Hepat Mon. 2020 February; 20(2):e98531.

Published online 2020 February 29.

doi: 10.5812/hepatmon.98531.

Research Article

The Incidence and Risk Factors of Non-Alcoholic Fatty Liver Disease: A Cohort Study from Iran

Nima Motamed¹, Mahmoodreza Khonsari², Mahshid Panahi², Nader Rezaie³, Mansooreh Maadi², Fahimeh Safarnezhad Tameshkel², Hossein Keyvani², Mohammad Hadi Karbalaie Niya², Hossein Ajdarkosh², Seyyed Mohammad Miri ¹/₁₀⁴ and Farhad Zamani^{2,*}

¹Department of Social Medicine, Zanjan University of Medical Sciences, Zanjan, Iran

²Gastrointestinal and Liver Disease Research Center, Iran University of Medical Sciences, Tehran, Iran

³Department of Pulmonology, Firouzgar Hospital, Iran University of Medical Sciences, Tehran, Iran

⁴Nedmedica, Heerlen, Netherlands

^{*} *Corresponding author*: Gastrointestinal and Liver Disease Research Center, Iran University of Medical Sciences, Firoozgar Hospital, Tehran, Iran. Tel: +98-9121547311, Email: zamani.f@iums.ac.ir

Received 2019 September 30; Revised 2020 February 09; Accepted 2020 February 14.

Abstract

Background: Non-alcoholic fatty liver disease (NAFLD) is a condition with a global prevalence of 24%. A broad spectrum of liver complications has been attributed to this condition.

Objectives: This study was conducted to determine the incidence of NAFLD and its potential risk factors in a seven-year follow-up study in Iran.

Methods: This is a prospective cohort population-based study conducted in the northern region of Iran. In phase I of the cohort study, 2,461 participants were selected, between 2009 and 2010, using the stratified randomization method based on the sex and age of individuals. Ultrasonographic examination was performed again after a seven-year follow-up between 2016 and 2017. The multiple binary regression analysis was applied to evaluate the association between the development of NAFLD and potential risk factors.

Results: The incidence of NAFLD was 27.88% (95% CI: 25.41% -30.35%) in men and 30.17% (95% CI: 27.40% -32.94%) in women (P=0.226) in a seven-year follow-up period. Based on the multiple binary logistic regression analysis, body mass index (BMI) [OR=1.219 (95% CI: 1.162 - 1.278) P < 0.001], triglyceride (TG) [OR=1.003 (95% CI: 1.001 - 1.005); P=0.005] and high-density lipoprotein (HDL) [OR=0.986 (95% CI: 0.972 - 0.999); P = 0.046] showed a significant association with the incidence of NAFLD in men. In contrast, the marital status [OR=2.141 (95% CI: 1.286 - 3.565); P = 0.003], BMI [OR=1.165 (95% CI: 1.121 - 1.211); P < 0.001] and homeostatic model assessment for insulin resistance (HOMA-IR) [OR=1.164 (95% CI: 1.041 - 1.301); P=0.007] had a significant relationship with the incidence of NAFLD in women.

Conclusions: It seems that NAFLD is markedly rising in the northern part of Iran. Higher levels of BMI, TG, and HDL are considered independent risk factors for the development of NAFLD in men, while the marital status, BMI, and HOMA-IR exhibited independent risk factors with the incidence of NAFLD in women.

Keywords: Non-Alcoholic Fatty Liver Disease (NAFLD), Risk Factor, Iran, Incidence

1. Background

Non-alcoholic fatty liver disease (NAFLD) is characterized by the accumulation of fat in the cytoplasm of hepatocytes in individuals without a history of excessive alcohol consumption, drugs, genetic diseases, or other causes of steatosis (1, 2). Patients with NAFLD can experience a wide range of clinical manifestations from simple steatosis to liver cirrhosis or hepatocellular carcinoma (3). In addition to liver disease, NAFLD correlates with other non-communicable disorders. It has been reported that NAFLD is associated with type 2 diabetes, hyperlipidemia, and metabolic syndrome (3-5). The disease may also lead to an increase in the risk of developing cardiovascular diseases, chronic renal disorder, or even death (6). One study conducted in Iran showed a significant correlation of NAFLD with age, body mass index (BMI), aspartate transaminase (AST), alanine aminotransferase (ALT), triglyceride, cholesterol, low-density lipoprotein (LDL) and fasting blood sugar (FBS) (7). Several studies have indicated that physical activity, socioeconomic status, diabetes, and hypertension are strongly associated with the develop-

Copyright © 2020, Author(s). This is an open-access article distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 International License (http://creativecommons.org/licenses/by-nc/4.0/) which permits copy and redistribute the material just in noncommercial usages, provided the original work is properly cited.

ment of NAFLD (8).

