

# Epidemiologic Profile and Predictors of Fatty Liver (A Hospital-Based Study)

## Abstract:

**Background:** Non-alcoholic fatty liver is the most common cause of chronic liver disease with increasing prevalence globally. Settings and design: The current study is an analytical case control study; conducted in ultrasonography outpatient clinic of Cairo University Hospital. Material and **Methods:** 150 consented fatty liver cases and 564 controls were screened for fatty liver infiltration using abdominal ultrasonography. Receiver Operating Characteristics (ROC) curve analysis was performed to explore the discriminant ability of the developed model.

**Results:** Among cases: 32.7%, 36% and 31.3% had mild, moderate and severe degree of fatty liver respectively. Cases showed significantly higher body mass index(BMI), waist circumference (WC), total cholesterol, triglyceride, low density lipoprotein (LDL), and lower high density lipoprotein (HDL) than controls. Cases demonstrated higher prevalence of hypertension(11.3%vs 8.3% respectively), and significantly higher prevalence of diabetes(22% vs. 9.2%)(  $p=0.03$ ). Severe fatty liver cases were significantly older and had significantly higher WC , BMI, significantly higher association with diabetes mellitus, significantly higher levels of total cholesterol, triglycerides and LDL than non-severe degree cases. The significant predictors of sever fatty liver were BMI, total cholesterol and LDL ( $P = <0.001$ ,  $R^2 = 0.543$ ).

**Conclusion:** The developed regression equation expressed good validation and calibration. It utilizes an algorithm that can quickly and easily address patients with of fatty liver. It would useful as a fast, inexpensive primary screening tool for severe fatty liver.

**Key words:** Fatty liver; predictors; regression model; algorithm

## Introduction:

Non-alcoholic fatty liver disease (NAFLD) is the excessive accumulation of fat (steatosis) in  $\geq 5\%$  of hepatocytes in individuals who consume little or no alcohol. Steatosis eventually leads to cellular stress, injury and apoptosis [1]. It is a major cause of morbidity and mortality. It is the most common cause of chronic liver disease in many parts of the world and is a leading cause of liver transplant in the US. Its incidence and prevalence are rising globally parallel to the increasing rates of obesity and diabetes. It is associated with other components of the metabolic syndrome [2].

NAFLD affects about one third of the US general population. The prevalence in Europe, Middle East and Japan ranges from 20% to 30%. In China, the prevalence is 15–30%, and in India is 16% to 32% [1]; however, limited data is available on the prevalence of NAFLD in Africa. A Nigerian study estimated the prevalence to be 9% [3]. In Egypt, a hospital-based study in Alexandria concluded that Fatty liver was prevalent in schoolchildren (15.8%) [4], also NAFLD was found in 52.17% of polycystic ovary syndrome (PCOS) patients [5].

It is now a global public health problem that requires the attention of policy makers to set plans for its prevention and control in countries where the prevalence is increasing [6].

The spectrum of pathologic changes in the liver ranges from simple steatosis to nonalcoholic steatohepatitis (NASH), early fibrosis, cirrhosis and may progress to hepatocellular carcinoma (HCC). It is the third most common risk factor for HCC after viral infection and alcohol [7].

40 Liver biopsy is a gold standard technique for diagnosis of hepatic steatosis. However, it cannot be used for  
41 screening due to its invasiveness and sampling risks. It is indicated after a diagnosis is established by non-  
42 invasive techniques. [8].

43  
44 Controlled attenuation parameters (CAP) measured using transient elastography is a non invasive tool for the  
45 diagnosis and grading of hepatic steatosis. However, Its use as a screening tool is limited due to the high cost and limited  
46 availability [8].

47  
48  
49 Trans-abdominal ultrasound is commonly used imaging technique for fatty liver diagnosis. This is  
50 because it is an available, non-invasive and a low cost technique [9]. At ultrasonography, the diffuse fatty liver is  
51 characterized by hyper echogenicity of the liver parenchyma relative to the adjacent right kidney or spleen (so-  
52 called bright liver) [10]. Other features of fatty liver described by ultrasound are decreased visualization of  
53 vascular margins, attenuation of the ultrasound beam, loss of definition of the diaphragm, and hepatomegaly  
54 [11].

55 On the other hand, there are several limitations of ultrasonography that include its inability to  
56 distinguish between diffuse and focal hepatic steatosis. In addition, it is not a quantitative method, so it is not  
57 possible to distinguish between simple steatosis, advanced fibrosis, and early cirrhosis. It is limited by  
58 abdominal gas and body habitus, and it is non reproducible as it is operator dependent [12].

59  
60 Despite the enormous work and resources spent on the study of NAFLD, no effective treatment is  
61 currently available [2]. Therefore, it is essential to explore its epidemiological features and potentially preventable  
62 risk factors. Although screening is crucial especially in communities at risk, yet the high cost of testing, the risk  
63 of liver biopsy, and the low predictive value for non-invasive tests should be considered [13].

