Effects of menopause and obesity on lipid profiles in middle-aged Taiwanese women: the Chin-Shan Community Cardiovascular Cohort Study

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Abstract

In this cross-sectional study, we examined the associations between lipid profiles and menopausal status, age, and obesity in Taiwanese women. The study population, established in 1990–91, consisted of 671 premenopausal and 872 postmenopausal women from the Chin-Shan Community Cardiovascular Cohort (CCCC). The associations of age, body mass index (BMI), and menopausal status with serum levels of total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), triglycerides (TG), apoproteins (Apo) A-1 and B, and lipoprotein (a) [Lp (a)] were evaluated. The results showed that menopause was associated with significant increases in TC, LDL-C, TG, and Apo B levels (all P < 0.001). Total cholesterol, LDL-C, TG, and Apo B levels increased consistently with BMI in middle-aged women, regardless of menopausal status. Among women aged 45–49, menopausal women had significantly higher levels of TC and LDL-C than premenopausal women (P < 0.01). However, TG and Apo B levels were higher in postmenopausal than in premenopausal women aged 50–54 years (P < 0.05). Standardized regression analyses showed all lipid variables, except those of Apo A1 and Lp (a) before menopause and TC, LDL-C, and Lp (a) after menopause, were significantly associated with BMI (all P < 0.01). We conclude serum lipid levels in Taiwanese women are no more strongly associated with menopause and BMI than with age. © 2000 Elsevier Science Ireland Ltd. All rights reserved.

Keywords: Menopause; Obesity; Lipids; Lipoproteins; Apoproteins; Taiwanese women

1. Introduction

The Framingham study reported that women usually lag men by 10 years in the development of coronary heart diseases (CHD) [1]. This difference is generally lost in women after menopause; the incidence of coronary events in menopausal women is quadruple that of premenopausal women of the same age [1]. Numerous authors consider menopause as a risk factor for CHD [1–3]. In contrast, Casiglia et al. [4] reported menopause did not influence cardiovascular disease risk in a 16-year longitudinal study.

Menopause is reported to be associated with increased serum levels of total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), triglycerides (TG), apoprotein B (Apo B) [5–10], and lipoprotein (Lp(a)) [5,6], and decreased levels of apoprotein A1 (Apo A1) [5] and high-density lipoprotein cholesterol (HDL-C) [5,7,9]. Several epidemiological studies have reported

Abbreviations: Apo A1, apoprotein A1; Apo B, apoprotein B; BMI, body mass index; CCCC, Chin-Shan community cardiovascular cohort; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; Lp (a), lipoprotein (a); TC, total cholesterol; TG, triglyceride.

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that high serum TC [2,11,14], TG [2], LDL-C [2], lipoprotein (a) [Lp (a)] [14], and Apo B [13] and low HDL-C [2,12,14] levels are risk factors for cardiovascular disease (CVD) mortality in females.

Obesity is also considered an independent risk factor for CVD [15] and the increased risk extends to even mildly or moderately overweight women [16]. Prospective studies have shown a strong correlation between CHD and the amount of abdominal visceral adipose tissue and LDL-C/HDL-C and Apo A1/Apo B ratios [17,18].

Previous studies delineating the relationship between menopause and serum lipid levels were usually based on small sample sizes, with a limited number of studies depicting the relationship of menopause with Apo A1, Apo B, and Lp (a) concentrations. Additionally, few studies have included Asian women in their cohorts; the study by Lyu et al., comparing Taipei and Framingham women, was based only on Taipei government employees [8]. The confounding effect of obesity has rarely been incorporated in comparisons of lipid profiles between menopausal stages. Of importance is that CVD prevalence in premenopausal Taiwanese women is lower than men, but this pattern is reversed for postmenopausal women. The hypothesis is significant changes in lipid profiles may occur in postmenopausal Taiwanese women [19].

The Chin-Shan Community Cardiovascular Cohort (CCCC) study, established in 1990–91, is a community-based investigation of CVD in adults consisting of 1899 men and 1703 women aged 35 and above. The purpose of the present study was to examine the relationships between serum lipid profiles and menopause, obesity, and age among women in Taiwan.