The NAFLD is a common condition with a global prevalence of 24% (9). Approximately 27% - 34% of the American population, and about 25% of the European population are afflicted with NAFLD (10, 11). One study reported that the annual incidence of NAFLD is 6.2% in a community cohort study in adult Sri Lankans who live in urban areas (12). A pooled estimate analysis showed that the incidence of NAFLD in Asia is 52.34 per 1,000 individuals, although only two studies were included in this analysis (9). It is estimated that the prevalence of NAFLD would be increasing in the Middle East as a result of increasing related risk factors (13). In a population-based study performed in the southern part of Iran, the prevalence of NAFLD was estimated at 21.5% (14). In a cohort study carried out in the northern region of Iran, the prevalence of NAFLD in men and women was estimated to be 40.1% and 44.2%, respectively (15).

Globally, there are longitudinal studies about the incidence of fatty liver disease and its association with other risk factors, particularly in developing countries. Thus, the estimation of the prevalence and particularly the incidence rate are of importance.

2. Objectives

This population-based study was conducted in the northern part of Iran to estimate the incidence rate of NAFLD and related risk factors in a seven-year follow-up period.

3. Methods

-2

3.1. Study Population

Considering that the expected annual incidence of NAFLD is equal to 10% as well as a precision of 0.02% - 1%, as the main aim, and expected drop-out rate of 50%, it was estimated that a total of 1728 to 6912 subjects could be recruited to estimate a related proportion based on the below formula (the sample size formula for a single proportion) (16):

$$n = \frac{Z_{1-\alpha/2}^{2} p\left(1-p\right)}{\delta^{2}}$$
(1)

$$\alpha = 0.05,$$

$$p = 10\%,$$

$$\delta = 0.1 \cdot 0.2p = 0.01 \cdot 0.02$$

In this prospective, population-based cohort study, in phase I of the cohort study, 2,461 individuals participated who were without a history of NAFLD. A schematic view of the inclusions and exclusions criteria are displayed in Figures 1 and 2.

The study procedures consisted of two phases; the phase I was performed in Amol: between 2009 and 2010, while the phase II was conducted between 2016 and 2017. Amol is a relatively populated city located in the midnorthern region of Iran. This city belongs to Mazandaran province and has 180 km distance from the capital of Iran, Tehran. Samples were obtained from district health centers related to primary health care settings in urban and rural areas. The primary data used to obtain an exact sampling frame are usually available in these settings. Based on these primary data, we stratified the sampling frame to 16 strata according to the sex and age of groups, including 10 - 19, 20 - 29, 30 - 39, 40 - 49, 50 - 59, 60 - 69, 70 - 79, and 80 - 89 years. The sample size in each stratum was calculated based on the size of individuals in the same stratum. The documented data of the primary healthcare settings were applied to obtain the sampling frame. Between the years 2009 and 2010, to begin phase II of the cohort study in 2016, we were annually in contact with the participants or their family members to provide the associated outcomes. We contacted households by means of telephone calls and cell phones. In the case of the unavailability of patients to respond to our calls, our team referred to where they lived in accordance with the district health centers where the sampling process was carried out. Next, the demographic, anthropometric, and clinical features of the former participants (documented in 2016) were recorded. Thus, former volunteers were actively invited to join in phase II of the cohort study. New cases of individuals with NAFLD were diagnosed by ultrasonographic examinations and a history of risk factors for each patient. Finally, 2,461 subjects who had not any history of NAFLD were included in the present study.

Further details of the study population are shown in Figures 1 and 2. In phase II of the cohort study, the related outcomes were also provided and compared with annually provided outcomes. The related outcomes were modified based on the valid documented data in the case of inconsistency.

3.2. Measurements

The demographic data, including the gender, age, marital status (single, married, and divorced/widowed), and residency status (urban and rural) were recorded upon the registration of participants in both phases of the study. A calibrated non-stretchable meter was utilized to measure the height of participants. The measurement was performed while participants were standing, and their heels and buttocks were in contact with a wall. Their weight was calculated by a calibrated scale with a precision of 100 g. The height and weight values of each subject were assessed after removing the excess cloths and shoes.

