Prevalence of cardiometabolic risk factors in patients with hypertension and subclinical hypothyroidism

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It is known that the presence of overt hypothyroidism carries additional risks of developing cardiovascular diseases due to impaired lipid and carbohydrate metabolism. But whether subclinical hypothyroidism (SH) has the same negative impact is still controversial. The assessment of its role is especially important in patients with existing arterial hypertension (AH) in the early stages of the disease in order to prevent future complications.

The aim of this work is to identify and assess the prevalence of early cardiometabolic risk factors in patients with AH combined with SH.

Materials and methods. 66 patients (55.4 % women) aged from 25 to 59 years with a median age of 51.1 years were included in the study during 2019–2020 years. All the patients were divided into 3 groups, randomized by age and sex: group 1 (n = 21) – volunteers without AH and SH; group 2 (n = 25) – euthyroid patients with stage 1–2 grade 1–2 AH and low-to-moderate cardiovascular risk (CVR); group 3 (n = 20) – patients with stage 1–2 grade 1–2 AH and low-to-moderate CVR in combination with SH. Blood pressure was measured, anthropometric data were assessed, glucose levels and lipid profile indicators were determined in all patients.

Results. Comparative characteristics of the groups showed a rise in the frequency of detecting increased waist circumference and the waist-to-hip ratio, obesity, metabolic syndrome, its individual components and lipid profile disorders, especially the levels of total cholesterol and high-density lipoprotein cholesterol in patients with SH even in the early stages of AH and CVR of low gradations. However, dyslipidemias in general and hypertriglyceridemia in particular were more common in euthyroid hypertensive patients compared to patients with AH and concomitant SH.

There was also a tendency towards an increase in gynoid obesity and a worsening of the lipid and carbohydrate profile disorders in SH patients in comparison to euthyroid patients with AH, although the differences were not statistically significant.

Conclusions. Evaluation of cardiometabolic risk factors revealed the increase in severity of female obesity and worsening of abnormalities in lipid and carbohydrate profiles with the SH development in patients even in the early stages of AH and low-CVR, that additionally increases the risk of cardiovascular complications.
Arterial hypertension (AH) is diagnosed in more than 40 % of the world population and this number is constantly growing, especially among the population of Central and Eastern Europe [1]. One of the negative consequences of this disease is disability. The average number of total years of life lost and disability adjusted life years (DALYs) due to disability and premature mortality caused by cardiovascular disease (CVD) was 10.000 per 100.000 Ukrainian population in 2017 [2]. The number of DALYs due to AH increased by 31 % over the period 2007 to 2017 [3]. At the same time, the prevalence of hypothyroidism in Ukraine has also been growing in recent years. Subclinical hypothyroidism (SH) is a pathological condition that precedes the development of overt (manifest) hypothyroidism. It is characterized by an increase in thyroid-stimulating hormone (TSH) levels above normal with normal laboratory values of thyroxine and triiodothyronine. The prevalence of SH is about 10 % among the population. The frequency of AH detection in patients with SH is quite high, which indicates the common links in the pathogenesis of both diseases. In the South Indian Population Study, AH was found in 24.6 % of patients with SH, and E. Plantanida et al. (2016) found that masked AH was revealed in 26.3 % of SH patients [4,5]. The negative impact of overt hypothyroidism on CVD has been proven by a large number of studies. For instance, it is known that increase in TSH is a statistically significant predictor of lipid and carbohydrate metabolism disorders, and decrease in thyroid function leads to the development of dyslipidemia, insulin resistance, obesity, metabolic syndrome (MS), hyperuricemia [6]. Lipid and carbohydrate metabolism disorders have also been found to be the most important cardiovascular risk (CVR) factors resulting in the development of age-associated diseases [7]. An assessment of subclinical thyroid dysfunction has been made possible only recently owing to new highly sensitive methods of laboratory diagnosis. Thus, current data on the impact of SH on the progression of CVD, namely AH, are limited and quite contradictory [8,9].

Diagnostic suspicion of SH in the population, especially among non-elderly patients without severe comorbid pathologies, is often absent in the routine practice of primary and secondary care physicians. Therefore, it is interesting to study the influence of SH on the CVR factors occurrence in patients with AH in the early stages of the disease. Existing studies suggest that elevated levels of TSH and low levels of free thyroxine, even within baseline ranges, are associated with a higher risk of carbohydrate metabolism disorders, the development of type 2 diabetes mellitus (DM) and an increased risk of AH[10,11]. The highest percentile of highly sensitive TSH among patients in Taiwan with a significantly increased risk of overweight, central obesity, high blood pressure, dyslipidemia, and MS is a case in point [12]. The occurrence and severity of many CVR factors, especially their combination, in patients with a combined course of AH and SH can cause more severe disorders and accelerate the progression of both diseases. Therefore, it is important to correctly assess the impact of SH on the CVR to optimize preventive measures for AH patients.

