Original Article

Impact of Apolipoprotein B on Hepatosteatosis in a Population Infected with Hepatitis C Virus: A Cross-Sectional Observational Study

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Key Words

Hepatitis C virus · Apolipoprotein-B · Non-alcoholic fatty liver disease

Abstract

Objective: Non-alcoholic fatty liver disease (NAFLD) is an established risk factor for diabetes, cardiovascular disease, antiviral treatment resistance, and progression of chronic hepatitis C virus (HCV) infection to fibrosis. Apolipoprotein-B 100 (ApoB-100) is a dyslipidemia marker and steatosis predictor. We assess the correlation between ApoB-100 and hepatosteatosis. **Methods:** This cross-sectional study enrolled 1,218 HCV-seropositive participants from a 2012–2013 health checkup in Taiwan. NAFLD was detected using ultrasound. All anthropometric and laboratory studies that included ApoB-100 were evaluated whether or not ApoB-100 predicts NAFLD. Logistic regression was also used to examine the association between ApoB-100 and NAFLD. **Results:** Participants were 47.16 ± 16.08 years old (mean age). The overall prevalence of NAFLD was 35.8% ($n = 436$; 32.8% men, 38.1% women). Participants with $\text{ApoB-100} \geq 8$ had a significantly higher incidence of NAFLD (39.4 vs. 29.4%; 95% CI 0.044–0.156; $p < 0.001$). After confounding factors had been adjusted for, ApoB-100 was significant-

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ly associated with NAFLD (OR 5.45; 95% CI 1.64–18.06; $p = 0.006$) and high-grade hepatosteatosis (OR 7.73; 95% CI 1.74–34.35; $p = 0.007$). **Conclusion:** ApoB-100 is strongly associated with NAFLD in people with non-genotype 3 HCV; greater ApoB-100 content is significantly correlated with higher-grade hepatosteatosis.

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Introduction

Hepatitis C virus (HCV) infection is an emerging health concern with an estimated global prevalence of 3% and 185 million chronic carriers worldwide [1]. Chronic HCV progresses from steatosis over fibrosis to cirrhosis and hepatocellular carcinoma. In addition, HCV increases insulin resistance and the incidence of metabolic syndrome and diabetes mellitus (DM) [2, 3], all of which contribute to the extrahepatic complications of systemic atherosclerosis and coronary artery disease [4].

Hepatosteatosis in chronic hepatitis C (CHC) is estimated to have a prevalence of 55–81%, which is significantly higher than its overall prevalence (15–40%) [5, 6]. Non-alcoholic fatty liver disease (NAFLD) is significantly associated with metabolic syndrome [7], which increases necroinflammation [8], accelerate the development of fibrosis, increases refractoriness to anti-viral therapy [9], and causes early atherosclerosis [10]. Therefore, the complex relationships between hepatosteatosis, insulin resistance, hepatic fibrosis and systemic inflammation contribute to higher mortality in CHC [11].

Many atherolipid profiles are associated with NAFLD. Apolipoprotein-B100 (ApoB-100) is the primary protein responsible for carrying cholesterol, including very low-density lipoprotein (VLDL) and low-density lipoprotein (LDL), to tissue, which might be inaccurately measured in hypertriglyceridemia and NAFLD [12, 13]. Studies have reported a strong association between ApoB, metabolic syndrome, DM [14] and cardiovascular events [15–17]. In addition, ApoB-100 is also a predictor of NAFLD in patients with [18] and without DM [19]. Despite the hypobetalipoproteinemia with hypocholesteremia observed in patients with genotype-3 HCV [20], there was no additional evidence of a correlation between ApoB-100 and steatosis in non-genotype 3 CHC.

To examine the involvement of ApoB-100 in HCV-related hepatosteatosis, we investigated the association between various measures of body weight (BW), BMI, waist-to-hip ratio (WHR), metabolic syndrome, and NAFLD in non-genotype 3 HCV patients.

Material and Methods

Sample and Study Design

The present study is a cross-sectional study based on an annual community health checkup. We aimed to determine the relationship between associated factors and hepatosteatosis in an HCV hyperendemic village. From the end of 2012 until August 2013, the health checkup screened 12,348 residents. The eligibility criteria for the present study included being > 20 years old and HCV-seropositive. Exclusion criteria were an incomplete health checklist as well as missing anthropometric records, laboratory data, and hepatitis viral markers, or a missing echogenic study. Finally, 1,218 eligible residents were identified and enrolled in the analysis (fig. 1). Ethical approval was granted by the hospital's Institutional Review Board (IRB: 103–6854B).