Uncorrected Proof



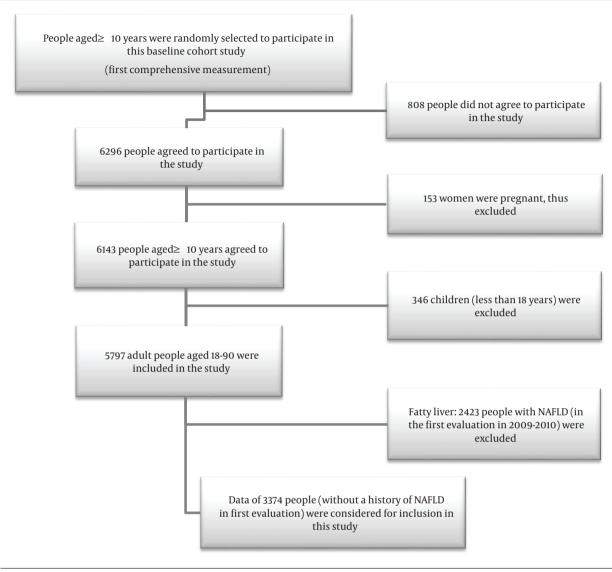


Figure 1. The flow diagram of selecting the individuals at the first step (beginning) of the study

The calibrated monitors with proper cuff sizes, according to the arm circumference of the subjects, were utilized to measure the blood pressure. At least a 5-min rest period was considered for each participant in a sitting position in a quiet room. The systolic and diastolic blood pressure values were determined based on the appearance and disappearance of Korotkof sounds, respectively. The systolic and diastolic blood pressure values were calculated based on the average measurements of systolic and diastolic blood pressure, respectively.

Approximately 10 mL of the whole blood was taken from each individual and placed in a serum separator tube (SST, tiger top tube). Samples were left until the clot appeared. All samples were centrifuged at 3000 rpm for 10 min. Thereafter, the serum was collected using the clean pipette technique and then kept at -80°C. For the separation of plasma samples, the whole blood specimens were poured in EDTA-containing tubes and centrifuged at 2000 rpm at 4°C for 20 min.

3.3. The Diagnosis of NAFLD

Ultrasonographic examinations were applied for the diagnosis of NAFLD in all participants between 2009 and 2010 and performed again between 2016 and 2017 as a follow-up procedure. This disease was defined as evidence of hepatic steatosis if participants had no history of excess

Uncorrected Proof

Motamed N et al.

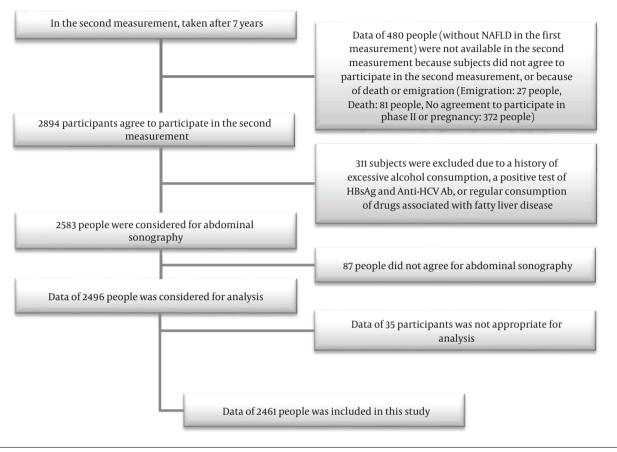


Figure 2. The flow diagram of the participants at the second step (follow-up) after seven years

consumption of alcohol, drug-related steatosis, and a history of hereditary steatogenic hepatic conditions. Ultrasonographic examinations were conducted by an expert radiologist was blinded to the study protocol and not directly involved in the study. The liver parenchyma was radiologically analyzed in different dimensions, including sagittal, longitudinal, lateral, and intercostal views using a 3-5 MHz transducer. The increased hepatic echogenicity, as well as the abnormal appearance of hepatic vessels and diaphragm in ultrasonographic images, was regarded as the required criteria to confirm the diagnosis of fatty liver.

3.4. Statistical Analysis

The outcome measure was defined as the incidence of the disease in a seven-year period (as a percentage), assuming that NAFLD is a common disease. Thus, the incidence rate of NAFLD in this study was calculated according to the below formula:

[(No. of new cases of NAFLD in the population during a 7-year period) / (No. of individuals who were at risk of de-

veloping NAFLD (subjects without NAFLD at the beginning of study))] \times 100

The incidence rate was expressed as the percentage, and the confidence intervals were set at 0.95. Continous values were presented as the means and standard deviations. The parametric values between the two groups were analyzed by the independent *t*-test in the case of the normal distribution of the data, while the non-parametric values between the two groups were analyzed by the Mann-Whitney U test if the data were not normally distributed. The two-group proportion test was utilized to compare the incidence of NAFLD between men and women.