**Aim**

The aim of this study was to identify and evaluate the prevalence of early cardiometabolic risk factors in patients with AH combined with SH.
**Materials and methods**

The study included 66 patients with low-to-moderate CVR and mean age of 51.1 [41.7; 56.2] years who underwent outpatient or inpatient treatment at the GI “L. T. Malaya Therapy National Institute of the National Academy of Medical Sciences of Ukraine” (L. T. Malaya NIT NAMSU) from 2019 to 2020 and signed an informed consent to participate in the study. Among the subjects, women accounted for 55.4 % (n = 41). The research protocol was approved at a meeting of the Ethics Commission of the L. T. Malaya NIT NAMSU.

All the patients underwent clinical examination that included blood pressure (BP) and anthropometric measurements (height, weight, waist (WC) and hip circumferences (HC), body mass index (BMI), WC to HC ratio). The gynoid morphotype (low WC to HC ratio: for women <0.8; for men <0.9) in patients was evaluated as one of the protective factors against cardiometabolic risks. Lipid profile (total cholesterol (TC); triglycerides (TG); very low-density lipoprotein cholesterol (VLDL-C); high-density lipoprotein cholesterol (HDL-C); atherogenic coefficient (AC); low-density lipoprotein cholesterol (LDL-C)); carbohydrate profile (fasting glucose level; homoeostasis model assessment of insulin resistance, HOMA-IR) were also determined. The presence of MS and its components as cardiometabolic risk factors were determined according to the criteria of the International Diabetes Federation (IDF) (2006).

Statistical processing of the results was performed using the application package Statistica (GRDKR-JFFPD-B34B-3GBV9-QTTHJ), with the serial number X12-53766. Since most of the variables were not normally distributed, the results were presented as the median (Me) and interquartile ranges – 25 (Q1) and 75 (Q3) percentiles – Me (Q1; Q3). The non-parametric Mann–Whitney U test was used to compare indicators between groups. Equality of group variances was checked using Fisher’s exact (one-sided) test in four-field tables. Statistically significant differences were determined at a level of P < 0.05.

**Results**

An objective examination showed that patients in group 3 had better systolic BP (sBP) control (P = 0.013) compared with group 2, as well as lower levels of diastolic BP (dBP), but the results were not statistically significant (Fig. 1). In our study, BP in patients with AH and SH was even lower compared to that in patients without SH. It is still unclear whether mild thyroid dysfunction affects BP, that requires further study using the results of daily BP monitoring.

According to the results of anthropometric measurements (Table 1), the highest mean values of weight, BMI, WC and HC to HC ratio were in group 2 patients. In patients with AH and SH, all anthropometric parameters, except for weight, were statistically higher than those in controls, but not as much as the corresponding values in group 2.

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**Table 1. Anthropometric characteristics of the studied patients**

| Indicators, units | Controls (n = 21) | AH without SH (n = 25) | AH with SH (n = 20) | p1 | p2 | p3 |
|-------------------|------------------|-----------------------|-------------------|----|----|----|
| Weight, kg        | 66.5 [60.0; 82.8] | 88.0 [75.8; 95.5]     | 78.0 [71.0; 88.0] | 0.004 | 0.053 | 0.284 |
| BMI, kg/m²        | 23.8 [21.5; 27.2] | 27.3 [25.4; 28.6]     | 27.1 [26.4; 28.9] | 0.022 | 0.005 | 0.423 |
| WC, m             | 0.805 [0.755; 0.921] | 0.938 [0.796; 1.013] | 0.930 [0.920; 1.030] | 0.008 | 0.003 | 0.875 |
| HC, m             | 0.985 [0.968; 1.053] | 1.010 [0.923; 1.063] | 1.050 [1.020; 1.100] | 0.541 | 0.004 | 0.027 |
| WC/HC             | 0.80 [0.77; 0.90] | 0.92 [0.88; 0.95]     | 0.88 [0.87; 0.93] | 0.003 | 0.035 | 0.125 |

*P1: controls vs AH without SH; P2: controls vs AH with SH; P3: AH without SH vs AH with SH.*
Noteworthy statistically significant WC increase and, as a consequence, the better WC to HC ratio were seen in group 3 patients compared with group 2.