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Fig. 1. Flow chart of inclusion criteria and participant selection.

Anthropometric, Blood Pressure, and Laboratory Measurements

Three measurements of waist circumference were taken midway between the lowest rib and the iliac crest to an accuracy of 0.5 cm. The hip was measured at the widest portion on the left side, and at the greater trochanters on the right. Both were calculated as the WHR. Measurements of weight and height were also taken to calculate the BMI: weight (kg) / height (m²). Blood pressure was measured using a standard electronic sphygmomanometer according to the Hypertension Detection and Follow-up Program protocol (HDFP) while the patient was sitting comfortably with their back supported for at least 5 min. Pulse pressure was calculated as systolic blood pressure (SBP) – diastolic blood pressure (DBP).

After the participants had fasted for at least 12 h, their blood was drawn for biochemical analyses of serum glucose, creatinine, uric acid, triglycerides, total cholesterol, high-density lipoprotein (HDL), VLDL, alkaline phosphatase, alanine aminotransferase (ALT), aspartate aminotransferase, gamma-glutamyl transferase, and ApoB-100 using an 7600 automatic biochemistry analyzer (Model 7699; Hitachi, Tokyo, Japan). ApoB-100 concentrations were measured with the Tina-quant apoB kits (Roche Diagnostics, Taipei, Taiwan) (version with the International Federation of Clinical Chemistry SP3-07 reference preparations) according to World Health Organization standardized methods [21].

Hepatitis B virus (HBV) and HCV seropositivity was assessed using sandwich radioimmunoassay (Elecsys; Cobas analyzer; Roche Diagnostics), a semiquantitative method to determine hepatitis B surface antigen and antibody to HCV using an electrochemiluminescence immunoassay test (Roche Diagnostics), respectively.

Personal Medical History

Participants completed a health checklist for baseline disease and lifestyle habits: smoking, alcohol drinking, and betel nut chewing. DM was defined by a participant's having a clinical diagnosis of the disease or by documented use of anti-diabetes drugs. Hypertension was defined by a participant's having a clinical diagnosis of high blood pressure or by documented use of anti-hypertension medication. Significant

alcohol consumption was defined as ongoing or recent alcohol consumption > 20 g/day. A smoker was defined as someone who had smoked at least 100 cigarettes in their lifetime and was a current smoker at the time of the study. Finally, a betel nut chewer was defined as someone who reported chewed betel nut > once a week.

Abdominal Echography

The ultrasound examinations of supine patients were done using three units with curved array transducers: Aloka SSD 4000 + UST-979–3.5 transducer (Hitachi Aloka Medical, Wallingford, CT, USA), Acuson S2000 + C4–1 MHz transducer (Siemens Limited, Taipei, Taiwan) and a CGM OPUS 5000 + CLA35 transducer (Chang Gung Medical Technology, Taipei, Taiwan). Technical parameters were adjusted for each patient based on a standard protocol. Livers with a homogenous echo texture, no acoustic attenuation, visible portal veins, a well-visualized diaphragm, and an echogenicity similar to or slightly higher than that of renal parenchyma were considered normal [22]. The diagnosis of the fatty liver was based on the brightness of the liver in comparison to the kidney, vascular blurring of hepatic vein trunk, and deep attenuation in the right hepatic lobe. The severity of fatty liver change was divided into four grades (0–3): Grade 0 = normal liver, normal echo texture, and no fatty change; Grade 1 = mild fatty liver change, mild increase in fine echoes in the parenchyma with slightly impaired visualization of the intrahepatic vessels and diaphragm; Grade 2 = medium grade diffuse increase in hepatic echogenicity, mild deterioration in the image of the diaphragm and intrahepatic vessels; Grade 3 = moderate to severe fatty liver change, marked increase in fine echoes in the parenchyma with poor or non-visualization of the intrahepatic vessel borders, diaphragm and posterior right lobe of the liver. All images were reviewed on a picture archiving and communication system (Centricity PACS; GE Healthcare, Ridgewood, NJ, USA).

Definitions

NAFLD

The definition of NAFLD requires i) evidence of hepatic steatosis by imaging and ii) the absence of secondary causes for hepatic fat accumulation such as significant alcohol consumption, use of steatogenic medication, or hereditary disorders. Low-grade NAFLD means grade 1 hepatosteatosis, and high-grade NAFLD means grades 2–3 hepatosteatosis.