2. Patients and methods

2.1. Study population

Chin-Shan is a suburban community located 20 miles outside metropolitan Taipei. The mortality pattern in Chin-Shan was similar to that of the general Taiwanese population from 1980 to 1994 [20]. Women undergoing hormone replacement therapy prior to, or during the CCC project study, were excluded from this report. This study was based on the baseline data on the year 1990–91. Women with a history of diabetes, fasting sugar > 140 mg, hysterectomy, oophorectomy, oral contraceptive pill use, or menopause before the age of 45 years were excluded from the current study. Only subjects with secondary amenorrhea of at least 1 year were defined as menopausal. Women who still experienced menstruation 3–11 months prior to this study were also excluded since it was uncertain whether they were at the stage of menopause.

2.2. Data collection and assays

A study clinic was set up at the Chin-Shan Community Health Center (CCHC). It consisted of ten senior residents and cardiologists (TCS, KLC, YTL) in conjunction with the surveillance group of 20 senior medical students, two assistant nurses, local practitioners, and CCHC officers. Trained medical students canvassed door-to-door with the assistance of community leaders to extend invitations for the baseline survey. The CCC cohort consisted of 4350 adult aged 35 and above. Among 3602 respondents (82.8%) in the baseline survey of the cohort, 47.3\% (n = 1703) were men and 52.7\% (n = 1899) were women.

Non-respondents included 95 (2.2\%) refusals and 652 (15.0\%) individuals who could not be reached. Information collected included sociodemographic characteristics, life style factors, dietary characteristics, personal and family histories of disease, and hospitalization records. Written consent was obtained during the face-to-face questionnaire interview, and physicians later conducted physical examinations and laboratory tests on participants invited to the clinic. Specimens for blood analyses were also collected at the clinic.

All venous blood samples were taken after a 10-h overnight fast, immediately refrigerated, and transported to the National Taiwan University Hospital within 6 h. Assay of serum TC and TG levels were performed within 24 h. Serum samples were then stored at −70°C for batch assay of LDL-C, HDL-C, Apo A1, Apo B, and Lp (a) as previously described [21,22]. These determinations were usually completed within 2 weeks. Few specimens were repeated for the determinations of HDL-C, LDL-C, Apo A1 and Apo B after storage of about 9 to 12 months and of Lp (a) after storage of about 1–2 years. Standard enzymatic methods for serum TC and TG were used (Merck 14354 and 14366, Germany). The HDL-C concentration in the supernatant was measured after precipitation with magnesium chloride phosphotungstate reagents (Merck 14993). The concentration of LDL-C was calculated as ‘total cholesterol — cholesterol in the supernatant’, and LDL-C was precipitated by heparin-citrate reagent (Merck 14992) [23]. Apo A1 and Apo B concentrations were measured by turbidimetric immunoassay using commercial kits (Sigma, USA). Lp(a) was measured with an enzyme-linked immunosorbent assay kit (Organon, USA). Lp(a) was measured with an enzyme-linked immunosorbent assay kit (Organon, USA). Lp(a) was measured with an enzyme-linked immunosorbent assay kit (Organon, USA). Lp(a) was measured with an enzyme-linked immunosorbent assay kit (Organon, USA). Lp(a) was measured with an enzyme-linked immunosorbent assay kit (Organon, USA). Lp(a) was measured with an enzyme-linked immunosorbent assay kit (Organon, USA).

BMI, an indicator related to the amount of visceral fat [24], was used as a measure of obesity and was calculated as body weight in kg divided by height in m^2; these parameters were measured at the time of blood sampling. A BMI of 26 kg/m^2 or greater was considered
an indication of obesity, based on a Chinese population study in Taiwan [25].

2.3. Statistical analyses

We attempted to isolate the effects of menopause by incorporating the effects of age and obesity in the statistical analyses. We first compared the overall average level of each lipid by menopausal status, and the average level of each lipid for every 5-year-age band from 35–39 through 55–59 years old for premenopausal women and from 45–49 through 60 and above for postmenopausal women. Pearson correlation analysis was performed to measure the relationships between the age bands and average lipid levels. We also compared the average menopause-specific lipid profiles for women aged 45–49 and those aged 50–54 years.

Lipid profiles with respect to obesity (BMI < 26 kg/m²), were further categorized into five groups (BMI < 21, 21–23, 23–25, 25–27, and ≥ 27 kg/m²) for women aged between 45 and 54 years. A multiple linear regression analysis was then used to examine the relationships between age, BMI, and each lipid profile for premenopausal and postmenopausal women. All statistical analyses were performed with SAS statistical software (Version 6.12, SAS Institute, Cary NC, USA). P values below 0.05 were considered statistically significant.