Further distribution of the obtained results into groups depending on the values of WC, WC to HC ratio and BMI revealed an increase in the degree of visceral obesity from the control group to the group of patients with AH and SH (Fig. 2). The proportion of individuals with normal WC among patients with AH and SH decreased, and the percentage of patients with increased and excessive WC as an obligatory component of MS, increased. In addition, the gynoid morphotype was predominant among controls, while this ratio changed in group 2, and in group 3, such a protective profile had only 9 % of patients.

Statistically significant differences were found in the levels of TG, LDL-C and HOMA-IR between the controls and AH patients without SH, and in the levels of HOMA-IR between the controls and AH patients with SH (Table 2). There was also a tendency for increased levels of glucose, TC, LDL-C, AC in group 3 patients compared with those in group 2.

Analysis of the MS prevalence, its components, as well as different types of dyslipidemia among patients of different groups (Table 3) showed that the best metabolic profile in terms of CVD prevention was more common in patients with AH without SH in comparison to AH patients with SH. The prevalence of MS was doubled in patients with SH. Notably, in group 3 patients, there was an increase in the frequency of lipid disorders and their combinations.
in addition to the tendency for deterioration of mean lipid values compared with other groups.

Discussion

Although patients with overt hypothyroidism often complain of weight gain, the association between SH and weight was less clear, especially in AH patients. Dey A., Kanneganti V., Das D. (2019) found that among SH patients with a mean age of 35.1 ± 10.26 years, 12 % were overweight, 24 % were obese, 44 % had a high waist-to-hip ratio, indicating on central obesity, 20 % had high both BMI and waist-to-hip ratio [13]. A significant relationship between central obesity and SH was found by T. M. J. Santhoshakumari, M. Sneha (2019) [14]. On the other hand, there was an assumption that weight gain contributed to the further development of SH. Zynat J. et al. (2020) found that abdominal obesity was a risk factor for elevated levels of antibodies to thyroid peroxidase in men [15]. The results of another study showed that the average level of TSH increased with increasing BMI [16]. The results of our study indicated a decrease in the severity of constitutional deviation in patients with AH and concomitant SH. However, we did not evaluate the level of antibodies to thyroperoxidase or antithyroid antibodies in patients. It is interesting, that in patients with AH and SH, the waist-to-hip ratio was better that in patients without SH. Therefore, there is a need for a more in-depth study of the “safe” limits of HC in AH patients with SH. After all, it is impossible to exclude the formation of a specific morphotype in patients with AH and SH in order to improve the processes of catabolism and protect the body from excessive accumulation of adipose tissue.

According to C. Xu et al. (2019), the proportion of individuals with high glucose levels in the group of people with SH was much higher than in controls. The risk of diabetes mellitus was likewise 2.29 times increased among people with SH [17]. In the current study, fasting glucose levels and HOMA-IR were increased, as well as the prevalence of elevated glucose levels and insulin resistance in people with AH and SH, but these changes were not significant.

The results of the available studies emphasize that SH patients have higher lipid levels, which may not correspond to the range of dyslipidemias, but contribute to the CVR in patients. Dey A., Kanneganti V., Das D. (2019) found that 92 % of SH patients had dyslipidemia. Among them, there were 36 % of hyperglycemia, 36 % – hypercholesterolemia, 32 % – increased LDL-C, 64 % – increased HDL-C, 68 % – reduced HDL-C [13]. Evaluation of our study results revealed the tendency for the lipid profile deterioration and increase in the incidence of dyslipidemia in patients with AH and SH. The levels of TG and LDL-C were slightly better in patients with AH and SH compared to those in group 2, but the differences were not statistically significant. The detected changes may be related to lipid metabolism in the liver and require careful study on the lipid profile changes in patients with AH and SH depending on the liver functional status in this category of patients.

In a study of R. Li et al. (2020), a multivariate analysis conducted among patients in China identified SH as an independent risk factor for the development of MS [18]. Interestingly, the multivariate logistic regression analysis performed by L. Jiang et al. (2020), demonstrated the opposite impact, namely BMI and BP were independent predictors of SH development in women [19]. In our study, the prevalence of MS among patients with AH and SH was also higher as compared with patients of other groups.

Conclusions

1. HC is statistically increased in patients with AH and SH compared with euthyroid AH patients.
2. In patients with AH and SH compared with the group of euthyroid AH patients, the increase in the values of fasting glucose, TC, LDL-C, AC is seen, while the levels of TG, HDL-C, on the contrary, tend to improve, although these changes are not statistically significant.