The simple and multiple logistic regression analyses were conducted in which the incidence rate of NAFLD (during a seven-year follow-up period) was considered a dependent variable. In multiple logistic regression analyses, age, residency, marital status, body mass index (BMI), homeostatic model assessment (HOMA-IR), triglyceride (TG), high-density lipoprotein (HDL), low-density lipoprotein (LDL), and the mean arterial blood pressure (MAP) were regarded as the potential predictors. The multicollinearity test between systolic blood pressure (SBP) and diastolic blood pressure (DBP) indicated that these two factors are positively correlated; thus we considered the MAP factor, instead of SBP and DBP, into the logistic regression model. The logistic regression model was separately conducted in both genders. The statistical significance was defined as the P value of being less than 0.05. All of the obtained data were analyzed using the SPSS software version 21.

4. Results

The results demonstrated that the incidence rate of NAFLD was 27.88% (95% CI = 25.41% - 30.35%) in men and 30.17% (95% CI = 27.40% - 32.94%) in women of Iranian population who live in the northern regions of Iran a seven-year follow-up period (P = 0.2263).

Table 1 shows the mean values of the basic characteristics of the population study at (in phase I) in individuals with NAFLD and without NAFLD after a seven-year followup. Although the mean age was significantly higher in women with NAFLD than those without NAFLD after seven years, there was no significant difference in the mean age between men with NAFLD and those without NAFLD. Moreover, BMI, MAP, DBP, SBP, FBS, TG, total cholesterol, LDL and HOMA-IR were significantly higher in men and women who are newly diagnosed with NAFLD (individuals with NAFLD after a seven-year follow-up period), while HDL-C was significantly higher in subjects without NAFLD after a seven-year follow-up period than the other groups.

Table 2 shows the results of the simple and multiple binary logistic regression analyses in which the incidence rate of NAFLD determines the outcome. The age, residency, marital status, BMI, HOMA-IR, TG, HDL, LDL, and MAP were regarded as the potential predictors. The analyses were performed separately in both genders. In the simple analysis method, all predictors mentioned above had a significant association with the incidence of NAFLD after a sevenyear follow-up period in women. In men, except for the age, residency, marital status, and other potential predictors showed a significant relationship with the incidence of NAFLD. In the multiple binary logistic regression analysis, BMI, TG, and HDL exhibited a significant correlation with the incidence of NAFLD in men. Moreover, marital status, BMI, and HOMA-IRhad a significant relationship with the incidence of NAFLD in women.

5. Discussion

In this prospective population-based study, the incidence of NAFLD was estimated in a seven-year follow-up peTable 1. The Mean Values of the Basic Characteristics of Individuals at Baseline (in the Phase I of the Cohort Study)^a

Variables	People with NAFLD After 7 Years, Mean \pm SD	People Without NAFLD After 7 Years, Mean ± SD	P Value
Male			
Age	40.68 ± 14.84	42.52 ± 16.99	0.161
BMI	26.27 ± 3.58	23.76 ± 3.39	< 0.001
MAP	88.74 ± 11.85	86.43 ± 11.68	0.001
DBP	75.47 ± 12.10	73.06 ± 11.90	0.005
SBP	115.26 ± 13.57	113.18 ± 13.98	< 0.001
FBS	93.27 ± 15.71	94.09 ± 26.71	0.002
TG	$\textbf{152.44} \pm \textbf{99.93}$	116.40 ± 69.16	< 0.001
Cholesterol	177.88 ± 39.61	170.73 ± 40.35	< 0.001
HDL	42.41 ± 11.46	46.73 ± 11.55	< 0.001
LDL	104.59 ± 29.92	99.60 ± 30.21	< 0.001
HOMA	2.09 ± 1.33	1.62 ± 1.46	< 0.001
Female			
Age	41.07 ± 11.90	35.73 ± 14.39	< 0.001
BMI	29.65 ± 4.30	26.11 ± 4.66	< 0.001
MAP	86.97 ± 13.27	83.93 ± 11.57	0.001
DBP	74.02 ± 12.58	$\textbf{71.48} \pm \textbf{11.93}$	0.005
SBP	112.87 ± 17.07	108.83 ± 14.41	< 0.001
FBS	95.92 ± 28.99	92.77 ± 25.04	0.002
TG	135.46 ± 85.98	106.69 ± 56.36	< 0.001
Cholesterol	188.76 ± 42.66	176.30 ± 40.40	< 0.001
HDL	46.56 ± 12.03	49.67 ± 11.60	< 0.001
LDL	110.60 \pm 30.77	101.72 ± 30.50	< 0.001
HOMA	2.56 ± 1.97	1.96 ± 1.21	< 0.001