3. Results

One thousand five hundred and forty three (81.3%) of the 1899 respondents met the entry criteria for this study, including 671 premenopausal and 872 postmenopausal women. The baseline average age (standard deviation) for women included in the analysis was 52.7 (12.2) years (range, 35–96 years). The mean age was 42.7 (5.8) years for premenopausal women and 61.2 (9.5) years for postmenopausal women. The mean age at the onset of menopause was 49.7 (4.3) years, and the average duration since menopause was 12.9 (9.1) years.

Data analyses revealed no significant differences in the effects of smoking or alcohol consumption on lipid profiles between premenopausal and postmenopausal women (data not shown). Only 4% of women were smokers, and few women drank regularly. Table 1 depicts the differences in lipid profiles between premenopausal and postmenopausal women. There was no difference in average BMI between premenopausal and postmenopausal women. However, the levels of TC, TG, LDL-C, and Apo B, but not HDL-C, Apo A1, or Lp (a), were significantly higher in postmenopausal women. The Apo B/Apo A1 and the TC/HDL-C ratio was thus higher in postmenopausal than in premenopausal women. However, the median value of Lp (a) was higher in postmenopausal women (9.7 mg/dl) than in premenopausal women (8.3 mg/dl). When CHD risk classifications were selected, significantly greater proportions of the premenopausal group had desirable levels of TC (< 200 mg/dl; 68.6 vs 41.0%) and LDL-C (< 130 mg/dl; 58.9 vs 33.0%) than did the postmenopausal group (both, P < 0.001). Postmenopausal women had a higher percentage of low HDL-C levels (< 35 mg/dl) than premenopausal women (11.6 vs 7.7%, P < 0.05) had.

Fig. 1 shows the serum lipid levels by age and menopausal status. TC, LDL-C, and TG tended to rise as age increased, with significant increases only for TC and LDL-C in premenopausal women (P < 0.05). However, premenopausal women had lower levels of TC, LDL-C, and TG than postmenopausal women at every age band between 45 and 59 years. The largest gap between the two groups (20 mg/dl for TC or LDL-C)

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Serum lipid concentrations by menopausal status among women in the Chin-Shan Community Cardiovascular Cohort Study*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Premenopause (n = 671)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>42.7 ± 5.8</td>
</tr>
<tr>
<td>Body height (cm)</td>
<td>154.7 ± 5.4</td>
</tr>
<tr>
<td>Body weight (kg)</td>
<td>56.9 ± 8.8</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>23.8 ± 3.4</td>
</tr>
<tr>
<td>TC (mg/dl)</td>
<td>186.6 ± 40.0</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>98.7 ± 67.6</td>
</tr>
<tr>
<td>HDL-C (mg/dl)</td>
<td>49.7 ± 11.3</td>
</tr>
<tr>
<td>LDL-C (mg/dl)</td>
<td>125.3 ± 38.4</td>
</tr>
<tr>
<td>TC/HDL-C</td>
<td>3.92 ± 1.21</td>
</tr>
<tr>
<td>Apo A1 (g/l)</td>
<td>1.30 ± 0.23</td>
</tr>
<tr>
<td>Apo B (g/l)</td>
<td>0.83 ± 0.26</td>
</tr>
<tr>
<td>Apo B/Apo A1</td>
<td>0.65 ± 0.22</td>
</tr>
<tr>
<td>Lp (a) (mg/dl) (median)</td>
<td>13.8 ± 15.9 (8.3)</td>
</tr>
</tbody>
</table>

* Data are expressed as the mean ± standard deviation (and median values for Lp (a)).

** P, P-value for difference between pre- and post-menopausal groups (t-test)
was in the 45–49 year age band. The change in HDL-C concentration was associated with age but not menopause. In both groups, the highest average HDL-C and Apo A1 concentrations were found in the age band from 50 to 54 years. Average Lp (a) levels tended to drop in the older age bands, but only the decrease in HDL-C was significant \((P < 0.05)\). Among postmenopausal women, the Lp (a) level increased again after the ages of 55–59.

Table 2 compares menopausal-specific average levels of serum lipid among women of two age groups, 45–49 and 50–54 years. Again, postmenopausal women had significantly higher TC and LDL-C levels than premenopausal women at younger (both \(P < 0.01\)) but not older ages. In the older group, TG and Apo B levels were significantly higher in postmenopausal women (both \(P < 0.05\)). The postmenopausal women also had a higher average Apo B/ApoA1 ratio (\(P < 0.05\) in the older group) and higher median values of Lp (a).

