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# Clinical and histological features of nonalcoholic steatohepatitis in Iranian patients

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## Abstract

**Background:** Although several studies have been performed on risk factors and natural course of NASH, it seems that NASH tends to be more than a disease confined to strict boundaries. The objective of this study was to assess the clinical and paraclinical features and risk factors for non-alcoholic steatohepatitis (NASH) patients in an Iranian population

**Methods:** Patients with histologically confirmed NASH who had elevated liver aminotransaminases, negative serologic markers of viral or autoimmune hepatitis and no findings in favor of metabolic liver disease were enrolled. A careful history was taken regarding alcohol intake.

**Results:** 53 patients consisting of 32 male and 21 female entered the study. The mean age was  $37.8 \pm 11.3$  years. Twenty-six patients (55.3%) were overweight, 15 (31.9%) obese, 40 (75.5%) dyslipidemic, and three patients (5.7%) were diabetic. Liver biopsy showed mild steatosis in 35.7%, moderate steatosis in 53.6%, and severe forms in 10.7%. In 80.2% of patients, portal inflammation was present, and 9.4% had cirrhosis. The amount of increase in liver enzymes bore no relationship with fibrosis, portal inflammation, and degree of steatosis.

**Conclusions:** The patients in our study showed a male predominancy and were somewhat younger than other studies.

## Background

*Non-alcoholic steatohepatitis* (NASH) is a disease of unknown origin characterized histologically by changes similar to what is seen in alcoholic-like liver injury but in the absence of significant alcohol intake [1,2]. NASH is considered a type of chronic hepatitis and is a severe form of a spectrum called *non-alcoholic fatty liver disease*

(NAFLD) [3]. NAFLD has four histological stages: (1) fatty infiltration of the liver (2) fatty infiltration plus inflammation (3) fatty infiltration with ballooning degeneration (4) fatty infiltration with lesions similar to alcoholic hepatitis and sinusoidal fibrosis, polymorphonuclear infiltration with or without Mallory hyaline. NASH is the name given to the third and fourth stages [4,5].

The prevalence and clinical significance of NASH appears to be more than what is generally believed [6–11]. Among patients who have had liver biopsies, NASH is seen in approximately 7 to 9 percent in Western countries [12,13]. Ultrasonographic studies on normal population show a prevalence of 25 percent for NAFLD in the United States [5,14]. The disease predominantly occurs between the ages of 40 and 60, [9,11,15] although there have been reports in children over the age of 10 [16,17] NASH is frequently associated with disorders such as insulin resistance, obesity, type 2 diabetes mellitus, hyperlipidemia, protein-calorie malnutrition, and jejeunoileal bypass surgery [5,9,12,18–21] It is important to consider that NASH can progress to cirrhosis and hepatic failure [7,8], further underscoring its importance as a potentially serious and life-threatening disease.

Regarding this fact that in Iran alcohol consumption is much less than western countries, we doubted whether the demography and risk factors of NASH in Iranian populations is the same as what has been seen in previous studies.

## Methods

### Patient selection

The cohort of patients reviewed here were selected from all patients between 15 and 65, referred to Gastroenterology and Hepatology Clinic of Imam Khomeini Hospital, Tehran, Iran, from March 20, 1999 to November 21, 2001. Inclusion criteria were: 1. elevated liver aminotransaminases 2. histologic confirmation of nonalcoholic steatohepatitis 3. negative serologic markers of viral or autoimmune hepatitis (including HBsAg, HCV antibody (ELISA), HIV antibody (ELISA), antinuclear antibodies, anti-smooth muscle antibodies, and anti-liver/kidney microsomes type 1 antibodies) 4. no findings in favor of metabolic liver diseases, including Wilson's disease and hemochromatosis) and negative alpha-1 antitrypsin. A careful history was taken with special attention to alcohol intake and ultrasonography was performed in all enrolled patients. Alcohol intake was asked from the patient and at least one of the patients' companions. In case of positive alcohol consumption, the questionnaire used for this purpose had more detail to help the interviewer to estimate the approximate amount of alcohol consumption. Exclusion criteria were: 1. recent gastrointestinal surgery 2. pregnancy 3. usage of drugs known to result in steatosis, including glucocorticoids, synthetic estrogens, aspirin, tamoxifen, amiodarone, Calcium-channel blockers, and methotrexate.

Of 213 patients (108 male and 105 female) who have primarily been studied for elevation of serum aminotransferases in the beginning, 72 patients who had inclusion criteria, except pathologic confirmation, consented to

undergo liver biopsy. The pathologic diagnosis of NASH was confirmed in 53 patients and these patients were analyzed. Also we used the demographic data of the other 21 patients to avoid any selection bias in this characteristics, which sounded to be different in our study, comparing with previous studies.