Abbreviations: BMI, body mass index; HDL, high-density lipoprotein; HOMA-IR, homeostatic model assessment for insulin resistance; LDL, low-density lipoprotein; MAP, mean arterial blood pressure; TG, triglyceride

^a Individuals include newly diagnosed NAFLD and healthy subjects without NAFLD after a seven-year follow-up period

riod in individuals without a history of NAFLD. Thus, individuals with NAFLD in phase I of the cohort study were excluded, and then the incidence of NAFLD during a one-year follow-up period was estimated in healthy participants. The present study showed that the incidence of NAFLD was 27.88% (25.41% - 30.35%) in men and 30.17% (27.40% -32.94%) in women. These results indicate that the annual rate of NAFLD is nearly 4% in men and 4.3% in women. Kanwal et al. (17) reported a 2% - 3% annual incidence of NAFLD in American veterans. In a study conducted by Zelber-Sagi and colleagues (18), they reported a 20% incidence of NAFLD in a seven-year follow-up period (similar to the follow-up period of our study) with an annual in-

Motamed N et al.

Variables	Simple Binary Logistic Regression Analysis			Multiple Binary Logistic Regression Analysis		
	Wald	OR (95% CI)	P Value	Wald	OR (95% CI)	P Value
Male						
Age	3.16	0.993 (0.986 - 1.001)	0.075	9.76	0.980 (0.968 - 0.993)	0.002
Residency ^a	0.327	0.930 (0.727 - 1.191)	0.567	0.549	0.893 (0.661 - 1.205)	0.459
Marriage status ^b	1.318	1.188 (0.885 - 1.594)	0.251	0.836	1.243 (0.780 - 1.982)	0.361
BMI	111.62	1.239 (1.191 - 1.289)	< 0.001	65.36	1.219 (1.162 - 1.278)	< 0.001
HOMA-IR	23.73	1.249 (1.142 - 1.365)	< 0.001	0.970	1.051 (0.952 - 1.160)	0.325
TG	39.29	1.005 (1.004 - 1.007)	< 0.001	8.00	1.003 (1.001 - 1.005)	0.005
HDL	32.89	0.967 (0.957 - 0.978)	< 0.001	3.92	0.986 (0.972 - 0.999)	0.046
LDL	6.68	1.005 (1.001 - 1.010)	0.010	2.25	1.004 (0.999 - 1.009)	0.134
MAP	9.63	1.016 (1.006 - 1.027)	0.002	0.712	0.995 (0.982 - 1.007)	0.399
emale						
Age	31.80	1.028 (1.018 - 1.037)	< 0.001	2.08	1.010 (0.997 - 1.023)	0.150
Residency ^a	13.46	1.641 (1.260 - 2.139)	< 0.001	2.51	1.308 (0.938 - 1.823)	0.113
Marriage status ^b	35.06	3.717 (2.407 - 5.739)	< 0.001	8.56	2.141 (1.286 - 3.565)	0.003
BMI	104.49	1.184 (1.146 - 1.223)	< 0.001	61.32	1.165 (1.121 - 1.211)	< 0.00
HOMA-IR	31.55	1.302 (1.187 - 1.428)	< 0.001	7.15	1.164 (1.041 - 1.301)	0.007
TG	31.27	1.006 (1.004 - 1.008)	< 0.001	2.86	1.002 (1.000 - 1.005)	0.089
HDL	14.70	0.977 (0.966 - 0.989)	< 0.001	1.17	0.992 (0.978 - 1.007)	0.280
LDL	17.39	1.009 (1.005 - 1.014)	< 0.001	0.433	0.998 (0.992 - 1.003)	0.433
MAP	13.47	1.020 (1.009 - 1.031)	< 0.001	0.832	1.006 (0.993 - 1.020)	0.362

Table 2. The Results of Simple and Multiple Binary Logistic Regression Analyses with Respect to the Incidence of NAFLD After a Seven-Year Follow-Up as the Outcome