Metabolic Syndrome

Metabolic syndrome based on the 2006 National Cholesterol Education Program Adult Treatment Panel III guideline [23] was defined as evidence of three or more of the following: i) waist circumference > 90 cm in men and > 80 cm in women for Asians; ii) fasting plasma glucose ≥ 110 mg/dl or taking anti-diabetes medication; iii) serum triglycerides ≥ 150 mg/dl; iv) serum HDL-cholesterol < 40 mg/dl in men and < 50 mg/dl in women; v) blood pressure $\geq 130/85$ mm Hg or taking anti-hypertension medication.

Statistical Analyses

SPSS 19.0 for Windows (SPSS, Inc., Chicago, IL, USA) was used for all analyses. Baseline characteristics and laboratory findings were compared between HCV-seropositive participants based on NAFLD and ApoB-100 level. Continuous variables, reported as means $\pm$ standard deviation (SD), were compared using Student's t test. Categorical variables are reported as number (%) and were compared using Fisher's Exact test. Factors associated with NAFLD were identified using a multivariate logistic regression analysis. Significance was set at $p < 0.05$ (two-sided).

Table 1. Baseline characteristics differences between HCV seropositive patients with or without NAFLD

Characteristic	All (n = 1,218)	NAFLD		p values
		yes (n = 436)	no (n = 782)	
Age, years	47.16 ± 16.08	47.62 ± 15.89	46.90 ± 16.19	0.459
Gender, female, n (%)	696 (57.1%)	265 (60.8%)	431 (55.1%)	0.055
BW, kg	65.89 ± 13.23	66.19 ± 13.27	65.73 ± 13.22	0.562
BMI ± SD	25.02 ± 4.12	25.24 ± 4.15	24.90 ± 4.09	0.163
BMI > 25, n (%)	577 (47.4%)	214 (49.1%)	363 (46.4%)	0.372
Waist circumference, cm	82.54 ± 11.23	82.86 ± 11.06	82.36 ± 11.32	0.455
WHR ± SD	0.86 ± 0.08	0.86 ± 0.09	0.86 ± 0.08	0.908
SBP, mm Hg	135.90 ± 20.65	138.12 ± 19.91	134.66 ± 20.97	0.005
DBP, mm Hg	80.72 ± 14.27	82.30 ± 13.52	79.84 ± 14.60	0.004
Pulse pressure, mm Hg	55.17 ± 13.94	55.82 ± 13.56	54.82 ± 14.14	0.230
Diabetes mellitus, n (%)	144 (11.8%)	88 (20.2%)	56 (7.2%)	<0.001
Hypertension, n (%)	249 (20.4%)	132 (30.3%)	117 (15.0%)	<0.001
HBV, n (%)	180 (14.8%)	61 (14.0%)	119 (15.2%)	0.563
Smoking, n (%)	131 (10.8%)	59 (13.5%)	72 (9.5%)	0.020
Betel-nut chewing, n (%)	75 (6.2%)	36 (8.3%)	39 (5.0%)	0.023
Creatinine, mg/dl	0.93 ± 0.33	0.92 ± 0.40	0.93 ± 0.28	0.673
Fasting glucose, mg/dl	109.93 ± 34.01	114.44 ± 36.84	107.41 ± 32.07	0.001
ALT, mg/dl	40.30 ± 49.89	41.14 ± 64.12	39.83 ± 39.86	0.662
AST, mg/dl	34.57 ± 32.31	33.42 ± 36.33	35.21 ± 29.84	0.355
Uric acid, mg/dl	5.99 ± 1.60	6.08 ± 1.61	5.94 ± 1.59	0.126
LDL-C, mg/dl	113.63 ± 31.13	115.66 ± 31.28	112.49 ± 31.02	0.089
HDL-C, mg/dl	51.01 ± 13.10	49.04 ± 12.18	52.10 ± 13.47	<0.001
TC, mg/dl	184.20 ± 36.47	187.09 ± 37.90	182.58 ± 35.57	0.038
TG, mg/dl	111.35 ± 67.56	129.66 ± 87.87	101.15 ± 50.21	<0.001
GGT, U/l	35.46 ± 55.99	36.59 ± 43.46	34.83 ± 61.91	0.600
ApoB-100, g/l	0.88 ± 0.23	0.91 ± 0.24	0.86 ± 0.22	<0.001
MetS, n (%)	346 (28.4%)	163 (37.4%)	183 (23.4%)	<0.001

Values are number (%) or mean ± SD; Fisher's Exact test for categorical variables and one-way ANOVA for continuous variables.

