A study conducted on 11,559 participants from the Republic of Korea, whose smoking status was verified with a nicotine urine test, revealed that the risk of developing metabolic syndrome was significantly higher among smokers.29 The women who smoked had a significantly higher risk of abdominal obesity and a higher level of triglycerides compared with non-smokers.29 These findings correspond with our results, showing that smokers significantly more often had symptoms typical of metabolic syndrome, such as lower HDL and higher LDL levels.

The outcomes of a lipid profile assessment performed in women in 1983 suggested a negative influence of smoking on lipid metabolism.28 Mean concentrations of triglycerides and total cholesterol in the smoking and non-smoking groups in that study were lower than those obtained in the current study in 2016 (total cholesterol – 1983: smoking

Table 1 Comparative analysis of lipid metabolism and hormones in non-smoking and smoking women in the late reproductive stage

<table>
<thead>
<tr>
<th>Variable</th>
<th>Non-smoking</th>
<th>Smoking</th>
<th>Z</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N=345</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>n=299 (86.67%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Median (IQR)</td>
<td>Min–Max</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lipid profile</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total cholesterol (mg/dL)</td>
<td>191 (44)</td>
<td>107–295</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HDL cholesterol (mg/dL)</td>
<td>55 (16)</td>
<td>21–109</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LDL cholesterol (mg/dL)</td>
<td>116 (38)</td>
<td>41–221</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Triglycerides (mg/dL)</td>
<td>79 (46)</td>
<td>35–327</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hormone profile</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>E2 (pg/mL)</td>
<td>85.9 (99.5)</td>
<td>4.3–841</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FSH (mIU/mL)</td>
<td>6.2 (3.4)</td>
<td>2.3–497.5</td>
<td></td>
<td></td>
</tr>
<tr>
<td>AMH (ng/mL)</td>
<td>1.33 (2.37)</td>
<td>0.15–11.59</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: Bold values denote statistical significance (p<0.05).

Abbreviations: IQR, interquartile range; Min, minimum value; Max, maximum value; HDL, high-density lipoprotein; LDL, low-density lipoprotein; E2, estradiol; FSH, follicle-stimulating hormone; AMH, anti-Müllerian hormone.

Table 2 Multiple regression analysis of lipid metabolism and hormones in smoking and non-smoking women in the late reproductive stage

<table>
<thead>
<tr>
<th>Smoking</th>
<th>Total cholesterol (mg/dL)</th>
<th>HDL cholesterol (mg/dL)</th>
<th>LDL cholesterol (mg/dL)</th>
<th>Triglycerides (mg/dL)</th>
<th>R/p</th>
</tr>
</thead>
<tbody>
<tr>
<td>AMH (ng/mL)</td>
<td>Yes</td>
<td>–</td>
<td>–0.015</td>
<td>–0.002</td>
<td>0.001</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>0.000</td>
<td>0.003</td>
<td>–</td>
<td>–0.001</td>
</tr>
<tr>
<td>E2 (pg/mL)</td>
<td>Yes</td>
<td>–</td>
<td>0.082</td>
<td>0.232</td>
<td>–0.036</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>0.112</td>
<td>0.583</td>
<td>–</td>
<td>0.172</td>
</tr>
<tr>
<td>FSH (mIU/mL)</td>
<td>Yes</td>
<td>–</td>
<td>–0.068</td>
<td>0.097</td>
<td>–0.039</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>0.004</td>
<td>0.000</td>
<td>–</td>
<td>–0.026</td>
</tr>
</tbody>
</table>

Abbreviations: HDL, high-density lipoprotein; LDL, low-density lipoprotein; AMH, anti-Müllerian hormone; E2, estradiol; FSH, follicle-stimulating hormone; R, coefficient of determination; p, level of significance.
Table 3 Multiple regression analysis of hormone and lipid metabolism determinants with age and BMI in the groups of smoking and non-smoking women in the late reproductive stage