Abbreviations: BMI, body mass index; HDL, high-density lipoprotein; HOMA-IR, homeostatic model assessment for insulin resistance; LDL, low-density lipoprotein; MAP, mean arterial blood pressure; TG, triglyceride

^aThe ratio of rural residency vs. urban residency

^bThe ratio of marital status to being single

cidence rate of 2.86%. Another study performed in Hong-Kong reported that 13.5% of new cases were afflicted with NAFLD within a period of 3 - 5 years (19). Two studies carried outin the Chinese population reported 43.4% and 36.7% incidence rates of NAFLD after seven-and six-year follow-up periods, respectively (20, 21). These results show that the incidence of NAFLD is markedly increasing in countries that previously had a low prevalence of NAFLD. This phenomenon may be due to the changes in the lifestyle of the people.

In the present study, the potential risk factors, including, age, residency, marital status, blood pressure, FBS, LDL, HDL, TG, and HOMA-IR were investigated in men and women with NAFLD to analyze the association of these factors with the disease development. In a simple analysis method, all potential predictors had a significant correlation with the incidence of NAFLD in a seven-year followup period in women. In men, except for the age, residency, and marital status, as well as other potential predictors, showed a significant relationship with the incidence of NAFLD. In adjusted models, BMI, TG, and HDL exhibited a significant association with the incidence of NAFLD in men, while the marital status, BMI, and HOMA-IR had a significant relationship with the incidence of NAFLD in women. Of note, the existing associations did not statistically seem strong, as one unit (1 mg/dL) increase in the concentration of TG leads to a 0.3% increase in the risk of developing NAFLD. Among the evaluated potential predictors, obesity had a strong association with NAFLD. Based on previous studies, an increase in BMI can lead to an increase in the prevalence and incidence of NAFLD (18, 22).

As mentioned above, marital status had a significant correlation with the incidence of NAFLD in female patients without a history of NAFLD. It is worth mentioning that married women may have a higher age and pregnancy rates that can cause pregnancy-associated diabetes, leading to higher abdominal obesity. All factors mentioned earlier could be considered the risk factors of NAFLD. Although the age was not independently associated with NAFLD in women, higher age can influence the development of NAFLD as a result of other factors, including insulin resistance, abdominal obesity, physical inactivity, and other potential contributing risk factors. Furthermore, other potential mediators, in the correlation of marital status with NAFLD in women, can also play a role; however, these potential factors were not evaluated in this study. The lifestyle of female patients can vary since, upon the marriage, they mostly devote themselves to perform housekeeping activities. Notably, being "in shape" might be less priority to married women compared with single women according to Iranian culture. An imbalance between the intake of high-calorie foods and energy expenditure may also be involved in this regard (12, 23).

Overall, the present study showed that the annual rate of NAFLD is estimated to be approximately 4% in men and 4.3% in women. This estimation is statistically higher than those of some investigations performed in Western populations (11), and it may have some confounding factors to be considered. Our study was limited by some restrictions, as the time of NAFLD onset was not recorded, and thus we were not able to analyze the survival models, such as the Cox proportional hazard model; however, predicting the precise time of NAFLD onset would not be easily feasible in a population-based study. The annual evaluation of patients with NAFLD by ultrasonographic examinations would be valuable. The application of this technique for the diagnosis of NAFLD was another limitation of this study. This diagnostic method is not capable of distinguishing between the changes in adipocytes of the liver and fibrosis. NAFLD has a wide spectrum of clinical manifestations from simple steatosis to liver fibrosis. Although the liver biopsy is a gold-standard method for the diagnosis of NAFLD, this procedure is invasive and cannot be considered the best diagnostic tool in population-based studies. The evaluation of alcohol consumption as a self-report variable was another limitation of the study due to social desirability bias. However, utilizing an anonymous questionnaire would be helpful in overcoming this obstacle. The inability to consider other socioeconomic factors may be another limitation of this study, as it would not be possible to collect truthful information about some variables, especially in developing countries. Household income and educational status could provide valuable information.

In conclusion, it seems that NAFLD is increasingly rising in this area of the Middle East. The potential risk factors for the incidence of NAFLD were not the same in both genders. Among the evaluated potential risk factors in men and women, only obesity was an independent risk factor for the incidence of NAFLD. Moreover, BMI, TG, and HDL had a significant independent relationship with the incidence of NAFLD in men, while marital status, BMI, and HOMA-IR exhibited an independent correlation with the incidence of the disease in women.