ApoB-100 = apolipoprotein-B 100; ALT = alanine aminotransferase; AST = aspartate aminotransferase; GGT = gamma-glutamyl transferase; LDL-C = low-density lipoprotein-cholesterol; HDL-C = high-density lipoprotein-cholesterol; TC = total cholesterol; TG = triglyceride.

Results

Study Population

After we had excluded patients who had not met the inclusion criteria, 1,218 enrollees were included for the final analysis (mean age 47.16 ± 16.08 years; females 57.1%). The prevalence of DM, hypertension, metabolic syndrome and BMI > 25 kg/m² was 11.8% (n = 144), 20.4% (n = 249), 28.4% (n = 346) and 47.4% (n = 577), respectively. The participants were divided into the NAFLD (n = 436) and non-NAFLD (n = 782) groups (table 1). There were non-significantly more women in the NAFLD group (60.8% vs. 55.1%; p = 0.055). Despite significantly higher percentages of DM, hypertension (p < 0.001), smoking (p = 0.020), betel-nut chewing (p = 0.023) in the NAFLD group, age, WHR, BW, and BMI were not significantly different. Patients with NAFLD had typical metabovascular features with significantly lower HDL (mean 49.04 ± 12.18 vs. 52.10 ± 13.47 mg/dl; p < 0.001) but significantly higher total cholesterol (187.09 ± 37.90 vs. 182.58 ± 35.57 mg/dl; p = 0.038), triglycerides (129.66 ± 87.87 vs. 101.15 ± 50.21 mg/dl; p < 0.001), fasting glucose (mean 114.44 ± 36.84 vs.

Table 2. Baseline characteristics differences between patients with HCV based on ApoB-100 content

Characteristic	ApoB $\geq$ 0.8 (n = 782)	ApoB < 0.8 (n = 436)	p values
Age, years	47.64 $\pm$ 16.01	46.30 $\pm$ 16.20	0.164
Gender, female, n (%)	440 (56.3%)	256 (58.7%)	0.408
BW, kg	66.19 $\pm$ 13.14	65.36 $\pm$ 13.40	0.299
BMI	25.11 $\pm$ 4.07	24.85 $\pm$ 4.19	0.298
BMI > 25, n (%)	364 (46.5%)	213 (48.9)	0.440
Waist circumference, cm	82.86 $\pm$ 11.25	81.97 $\pm$ 11.18	0.184
WHR	0.86 $\pm$ 0.08	0.85 $\pm$ 0.08	0.104
SBP, mm Hg	136.67 $\pm$ 20.74	134.51 $\pm$ 20.45	0.080
DBP, mm Hg	81.28 $\pm$ 14.43	79.72 $\pm$ 13.93	0.066
Pulse pressure, mm Hg	55.39 $\pm$ 14.17	54.79 $\pm$ 13.52	0.477
Diabetes mellitus, n (%)	84 (10.7%)	60 (13.8%)	0.118
Hypertension, n (%)	172 (22.0%)	77 (17.7%)	0.072
HBV, n (%)	121 (15.5%)	59 (13.5%)	0.360
Smoking, n (%)	79 (10.1%)	52 (11.9%)	0.325
Betel-nut chewing, n (%)	48 (6.1%)	27 (6.2%)	0.970
Creatinine, mg/dl	0.94 $\pm$ 0.36	0.91 $\pm$ 0.27	0.182
Fasting glucose, mg/dl	110.64 $\pm$ 34.45	108.64 $\pm$ 33.19	0.323
ALT, mg/dl	44.60 $\pm$ 44.31	37.90 $\pm$ 52.63	0.025
AST, mg/dl	39.64 $\pm$ 35.49	31.74 $\pm$ 30.04	< 0.001
Uric acid, mg/dl	6.11 $\pm$ 1.62	5.78 $\pm$ 1.55	0.001
LDL, mg/dl	129.01 $\pm$ 26.20	86.05 $\pm$ 17.07	< 0.001
HDL, mg/dl	49.82 $\pm$ 12.32	53.13 $\pm$ 14.16	< 0.001
TC, mg/dl	200.80 $\pm$ 31.75	154.42 $\pm$ 22.97	< 0.001
TG, mg/dl	125.35 $\pm$ 70.41	86.24 $\pm$ 53.72	< 0.001
GGT, U/l	34.13 $\pm$ 47.92	37.86 $\pm$ 68.12	0.265
MetS, n (%)	256 (32.7%)	90 (20.6%)	< 0.001

Values are number (%) or mean $\pm$ SD; Fisher's Exact test for categorical variables and one-way ANOVA for continuous variables.