<table>
<thead>
<tr>
<th>Smoking and BMI</th>
<th>Age</th>
<th>BMI</th>
<th>R/p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total cholesterol</td>
<td>Yes</td>
<td>0.247 ± 0.040</td>
<td>0.067</td>
</tr>
<tr>
<td>No</td>
<td>0.154 ± 0.030</td>
<td>0.021</td>
<td></td>
</tr>
<tr>
<td>HDL cholesterol</td>
<td>Yes</td>
<td>0.193 ± 0.014</td>
<td>0.014</td>
</tr>
<tr>
<td>No</td>
<td>-0.066 ± 0.000</td>
<td>0.000</td>
<td></td>
</tr>
<tr>
<td>LDL cholesterol</td>
<td>Yes</td>
<td>0.197 ± 0.255</td>
<td>0.000</td>
</tr>
<tr>
<td>No</td>
<td>0.169 ± 0.002</td>
<td>0.002</td>
<td></td>
</tr>
<tr>
<td>Triglycerides</td>
<td>Yes</td>
<td>-0.018 ± 0.671</td>
<td>0.000</td>
</tr>
<tr>
<td>No</td>
<td>0.108 ± 0.000</td>
<td>0.000</td>
<td></td>
</tr>
<tr>
<td>AMH (ng/mL)</td>
<td>Yes</td>
<td>-0.402 ± 0.018</td>
<td>0.000</td>
</tr>
<tr>
<td>No</td>
<td>-0.347 ± 0.000</td>
<td>0.000</td>
<td></td>
</tr>
<tr>
<td>E2 (pg/mL)</td>
<td>Yes</td>
<td>0.371 ± 0.076</td>
<td>0.007</td>
</tr>
<tr>
<td>No</td>
<td>0.042 ± 0.369</td>
<td>0.000</td>
<td></td>
</tr>
<tr>
<td>FSH (mIU/mL)</td>
<td>Yes</td>
<td>0.243 ± 0.083</td>
<td>0.000</td>
</tr>
<tr>
<td>No</td>
<td>0.069 ± 0.245</td>
<td>0.000</td>
<td></td>
</tr>
</tbody>
</table>

Notes: *p<0.1; **p<0.05; ***p<0.001.
Abbreviations: BMI, body mass index; HDL, high-density lipoprotein; LDL, low-density lipoprotein; AMH, anti-Müllerian hormone; E2, estradiol; FSH, follicle-stimulating hormone; R, coefficient of determination; p, level of significance.

Research conducted among 141 infertile women in Chile proved that cigarette smoking is a risk factor leading to decreased AMH concentrations in blood plasma. After ruling out the influence of other factors, it was shown that smoking caused a decrease of AMH by 2.29 ng/mL, irrespective of the number of cigarettes smoked in the week preceding the test. However, a relationship between smoking and FSH and estradiol levels on the third day of the menstrual cycle was not proven. In contrast, the level of AMH in our study was similar in both groups, but a significant relationship between smoker status and estradiol and FSH levels in the initial follicular phase was demonstrated. Divergence between these results was probably associated with the fact that the Chilean study included infertile women with hormonal disorders, whereas the present study involved healthy women in the late reproductive stage (FSH: smokers 14.86 mIU/mL vs 7.4 mIU/mL and non-smokers 14.03 mIU/mL vs 6.2 mIU/mL, in the Chilean and the present study, respectively; E2: smokers 28.89 pg/mL vs 64.4 pg/mL and non-smokers 30.67 pg/mL vs 85.9 pg/mL). Comparison between these two studies reveals the influence of smoking on female hormone metabolism, depending on whether hormonal function was normal or disordered.

A study among Chinese women showed that BMI has a significant impact on the occurrence of dyslipidemia. It was observed that people with dyslipidemia had a BMI of 25.1 ± 3.3 kg/m² (mean ± SD), while for people with no lipid metabolism abnormalities, the BMI was 23.2 ± 3.1 kg/m². This relationship was particularly apparent for people with hypertriglyceridemia, in whom the BMI was 25.5 ± 3.4 kg/m². Moreover, results of multivariable logistic regression revealed that the level of education, presence of obesity, and cigarette smoking significantly increased the risk of dyslipidemia, while regular physical activity turned out to be an independent protective factor. Our study also demonstrated a significant negative impact of increased BMI.
on lipid metabolism, which especially refers to the levels of HDL and triglycerides. This effect was observed in both smoking and non-smoking women.

In the study by Qi et al, the age-adjusted prevalence of dyslipidemia was higher in women (37.6%) than in men (34.4%). This result may be due to differences in the frequency of overweight, obesity, and central obesity, which affect women much more often than men in the Chinese population. In the current study, the level of cholesterol in smoking and non-smoking women in the late reproductive stage increased significantly with age. Confirmation of these findings requires further analyses and a larger number of studied patients.

Limitations
Our study was not free from limitations. The most important of these were the small number of women categorized as smokers, and the fact that women were included in the non-smoking group exclusively on the basis of the ICD-10 dependence syndrome. It would be interesting to conduct more extensive research, taking into consideration duration and intensity of smoking, as well as exposure to passive smoking. Despite these limitations, research outcomes indicate the presence of mutual relationships between some hormone and lipid metabolism markers and cigarette smoking in the group of women in the late reproductive stage.

Conclusion
In relation to our earlier aims we conclude that 1) cigarette smoking in women in the late reproductive stage has a negative influence on lipid and hormone metabolism; 2) age affects both lipid and hormone metabolism in women in the late reproductive stage. This relationship can be more clearly seen in non-smoking women; and 3) higher BMI values have a negative influence on lipid metabolism in both smoking and non-smoking women.

Disclosure
The authors declare no conflicts of interest in this work.

References