Fig. 2 shows that, with the exception of HDL-C, Apo A1, and Lp (a) concentrations, the levels of serum lipids and Apo B were elevated in postmenopausal women, compared with premenopausal women, regardless of their obesity status. These differences were greater in non-obese than in obese women.

Fig. 3 shows the lipid profile changes with BMI and menopausal status for women 45–54 years old. The average LDL-C level in premenopausal women increased steadily from 120 to 146 mg/dl when BMI was stratified into levels of <21, 21–23, 23–25, 25–27, and \(\geq 27\) kg/m². The average TC value in premenopausal women also increased as BMI increased, but with a weaker trend. TG and Apo B levels were also elevated in postmenopausal women, except for TG in women with BMI \(\geq 27\) kg/m².

Standardized regression analyses by age and BMI shown in Table 3 suggest that age had a positive correlation with premenopausal TC, LDL-C levels and postmenopausal Apo B/Apo A1 ratio (\(P < 0.05\)), and a negative correlation with postmenopausal HDL-C (\(P < 0.05\)). BMI was positively correlated with TG and Apo B concentration, as well as the TC/HDL-C and the Apo B/Apo A1 ratio, and negatively correlated with HDL-C concentration in both pre- and postmenopausal women (all \(P < 0.001\)). Significant trends between BMI and TC and LDL-C were found only in premenopausal women (both \(P < 0.001\)), and between BMI and Apo A1 in only postmenopausal women (\(P < 0.01\)).
Table 2
Menopause-specific average serum lipid concentrations of women aged 45–49 and 50–54 years in the Chin-Shan Community Cardiovascular Cohort Study

<table>
<thead>
<tr>
<th>Age</th>
<th>45–49</th>
<th>Postmenopause</th>
<th>50–54</th>
<th>Postmenopause</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Premenopause</td>
<td></td>
<td>Premenopause</td>
<td></td>
</tr>
<tr>
<td></td>
<td>n = 147</td>
<td>n = 63</td>
<td>n = 56</td>
<td>n = 162</td>
</tr>
<tr>
<td>TC (mg/dl)</td>
<td>186.6 ± 40.6*</td>
<td>205.0 ± 42.0*</td>
<td>200.3 ± 30.3</td>
<td>209.3 ± 47.3</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>101.9 ± 75.5</td>
<td>114.1 ± 64.6</td>
<td>98.1 ± 54.5**</td>
<td>119.5 ± 76.5**</td>
</tr>
<tr>
<td>HDL-C (mg/dl)</td>
<td>48.7 ± 11.5</td>
<td>49.2 ± 11.0</td>
<td>52.3 ± 10.1</td>
<td>51.6 ± 13.2</td>
</tr>
<tr>
<td>LDL-C (mg/dl)</td>
<td>127.0 ± 40.5*</td>
<td>144.5 ± 40.5*</td>
<td>136.5 ± 31.8</td>
<td>145.2 ± 46.9</td>
</tr>
<tr>
<td>TC/HDL</td>
<td>4.05 ± 1.32</td>
<td>4.33 ± 1.13</td>
<td>3.98 ± 1.03</td>
<td>4.31 ± 1.41</td>
</tr>
<tr>
<td>Apo A1 (g/l)</td>
<td>1.31 ± 0.27</td>
<td>1.31 ± 0.21</td>
<td>1.37 ± 0.20</td>
<td>1.31 ± 0.22</td>
</tr>
<tr>
<td>Apo B (g/l)</td>
<td>0.87 ± 0.29</td>
<td>0.94 ± 0.26</td>
<td>0.87 ± 0.26**</td>
<td>0.99 ± 0.31**</td>
</tr>
<tr>
<td>Apo B/Apo A1</td>
<td>0.68 ± 0.24</td>
<td>0.72 ± 0.21</td>
<td>0.63 ± 0.22**</td>
<td>0.77 ± 0.23**</td>
</tr>
<tr>
<td>Lp (a) (mg/dl) (median)</td>
<td>16.2 ± 17.8 (8.7)</td>
<td>16.4 ± 16.6 (9.8)</td>
<td>12.6 ± 14.6 (7.4)</td>
<td>12.1 ± 11.8 (9.0)</td>
</tr>
</tbody>
</table>

* Data are expressed as the mean ± standard deviation (and median values for Lp (a)).
* P < 0.01 (t-test).
** P < 0.05 (t-test).