### Laboratory and pathological studies

Levels of aminotransferases, alkaline phosphatase, total bilirubin, serum cholesterol and triglycerides, fasting blood sugar, and prothrombin time were measured using standard techniques. The definite diagnosis of NASH was based on a histology showing moderate to gross macrovesicular fatty change with inflammation (lobular or portal), with or without Mallory bodies, fibrosis, or cirrhosis. The grading and staging of all biopsy specimens were determined based on the method proposed by Brunt et al. [2] All patients who had mild steatosis in their biopsy were re-checked for HCV infection using radioimmunoabsorbant assay (RIBA), which was negative in all. Because of financial limitation we did not recheck those with higher degrees of steatosis, since the diagnosis of NASH was more probable in them. A single radiologist performed all ultrasonographic studies and all pathologic studies were performed and reported by one pathologist. Overweight was defined as a BMI between 25 and 29.9 kg/m<sup>2</sup>, and obesity as BMI equal or above 30 kg/m<sup>2</sup>. Ideal weight for height was calculated by Hamwi Method [22] (Men: 48.18 kg for 150 cm +/- 1.1 kg per cm over/under 150 cm; Women: 45.45 kg for 150 cm +/- 0.91 kg per cm over/under 150 cm). Diabetes mellitus was defined as either 1) fasting blood glucose concentration of 126 mg/dL or higher in two occasions, or 2) a random value of 200 mg/dL or higher, confirmed on another occasion, or 3) having a confirmed history of diabetes mellitus for which the patient is currently receiving antidiabetic drugs. Dyslipidemia and insulin resistance was defined according to guidelines of ATP III. [23] Thus, patients with one of the criteria of LDL-C ≥ 160 mg/dL, total cholesterol ≥ 200 mg/dL, triglycerides ≥ 150 mg/dL, or HDL-C < 40 mg/dL were considered dyslipidemic. Insulin resistance was defined with presence of three of the following:

1. Abdominal obesity, defined as a waist circumference in men >102 cm (40 in) and in women >88 cm (35 in)
2. Triglycerides ≥ 150 mg/dL
3. HDL cholesterol < 40 mg/dL in men and < 50 mg/dL in women
4. Blood pressure ≥ 130/ ≥ 85 mmHg
5. Fasting glucose ≥ 110 mg/dL

**Table 1: Anthropometric and biochemical characteristics of 53 patients with NASH**

	Male	Female	Total
Number	32	21	53
Age (y)	36.0 ± 10.6	40.5 ± 11.9	37.8 ± 11.3
Weight (kg)	93.6 ± 16.5*	69.3 ± 12.0	83.8 ± 19
BMI (kg/m <sup>2</sup> )	30.7 ± 4.9†	27.3 ± 3.5	29.3 ± 4.7
Normal BMI, n (%)	7 (21.9%)	5 (23.8%)	12 (22.6%)
Obesity, n (%)	13 (46.4)*	2 (10.5)	15 (31.9)
Overweight, n (%)	12 (42.9)*	14 (73.7)	26 (55.3)
Total cholesterol (mg/dL)	229.6 ± 73.5	223.5 ± 62.5	227.0 ± 68.5
LDL-C (mg/dL)	139.5 ± 42.3	123.0 ± 43.3	134.8 ± 42.2
HDL-C (mg/dL)	47.6 ± 16.1	43.1 ± 13.6	46.3 ± 15.2
Triglycerides (mg/dL)	207 ± 83.6	184 ± 89.1	197.3 ± 85.8
FBS (mg/dL)	102.1 ± 11.4	112.8 ± 42.5	106.5 ± 28.1
ALT (IU/L)	80.3 ± 27.5	82.0 ± 38.8	80.9 ± 32.0
AST (IU/L)	68.8 ± 22.4	71.3 ± 37.5	69.8 ± 28.9
AST/ALT	0.9 ± 0.3	0.9 ± 0.5	0.9 ± 0.4

\*p < 0.001, †p < 0.05 compared to female patients. ALT: Alanine Aminotransferase, AST: Aspartate Aminotransferase, BMI: Body Mass Index, FBS: Fasting Blood Sugar, HDL-C: High Density Lipoprotein Cholesterol, LDL-C: Low Density Lipoprotein Cholesterol.

Risk factors for NASH, according to previous studies, included diabetes mellitus, hyperlipidemia, obesity, history of significant weight loss or weight gain, history of consumption of estrogens or androgens, and extensive abdominal surgery.