ALT = Alanine aminotransferase; AST = aspartate aminotransferase; GGT = gamma-glutamyl transferase; LDL-C = low-density lipoprotein-cholesterol; HDL-C = high-density lipoprotein-cholesterol; TC = total cholesterol; TG = triglyceride.

107.41 $\pm$ 32.07 mg/dl; $p = 0.001$), higher SBP (138.12 $\pm$ 19.91 vs. 134.66 $\pm$ 20.97 mm Hg; $p = 0.005$), DBP (82.30 $\pm$ 13.52 vs. 79.84 $\pm$ 14.60 mm Hg; $p = 0.004$), and ApoB-100 content (0.91 $\pm$ 0.24 vs. 0.86 $\pm$ 0.22 g/l; $p < 0.001$).

Nonsignificantly more women had ApoB-100 content $\geq$ and < 0.8 (56.3% and 58.7%; $p = 0.408$) (table 2). Age distribution, BW, BMI, waist size, and WHR were not significantly different. The proportion of metabolic syndrome was significantly higher in patients with ApoB-100 $\geq$ 0.8 (32.7 vs. 20.6%; $p < 0.001$); these patients also had significantly higher serum LDL, total cholesterol, triglycerides, and significantly lower HDL. However, SBP, DBP, pulse pressure, serum creatinine, fasting glucose, and GGT were not significantly different.

The incidences of NAFLD (39.4 vs. 29.4%; $p < 0.001$) and high-grade NAFLD (17.3 vs. 11.0%; $p = 0.003$) were significantly higher in patients with ApoB-100 $\geq$ 0.8 (fig. 2), and the mean value of ApoB-100 significantly increased at higher grades of NAFLD: Grade 1 = 0.86 g/l, Grade 2 = 0.88 g/l, Grade 3 = 0.95 g/l; $p < 0.001$ (fig. 3).

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Fig. 2. Incidence of NAFLD and high-grade hepatosteatosis based on the ApoB-100 level.

Fig. 3. The ApoB-100 levels by NAFLD grade.

Independent Factors of Hepatosteatosis in HCV-Seropositive Patients

All variables were evaluated for predicting NAFLD and high-grade hepatosteatosis. ApoB-100 (OR 5.45; 95% CI 1.64–18.06; $p = 0.006$;) and metabolic syndrome (OR 1.58; 95% CI 1.18–2.11; $p = 0.002$) were independently associated with NAFLD (Grades 1–3) (fig. 4A), and ApoB-100 (OR 7.73; 95% CI 1.64–34.35; $p = 0.006$), metabolic syndrome (OR 1.79; 95% CI 1.24–2.59; $p = 0.002$), and uric acid (OR 1.17; 95% CI 1.05–1.31; $p = 0.006$) were strongly correlated with high-grade NAFLD (grades 2–3) (fig. 4B).

Discussion

This is the first cross-sectional population study that shows a strong association between ApoB-100 and NAFLD in an HCV-seropositive population. The higher ApoB-100 levels indicated a significant risk (OR: 7.73) for high-grade hepatosteatosis.

Fig. 4. **A** Multivariate logistic regression of factors associated with NAFLD. **B** Multivariate logistic regression of factors associated with a high-grade NAFLD. Adjusted for age, gender, WHR, alanine aminotransferase, aspartate aminotransferase, BMI, metabolic syndrome, gamma-glutamyl transferase, HBV, LDL-cholesterol, total cholesterol; uric acid, creatinine, betel-nut chewing, smoking, ApoB-100.

The prevalence of steatosis in patients with hepatitis C (35.8% in all patients) was lower than in another study [6]. A relatively lower incidence of steatosis was thought to be related to limited echogenic screening and non-genotype 3 distributions, according to an HCV line probe assay showing that genotypes 1 and 2 were predominant in Taiwan [24].