Fig. 2. Serum concentrations of lipids (mg/dl, and g/l for apoproteins) by menopausal status and obesity (body mass index (BMI) < 26 kg/m²: non-obese; BMI ≥ 26 kg/m²: obese) in women in Chin-Shan Community, Taiwan. **, P < 0.01; ***, P < 0.001 (obese vs non-obese groups, t-test).
4. Discussion

4.1. Lipid profiles in premenopausal and postmenopausal women

Lyu et al. [8] reported female age-adjusted CHD mortality was two-fold higher in the Framingham than in the Taipei population [8]. He attributed this difference to lower LDL-C and higher HDL-C levels in Taipei women. We found the population in Chin-Shan had a CHD mortality rate similar to that of women in Taipei, but had higher TG and LDL-C levels, and lower HDL-C levels, than pre- and postmenopausal women in both Taipei and Framingham. While Framingham women had the highest TC concentrations and Chin-Shan women the lowest, Taipei women had the highest HDL-C levels and Chin-Shan women the lowest. Compared with Chin-Shan women, both Taipei and Framingham women were younger, with 2/3 of these women in the premenopausal state, these women accounted for only 44% of the current study population.

The levels of TC, LDL-C, HDL-C, Apo A1, Apo B, and Lp (a) were significantly higher in postmenopausal than in premenopausal women in Taipei, while only TC, TG, LDL-C, and Apo B levels were higher in postmenopausal women the in Framingham study. Several studies [5–10] reported changes in lipid profiles after menopause similar to those of the Chin-Shan women. However, a decrease in serum HDL-C levels after menopause was not seen in this study. This result may be controversial because the subclasses of HDL-C [26], which may be individually associated with menopause [7] not measured in our study.

Table 3
Regression coefficients (Pearson’s $r$) for associations of lipid concentrations with age and BMI (kg/m²) by menopausal status for women in Chin-Shan Community Cardiovascular Cohort Study

<table>
<thead>
<tr>
<th></th>
<th>Premenopause</th>
<th>Postmenopause</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Age</td>
<td>BMI</td>
</tr>
<tr>
<td>TC (mg/dl)</td>
<td>0.70*</td>
<td>2.10**</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>0.74</td>
<td>6.46**</td>
</tr>
<tr>
<td>HDL-C (mg/dl)</td>
<td>0.01 -0.85**</td>
<td>-0.11*</td>
</tr>
<tr>
<td>LDL-C (mg/dl)</td>
<td>0.67*</td>
<td>2.76**</td>
</tr>
<tr>
<td>Apo A1 (g/l)</td>
<td>0.0009</td>
<td>-0.006</td>
</tr>
<tr>
<td>Apo B (g/l)</td>
<td>0.003</td>
<td>0.02**</td>
</tr>
<tr>
<td>Apo B/Apo A1</td>
<td>0.004</td>
<td>0.04**</td>
</tr>
<tr>
<td>Lp (g) (mg/dl)</td>
<td>-0.02</td>
<td>0.14</td>
</tr>
</tbody>
</table>

* $P<0.05$.
** $P<0.001$.
*** $P<0.01$. 

Fig. 3. Serum concentrations of lipids (mg/dl, and g/l for apoproteins) by body mass index (BMI) category (< 21, 21–23, 23–25, 25–27, and ≥ 27 kg/m²) in premenopausal and postmenopausal women 45–54 years of age in Chin-Shan Community, Taiwan. There were 25, 55, 54, 40, and 37 premenopausal women and 39, 57, 50, 36, and 35 postmenopausal women in the respective BMI categories.
The HDL-C level is universally related to CHD, although CHD is influenced by a variety of factors including biological, environmental, and behavioral characteristics [28]. Thus, differences in the levels of LDL-C and HDL-C between the Chin-Shan and Framingham cohorts cannot explain the lower rate of CHD mortality in the Chin-Shan study group. However, the average BMI of Framingham women was higher than that of Chin-Shan women [8]. It may be of interest to note that, due to religious beliefs, traditional families in Chin-Shan may consume more frequently a Taiwanese vegetarian diet. These meals consist mainly of soybean products, which have been claimed as the phyto-oestrogens products that may improve menopausal syndrome and prevent atherosclerosis [29]. It has been found the consumption of soybean may decrease plasma TC and LDL-C levels [30], though the other study found no alteration in plasma lipids in menopausal and perimenopausal women [31]. In addition, fish is also a primary protein source in Chin-Shan families as a fishing village is near by. Women in this study cohort are also active in their daily routines.