#### Statistical analysis

The results of quantitative variables are presented as mean ± SD and those of qualitative variables as numbers and percentages. T test was used to compare quantitative variables between subgroups of patients, and differences between categorical variables were analyzed by Chi Square. Logistic regression analysis was used to assess the effect of different factors on histological findings in liver biopsy specimens. A significance level of 0.05 was used.

#### Results

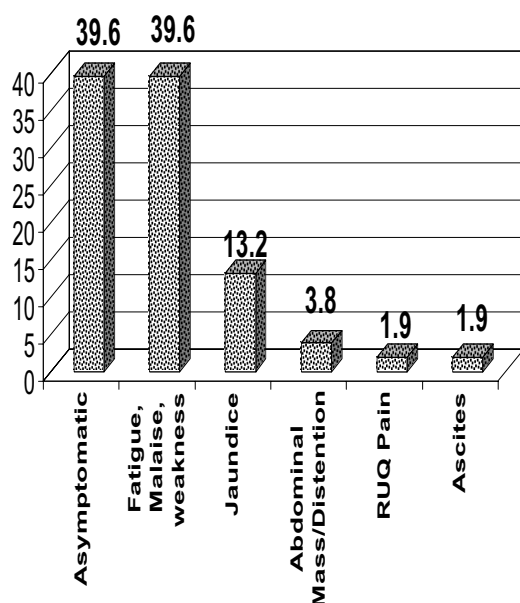
Of the 53 patients whose diagnosis was confirmed by biopsy, 32 (60.4%) were male and 21 (39.6%) were female with the mean age of 37.8 ± 11.3 years. Mean weight of the patients was 83.8 ± 19 kg and mean BMI was 29.3 ± 4.7 kg/m<sup>2</sup>. Twenty-six patients (55.3%) were overweight and 15 (31.9%) were obese (Table 1). None of the patients had weights lower than the "ideal weight for height". On the average, the patients had weights 36.3 ± 20.5% (ranging from 0 to 125.2%) higher than the ideal weight and 94.3% of the patients had weights more than 10% higher than the ideal weight for their height (the mean ideal weight for height of the patients was 61.4 ± 9.6 kg). Six patients (11.3%) had a history of considerable weight loss and 12 patients (22.7%) reported an increase in weight during the last year.

Three patients (5.7%) had overt diabetes mellitus. Forty patients (75.5%) had some form of dyslipidemia; in 66.0% serum cholesterol was high, in 60.4% hypertriglyceridemia was present, and both cholesterol and triglycerides were high in 50.9%. Insulin resistance was observed in 29 patients (54.7%). Seven patients (13.2%) had none of the risk factors usually associated with NASH. The mean ALT and AST levels were 80.9 ± 32.0 IU/L and 69.8 ± 28.9 IU/L, respectively. Mean AST to ALT ratio was 0.9 ± 0.4. This ratio was below 1 in 65.3% of patients and below 2 in 96.2%.

In 31 patients (58.5%) abnormal ultrasound findings were observed; 17% of all patients had increased liver size, and 50.9% had hyperechogenicity, indicating fatty liver.

79.2% of patients had no or very unspecific clinical manifestations, like fatigue, malaise and weakness. Different clinical manifestations are shown in Figure 1.

Liver biopsy showed that steatosis was mild in 35.7%, moderate in 53.6%, and severe in 10.7%. 22 patients (41.5%) had mild, 26 patients (49.1%) moderate and 5 patients (9.4%) had severe necroinflammatory grades. In histological staging of fibrosis, 11.3%, 18.9%, 22.6%, and 37.7% of patients had 0 through 3 scores, respectively and in 9.4% (5 patients), liver biopsy demonstrated cirrhosis (stage 4). The results of logistic regression analysis showed no relationship between the level of liver enzymes and fibrosis, necroinflammatory grading, and steatosis. Presence of insulin resistance did not have any significant relationship with the staging and grading of the disease, as well.



**Figure 1**  
Relative Frequency (Percent) of Different Clinical Manifestations in Patients with NASH, Tehran, 2002

In an analysis of demographic characteristics of 53 histologically-confirmed NASH cases and 21 patients with clinical diagnosis of NASH whose diagnosis was not confirmed in biopsy, 56.8% of patients (42/74) were male and 43.2% (32/74) were female. Mean age was  $39.2 \pm 12.1$  years and mean BMI was  $29.1 \pm 4.1$  kg/m<sup>2</sup>. There was no significant difference in these characteristics between those who had positive biopsy and those who had negative.