Acknowledgments

All authors acknowledge the financial support of the Iran University of Medical Sciences, Tehran, Iran.

Footnotes

Authors' Contribution: Study concept and design: Farhad Zamani and Nima Motamed; analysis and interpretation of data: Seyyed Mohammad Miri, Hossein Keyvani, and Nader Rezaie; drafting of the manuscript: Mahshid Panahi and Hossein Ajdarkosh; critical revision of the manuscript for important intellectual content: Mohammad Hadi Karbalaie Niya and Fahimeh Safarnezhad Tameshkel; statistical analysis: Mansooreh Maadi and Mahmoodreza Khonsari.

Conflict of Interests: There is no conflict of interest.

Ethical Approval: The protocol of the study was approved by the Ethics Committee of the Iran University of Medical Sciences, Tehran, Iran (IR.IUMS.REC.1398.852).

Funding/Support: The study was financially supported by a research grant (98-3-30-16276).

Informed Consent: Written informed consent was obtained from all individuals before the initiation of the study. The information related to the participants remained confidential and secured.

References

- Angulo P, Lindor KD. Non-alcoholic fatty liver disease. J Gastroenterol Hepatol. 2002;17 Suppl: S186–90. doi: 10.1046/j.1440-1746.17.51.10.x. [PubMed: 12000605].
- Neuschwander-Tetri BA, Caldwell SH. Nonalcoholic steatohepatitis: Summary of an AASLD Single Topic Conference. *Hepatology*. 2003;37(5):1202–19. doi: 10.1053/jhep.2003.50193. [PubMed: 12717402].
- Hafeez S, Ahmed MH. Bariatric surgery as potential treatment for nonalcoholic fatty liver disease: A future treatment by choice or by chance? J Obes. 2013;2013:839275. doi: 10.1155/2013/839275. [PubMed: 23431426]. [PubMed Central: PMC3569911].
- Kim HJ, Kim HJ, Lee KE, Kim DJ, Kim SK, Ahn CW, et al. Metabolic significance of nonalcoholic fatty liver disease in nonobese, nondiabetic adults. *Arch Intern Med.* 2004;**164**(19):2169–75. doi: 10.1001/archinte.164.19.2169. [PubMed: 15505132].
- Bedogni G, Miglioli L, Masutti F, Tiribelli C, Marchesini G, Bellentani S. Prevalence of and risk factors for nonalcoholic fatty liver disease: The Dionysos nutrition and liver study. *Hepatology*. 2005;42(1):44–52. doi: 10.1002/hep.20734. [PubMed: 15895401].
- Musso G, Gambino R, Tabibian JH, Ekstedt M, Kechagias S, Hamaguchi M, et al. Association of non-alcoholic fatty liver disease with chronic kidney disease: A systematic review and meta-analysis. *PLoS Med.* 2014;11(7). e1001680. doi: 10.1371/journal.pmed.1001680. [PubMed: 25050550]. [PubMed Central: PMC4106719].

- Adibi A, Maleki S, Adibi P, Etminani R, Hovsepian S. Prevalence of nonalcoholic fatty liver disease and its related metabolic risk factors in Isfahan, Iran. *Adv Biomed Res.* 2017;6:47. doi:10.4103/2277-9175.204590. [PubMed: 28503502]. [PubMed Central: PMC5414408].
- Zarean E, Goujani R, Rahimian G, Ahamdi A. Prevalence and risk factors of non-alcoholic fatty liver disease in southwest Iran: A population-based case-control study. *Clin Exp Hepatol.* 2019;5(3):224– 31. doi: 10.5114/ceh.2019.87635. [PubMed: 31598559]. [PubMed Central: PMC6781826].
- Younossi ZM, Koenig AB, Abdelatif D, Fazel Y, Henry L, Wymer M. Global epidemiology of nonalcoholic fatty liver disease-Meta-analytic assessment of prevalence, incidence, and outcomes. *Hepatology*. 2016;64(1):73-84. doi: 10.1002/hep.28431. [PubMed: 26707365].
- Vernon G, Baranova A, Younossi ZM. Systematic review: The epidemiology and natural history of non-alcoholic fatty liver disease and non-alcoholic steatohepatitis in adults. *Aliment Pharmacol Ther.* 2011;34(3):274–85. doi: 10.1111/j.1365-2036.2011.04724.x. [PubMed: 21623852].
- Bedogni G, Miglioli L, Masutti F, Castiglione A, Croce LS, Tiribelli C, et al. Incidence and natural course of fatty liver in the general population: The Dionysos study. *Hepatology*. 2007;46(5):1387–91. doi: 10.1002/hep.21827. [PubMed: 17685472].
- Niriella MA, Pathmeswaran A, De Silva ST, Kasturiratna A, Perera R, Subasinghe CE, et al. Incidence and risk factors for non-alcoholic fatty liver disease: A 7-year follow-up study among urban, adult Sri Lankans. *Liver Int.* 2017;**37**(11):1715–22. doi: 10.1111/liv.13478. [PubMed: 28544258].
- Ahmed MH, Noor SK, Bushara SO, Husain NE, Elmadhoun WM, Ginawi IA, et al. Non-alcoholic fatty liver disease in Africa and Middle East: An attempt to predict the present and future implications on the healthcare system. *Gastroenterology Res.* 2017;**10**(5):271– 9. doi: 10.14740/gr913w. [PubMed: 29118867]. [PubMed Central: PMC5667692].
- Lankarani KB, Ghaffarpasand F, Mahmoodi M, Lotfi M, Zamiri N, Heydari ST, et al. Non alcoholic fatty liver disease in southern Iran: A population based study. *Hepat Mon.* 2013;13(5). e9248. doi: 10.5812/hepatmon.9248. [PubMed: 23922564]. [PubMed Central: PMC3734894].
- 15. Motamed N, Sohrabi M, Ajdarkosh H, Hemmasi G, Maadi M, Sayeedian FS, et al. Fatty liver index vs waist circumference for predicting non-