Table 2. Fasting glucose levels and lipid profile in the studied patients

| Indicators, units | Controls (n = 21) | AH without SH (n = 25) | AH with SH (n = 20) | p 1 | p 2 | p 3 |
|-------------------|-------------------|------------------------|--------------------|-----|-----|-----|
| Glucose, mmol/l   | 5.05 [4.87; 5.37] | 5.27 [4.86; 5.60] | 5.36 [4.85; 5.74] | 0.315 | 0.320 | 0.607 |
| HOMA-IR           | 2.52 [2.26; 2.97] | 3.92 [2.96; 6.12] | 4.32 [2.87; 7.25] | 0.034 | 0.021 | 0.114 |
| TC, mmol/l        | 5.08 [4.68; 5.84] | 5.47 [4.58; 6.11] | 6.12 [4.37; 7.41] | 0.460 | 0.252 | 0.336 |
| TG, mmol/l        | 1.15 [0.88; 1.37] | 1.48 [1.12; 2.25] | 1.17 [1.02; 1.51] | 0.027 | 0.433 | 0.266 |
| VLDL-C, mmol/l    | 0.58 [0.39; 0.62] | 0.67 [0.51; 1.02] | 0.67 [0.46; 0.72] | 0.045 | 0.181 | 0.578 |
| HDL-C, mmol/l     | 1.51 [1.18; 1.67] | 1.23 [1.08; 1.59] | 1.31 [1.19; 1.62] | 0.071 | 0.528 | 0.343 |
| AC                | 2.76 [2.07; 3.34] | 3.12 [2.02; 4.27] | 3.41 [2.21; 4.40] | 0.121 | 0.214 | 0.866 |
| LDL-C, mmol/L     | 3.13 [2.84; 3.58] | 3.34 [2.59; 4.18] | 4.05 [2.28; 4.96] | 0.307 | 0.207 | 0.297 |

P1: controls vs AH without SH; P2: controls vs AH with SH; P3: AH without SH vs AH with SH.

Table 3. Frequency of metabolic syndrome detection, its components and different types of dyslipidemia in the studied groups

| Indicators, units | Controls (n = 21) | AH without SH (n = 25) | AH with SH (n = 20) |
|-------------------|-------------------|------------------------|--------------------|
| MS, %             | 0.0               | 9.1                    | 18.2               |
| Dyslipidemia, %   | 68.2              | 78.8                   | 72.7               |
| Components of MS  |                   |                        |                    |
| ↑TG + ↑LDL-C, %   | 0.0               | 12.1                   | 18.2               |
| ↑TG + ↑glucose, % | 4.5               | 15.2                   | 9.1                |
| ↑HDL-C + ↑glucose, % | 4.5         | 15.2                   | 9.1                |
| ↑glucose, %       | 18.2              | 30.3                   | 36.4               |
| ↑HOMA-IR, %       | 9.0               | 36.0                   | 65.0               |
| Types of dyslipidemia |                   |                        |                    |
| ↑TG, %            | 45.5              | 60.6                   | 63.6               |
| ↑HDL-C, %         | 4.5               | 36.4                   | 18.2               |
| ↑LDL-C, %         | 22.7              | 27.3                   | 36.4               |
| ↑TC + ↑HDL-C, %   | 50.0              | 63.6                   | 63.6               |
| ↑TC + ↑LDL-C, %   | 0.0               | 24.2                   | 18.2               |
| ↑TC + ↑HDL-C, %   | 40.9              | 57.6                   | 63.36              |
| ↑TC + ↓HDL-C, %   | 9.1               | 15.2                   | 27.3               |
| ↑LDL-C + ↑HDL-C, %| 13.6              | 21.2                   | 27.3               |
| ↑LDL-C + ↑VLDL-C + ↑HDL-C, % | 0.0     | 9.1                    | 18.2               |
3. The prevalence of combined lipid profile disorders is higher among patients with AH and SH compared with euthyroid AH patients.

4. Metabolic profile is generally deteriorated in patients with AH even in the early stages and low-to-moderate CVR in combination with SH, that further increases the risk of other CVD and type 2 diabetes mellitus development.

5. Timely detection of SH in AH patients and a better monitoring of anthropometric parameters and the lipid and carbohydrate profile indices are necessary in order to prevent complications.

Prospects for further research are to study the relationship between inflammatory markers, oxidative stress indicators and cardiometabolic parameters in AH patients depending on thyroid dysfunction.

Conflicts of interest: authors have no conflict of interest to declare.

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