## Discussion

Nonalcoholic fatty liver disease (NAFLD) seems to have become an important medical entity and this importance is mainly resulted from its potential to progress to cirrhosis and liver failure and its common occurrence in general population [6]. Because of the low rate of alcohol consumption in Iran, the prevalence might be even higher than other countries. Regardless of the religious beliefs, the habit of alcohol consumption in Iranian population, in most instances, is not so heavy and frequent as what is seen in most of the western countries.

Our patients were somewhat younger than those in previous studies (40–60 years old) [9,12,15]. Also most of our patients were male (60.4%), while most of the previous series showed a female predominance [10–12,19] and only a few had more male patients than females [9]. One of the reasons for these differences might be the failure to

rigorously rule out hepatitis C in some of the studies. We rechecked all patients with mild steatosis for hepatitis C infection. In the report mentioned before as showing a male preponderance [9], also hepatitis C was excluded with precision, and the patients had demographic characteristics different from those usually reported. The male predominance in our study, prompted us to evaluate whether this difference is related to the disease, itself, or is a selection bias in patients who are referred to our clinic. We compared these patients with other patients who primarily had elevation of serum aminotransferases but did not meet the inclusion criteria of the study. From total number of 127 patients with aminotransferases elevation who were excluded from the study, 52.8% were male. Although not exclusively rejecting the selection bias, these findings are against the assumption of a selection bias, since there is not a significant male predominance in other patients of our clinic. But more investigations are mandatory in this regard. Another probable justification of the different sex composition of our patients might be attributed to the lower rate of alcohol consumption in our society. Since men usually consume more amount of alcohol and are more at risk of alcoholic liver disease, in societies with higher prevalence of alcohol consumption, more men are excluded from the diagnosis of nonalcoholic steatohepatitis because of their alcohol consumption.

The mean weight of the patients was higher than normal as in previous reports. [11,12,18] Most patients had abnormally high BMI's and 94.2% had weights more than 10% above their ideal body weights. 11.3% of the patients had a history of weight loss in the year before diagnosis, which is similar to other results [24,25].

Another difference between our results and other reports is the relatively low prevalence of diabetes. Although some reports have shown a prevalence of 2–5% for diabetes mellitus in this group [19,26], in most of them this prevalence is as high as 21–55% [12,9–11,18–21]. An interesting finding is that 13.2% of our patients had none of the known risk factors for development of NASH, and NASH should be considered as an entity not being confined to obese and diabetic patients.

Liver enzyme levels in our patients were similar to previous reports [9]. The mean AST to ALT ratio was less than 1 as expected, and in 96.2% this ratio was below 2. In alcoholic liver disease, this ratio is usually above two, averaging from 2.6 to 2.85 [24,27,28].

We observed dyslipidemia in 75.5% of our patients. Others have reported similar prevalences (20–80%) [9–12,20,21]. The ultrasonographic findings, such as fatty infiltration, are usually nonspecific [29].

It is not usually advisable to make a diagnosis of NASH before liver biopsy. The possibility of making an erroneous diagnosis is higher in NASH as compared to alcoholic liver disease. In our study, from 74 patients selected on clinical basis, 53 ones had NASH on liver biopsy. This figure is larger than the ones mentioned in other studies [30–33], which is probably due to precise patient selection in the present study, exclusion of cases with low aminotransaminase levels, or positive markers for viral hepatitis.

Factors affecting the pathogenesis of NASH are steatosis, inflammation, and fibrosis [2,34–38]. The severity of steatosis was similar to other reports. Also, the majority of patients had portal inflammation. The findings on liver biopsy were not correlated to liver enzymes, thus the increase in liver enzymes does not seem a good estimate of the severity of liver disease.

NASH can cause fibrosis and progress to cirrhosis. It has been shown that this progress is seen in 8–26% of patients [6,8,9,19]. Hepatic fibrosis, Mallory hyaline, or ballooning degeneration are findings that indicate higher likelihood of progression to cirrhosis [39]. Altogether, these findings further underscore the importance of NASH and the need to diagnose it before it causes irreversible liver damage.

## Conclusion

The epidemiology and risk factors of nonalcoholic steatohepatitis might be different in societies with lower rate of alcohol consumption, although this is just a preliminary hypothesis that should be evaluated in further studies. Regarding the considerable prevalence of the disease, confinement of the diagnosis to some specific epidemiologic and demographic subgroups does not seem so reasonable, particularly in regions where the distribution of the disease is not well-known.

## Competing Interests

None declared.

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