alcoholic fatty liver disease. *World J Gastroenterol*. 2016;**22**(10):3023-30. doi: 10.3748/wjg.v22.i10.3023. [PubMed: 26973398]. [PubMed Central: PMC4779925].

- Ong JP, Pitts A, Younossi ZM. Increased overall mortality and liver-related mortality in non-alcoholic fatty liver disease. J Hepatol. 2008;49(4):608–12. doi: 10.1016/j.jhep.2008.06.018. [PubMed: 18682312].
- Kanwal F, Kramer JR, Duan Z, Yu X, White D, El-Serag HB. Trends in the burden of nonalcoholic fatty liver disease in a United States Cohort of Veterans. *Clin Gastroenterol Hepatol.* 2016;14(2):301-8 e1-2. doi: 10.1016/j.cgh.2015.08.010. [PubMed: 26291667]. [PubMed Central: PMC4918764].
- Zelber-Sagi S, Lotan R, Shlomai A, Webb M, Harrari G, Buch A, et al. Predictors for incidence and remission of NAFLD in the general population during a seven-year prospective follow-up. *J Hepatol.* 2012;56(5):1145–51. doi: 10.1016/j.jhep.2011.12.011. [PubMed: 22245895].
- Wong RJ, Ahmed A. Obesity and non-alcoholic fatty liver disease: Disparate associations among Asian populations. *World J Hepatol.* 2014;6(5):263-73. doi: 10.4254/wjh.v6.i5.263. [PubMed: 24868320]. [PubMed Central: PMC4033284].
- Toshimitsu K, Matsuura B, Ohkubo I, Niiya T, Furukawa S, Hiasa Y, et al. Dietary habits and nutrient intake in non-alcoholic steatohepatitis. *Nutrition*. 2007;23(1):46–52. doi: 10.1016/j.nut.2006.09.004. [PubMed: 17140767].
- Jin YJ, Kim KM, Hwang S, Lee SG, Ha TY, Song GW, et al. Exercise and diet modification in non-obese non-alcoholic fatty liver disease: Analysis of biopsies of living liver donors. J Gastroenterol Hepatol. 2012;27(8):1341–7. doi: 10.1111/j.1440-1746.2012.07165.x. [PubMed: 22554085].
- Wong VW, Wong GL, Yeung DK, Lau TK, Chan CK, Chim AM, et al. Incidence of non-alcoholic fatty liver disease in Hong Kong: A population study with paired proton-magnetic resonance spectroscopy. *J Hepatol.* 2015;62(1):182–9. doi: 10.1016/j.jhep.2014.08.041. [PubMed: 25195550].
- Wu J, He S, Xu H, Chi X, Sun J, Wang X, et al. Non-alcoholic fatty liver disease incidence, remission and risk factors among a general Chinese population with a 6-year follow-up. *Sci Rep.* 2018;8(1):7557. doi: 10.1038/s41598-018-25641-z. [PubMed: 29765064]. [PubMed Central: PMC5954048].