Factors Associated with NAFLD in HCV

The prevalence of steatosis in patients with genotype 3 HCV is significantly higher than in patients with other forms of chronic liver disease, e.g., hepatitis B or autoimmune hepatitis, which suggests a direct effect of HCV replication on excess fat accumulation in the liver. The severity of steatosis is associated with the viral load in patients infected with genotype 3 HCV, which subsides with sustained response to antiviral therapy but recurs after a virological relapse. In patients infected with other HCV genotypes, a significant relationship exists between steatosis, elevated BMI and waist circumference, which imply the role of host metabolic factors such as insulin resistance. Our study showed that patients with NAFLD who had similar waist circumference, BW, and BMI also had higher fasting blood glucose, TG, and blood pressure as symptoms of metabolic syndrome ($p = 0.001$). The metabovascular features were still significant in the analyses of various grades of NAFLD. Moreover, the nonsignificant anthropometric measures may be related to the complex mechanisms, including the effect of HCV activity, that contribute to hepatosteatosis. Therefore, underlying dyslipidemia and metabolic disturbance were more essential in the study population;

ApoB-100 might be a more important predictor of NAFLD in people with a similar waist circumference or BW.

NAFLD in chronic HCV infection increases cardiovascular events and extrahepatic mortality through metabolic disturbances and systemic atherosclerosis, which is induced by excessive oxidative stress and advanced necroinflammation. In patients with CHC infection, hypoadiponectinemia increases hepatic steatosis, inflammation, fibrosis, and hepatocarcinogenesis in animal models and human studies [25]. Interestingly, in our study, hyperuricemia was significantly increased in patients with HCV and advanced steatosis (OR 1.172; 95% CI 1.047–1.311; $p = 0.006$), which supported the notion that NAFLD and hyperuricemia are risk markers of hepatic damage, systemic inflammation, and cardiovascular mortality [26]. Taken together, these findings suggest that, in patients with CHC, high-grade NAFLD reflects additional atherogenic risks instead of metabolic syndrome.

Apolipoprotein B-100 on NAFLD

Apolipoproteins, such as ApoE, ApoB, ApoA1, and several ApoC proteins, are associated with the impact of HCV lipo-viro-particles on viral entry, replication, and infection. The ApoB-100 level was relatively lower in genotype-3 HCV patients, and hypocholesteremia was correlated with the virus load [20].

However, in people with non-genotype 3 HCV, hepatosteatosis is primarily related to the host factors of obesity, insulin resistance and DM, and antiviral therapy does not ameliorate steatosis in these patients [26]. Established studies [28–30] showed higher ApoB levels or ApoB/AI ratios associated with fatty liver in patients with DM and in the general population. Greater ApoB expression is associated with insulin resistance [31], and delayed ApoB clearance is associated with triglyceride lipolysis, greater oxidative stress, and greater glycosylation oxidation of lipoprotein particles [32]. In our study, patients with HCV and NAFLD had typical features of metabolic syndrome, and ApoB-100 was strongly associated with NAFLD after traditional metabovascular factors had been adjusted for; the value of ApoB-100 might be more closely associated with advanced hepatosteatosis (OR 7.73; 95% CI 1.64–34.35; $p = 0.006$) in patients with non-genotype 3 HCV.

Limitations

This study has some limitations. First, because of its cross-sectional design, the present findings are inherently limited in their ability to eliminate causal relationships between ApoB-100 and NAFLD. Second, confirmatory HCV testing, e.g., HCV RNA and HCV core antigen testing was not done; thus, we were unable to accurately characterize each patient's carrier status. However, these tests are expensive and infeasible for community health examinations. Third, although a liver biopsy is more sensitive than ultrasonography and is, therefore, the gold standard for this kind of investigation, it is expensive and bears the risks of bleeding and death. Using liver biopsies in population-based studies is unsuitable. Ultrasonography is feasible in a population-based epidemiological study, and interobserver variation can be reduced by experienced operators. Fourth, ApoB-100 was measured only once in the annual examination, and the coefficients of variability are unavailable because the check-up was unrepeatable. Moreover, we did not test for ApoAI, ApoE, or ApoC because the tests are too expensive and give information similar to that provided by tests of ApoB-100 and the ApoB/ApoAI ratio [17]. Finally, some medications or diseases related to different lipid profiles and status of anti-viral therapy were missing despite our careful personal medical survey. Additional large prospective investigations of the mechanism are needed to confirm the correlation.

Conclusion

We found a strong association between ApoB-100 and fatty liver in patients with non-genotype 3 HCV. Despite metabolic syndrome contributing to hepatosteatosis in the study, ApoB-100 is another independent factor for NAFLD. Physicians should pay more attention to patients with HCV and a high ApoB-100 level; they might indicate advanced steatosis, resistance to antiviral therapy, and poor extrahepatic outcomes.

Disclosure Statement

No conflicts of interest to declare.

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