4.2. Effect of age on lipid changes in the perimenopausal period

The present study revealed changes in lipid profiles in menopausal women are age-associated. The TC and LDL-C levels were significantly higher in post-menopausal women aged 45–49 years than in pre-menopausal women in the same age group, as were the levels of TG and Apo B among women aged 50–54 years. Because no significant HDL-C change was observed in menopausal women, the TC change reflects the LDL-C change. This age-associated increase in TC and in LDL-C appeared to be significant only in pre-menopausal women, i.e. once the LDL-C level increases in response to menopause, the age association with LDL-C became less apparent. Other cross-sectional and longitudinal studies have examined similar patterns of TC, TG, and LDL-C levels in menopausal women. Both TC and LDL-C levels may be remarkably elevated from 1 to 2 years before menopause [10,32] to years after menopause [32,33]. Serum TG levels can show a similar pattern of increase before menopause [10]. The LDL-C level was reported to rise substantially with age in women < 53 years old [6]. In CCCC study, the association between LDL-C concentration with menopause was greatest if menopause occurred between the ages of 45 and 49 years.

Recent studies have debated the association between obesity, age and lipids and circulating estrogen levels [34–36]. The change in estrogen levels has a significant effect on the redistribution of lipids [37]. The estrogen levels are unknown for the women in this study.

4.3. The effect of obesity on lipid changes

Obesity in women has been found to be strongly associated with elevated levels of TC, LDL-C, [18,38] and TG, and lowered HDL-C [18,36,38], even after controlling for age [38] and estrogen levels [36]. This correlation is strongest in women 35–54 years of age [18]. In this study, we found significant associations between BMI and TC, LDL-C, TG, HDL-C, Apo B levels. This correlation was independent of age, and the rising trend was closely associated with the increase in BMI. This trend was even more obvious at the ages of perimenopausal. A substantial decrease in Apo A1 concentration with decreasing estrogen levels after menopause also has been reported [5,39]. However, we found our Apo A1 levels to be highest in the 50–54 years bands, contrary to the reported declining estrogen levels before [10,40] and after [41] menopause. We also found that after menopause Apo A1 concentration was negatively related to BMI; this has not been reported elsewhere.

Reeder et al. [18] reported that TG and HDL-C levels are most strongly associated with abdominal obesity, and the least strongly associated with LDL-C and TC. They did not consider the effect of menopause. In our study, the associations between BMI and both TC and LDL-C were stronger before than after menopause. This suggests menopause also affects LDL-C and TC rather than obesity itself. Menopausal changes are less dramatic in obese women for TC, TG and LDL-C, but not for Apo B, TC/HDL-C or HDL-C (Fig. 2).

Body fat shifts from the subcutaneous to the intra-abdominal area during menopause, and results in an android-type fat distribution [42,43]. Women with android-type obesity show less favorable lipid and lipoprotein profiles than those with gynoid-type obesity [44]. BMI has long been claimed to be a simple, precise predictor for assessing the risk of abdominal obesity [18]. Obesity itself presents a rather complex metabolic status. Obesity in women is associated with insulin resistance and an altered lipoprotein composition [45]. In our study, the change in lipid profiles (in TC, TG, and LDL-C) after menopause was slightly less dramatic in obese than in non-obese women. No BMI change was seen in Chin-Shan women after menopause. This may suggest that metabolic characteristics causing dyslipidemia in obese subjects are more important than the occurrence of menopause.
4.4. Unique feature of Lp (a) in perimenopausal women

Previous studies generally showed small changes in Lp (a) levels in postmenopausal women [2,46,47]. Our data also demonstrated no changes in Lp (a) when overall average values between premenopausal and postmenopausal women were compared, regardless of whether BMI was considered; although, postmenopausal women had a higher median value of Lp (a). We also found Levels of LDL-C, TG and HDL-C corresponded to Lp (a) levels [48]. Serum Lp (a) level has been reported to increase gradually with age in women [6], especially those aged 44–58 years [49]. In this study, the distribution of Lp (a) showed a peak at 45–49 years and did not climb again until after menopause at 55–59 years old. We believe that this unique pattern of Lp levels in Chin-Shan women is influenced by both age and menopause and has no relationship with BMI.

4.5. Conclusions

The changes in lipid profiles after menopause, particularly apoprotein levels, are more complicated for obese women than for non-obese women, and are probably multifactorial in origin. However, this study demonstrated the obvious influence of obesity on TC, LDL-C, TG, and Apo B levels in middle age Chin-Shan women who were experiencing menopause. It is not necessary that serum lipid levels have stronger association with menopause and BMI than with age.

Acknowledgements

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