64 In the light of the information mentioned above, it is clear that a noninvasive, reliable, fast, and  
65 inexpensive tool for screening and staging of fatty liver is urgently needed. It would be useful particularly in  
66 clinics where ultrasound and or specialist is not available.

## 67 **Patients and methods:**

68 **Study design:** A hospital based case control analytical study.

69 **Study setting:** This study was conducted in ultrasonography outpatient clinic (under the supervision of Tropical  
70 Medicine Department) of Cairo University (Kasr-Alainy) Hospital.

71 **Study period:** from October 2013 till March 2016

72 **Participants:**

73 **A- Cases**

74 **Eligibility criteria:**

75 Inclusion criteria:

76 All attendants of the clinic who were confirmed to fulfill the criteria of bright liver through abdominal ultrasound  
77 were included.

78 Exclusion criteria:

79 -Patients with advanced comorbidities e.g., heart disease, renal failure.

80 - Patients with advanced hepatic disease (chronic hepatitis B or C) were excluded to avoid any confounding  
81 factors.

82 -Patients with significant alcohol consumption (more than once per day for women, and more than twice per day  
83 for men)

84 - Patients receiving any medications that may induce hepatic steatosis.

85

86 **B- Controls:** Healthy relatives (as proved by abdominal ultrasonography) of the study participants; who  
87 approved to participate in the study. They matched to cases as regards age and sex.

88 **Sampling:**

89 Sampling type was non-probability purposive sampling including all patients that fit the criteria of fatty liver as  
90 detected by ultrasonography. In order to minimize variation in scans interpretation, and to ensure consistency,  
91 the same expert Tropical Medicine consultant performed and graded all the scans. The average number of  
92 patients detected to have fatty liver by the same consultant was three patients per week; accordingly, this  
93 sample was collected over a period of two years as the ultrasound list was assigned every other week for the  
94 same Tropical Medicine consultant. A total of 150 patients were recruited to the study. Individuals who proved  
95 free from any fatty liver infiltration were recruited as the control group with a total of 564 cross matched controls.

96 **Study tools:**

97 1- An interview questionnaire was designed to collect data. The questions were close ended and were pre-  
98 coded prior to data collection to facilitate data entry and analysis. It included socio-demographic data, smoking  
99 history, and history of co-morbidities.

100 2- Anthropometric assessment included:

101 The weight that was measured in kilograms using traditional (non-digital weighing scale).

102 The height which was measured in meters using full length stadiometer.

103 BMI was calculated as follows : $BMI = \frac{Weight (Kg)}{Height (m)^2}$ . BMI was interpreted according to CDC 2015<sup>[14]</sup>.

104 3- Blood pressure was measured using mercury Sphygmomanometer. Blood pressure was interpreted  
105 according to Mayo clinic 2015<sup>[15]</sup>.

106 4- **Biochemical tests**: All patients were subjected to measurements of:

107 Liver enzymes: aspartate aminotransferase (AST), and alanine aminotransferase (ALT). The cutoff  
108 points for normal liver enzymes were interpreted according to Mayo clinic 2015<sup>[16]</sup>.

109 Lipid profile: triglycerides, total cholesterol, low density lipoprotein (LDL) and high density lipoprotein  
110 (HDL) were measured in (mg/dL). The cutoff points were interpreted according to Mayo clinic 2017<sup>[17]</sup>.

111 Virology markers: hepatitis B virus surface antigen (HBs Ag), HBV core antibody (HBc Ab Ig G) and  
112 hepatitis C virus antibodies (HCV Abs) (+VE/-VE).

113 Radiological investigations:

114 All participants were screened for fatty liver infiltration by abdominal ultrasonography (Famio5 TOSHIBA). The  
115 patients with fatty liver were classified into three groups according to the degree of their liver ultrasound  
116 echogenicity;

117 (1) The mild fatty liver was defined as a slight increase in liver echogenicity and the relative  
118 preservation of echoes from the walls of the portal vein.

119 (2) The moderate fatty liver was defined as moderate loss of echoes from the walls of the portal vein,  
120 particularly from the peripheral branches, and moderate diffuse abnormally bright echoes.

121 (3) The severe fatty liver was defined as a greater reduction in beam penetration, loss of echoes from  
122 most of the portal vein wall, and extensive, abnormally bright, echoes<sup>[18]</sup>. Beside hepatic echo pattern, liver  
123 size was also determined in addition to the other hepatic findings.

124 Data management and analysis:

125 All collected questionnaires were revised for completeness and consistency. Pre-coded data was entered on  
126 the computer using "Microsoft Office Excel Software" program for windows version 2010. Data was then

127 transferred to the Statistical Package of Social Science Software program, version 23 (SPSS) for statistical  
128 analysis.

129 Qualitative data was summarized using frequency and percentage, while quantitative data was checked for  
130 normality using Kolmogorov Smirnov test. Then normally distributed data was summarized using mean  $\pm$   
131 standard deviation and data that was not normally distributed was summarized using median and interquartile  
132 range (IQR).

133 Cases of fatty liver were classified into severe group vs. non severe group (mild and moderate fatty infiltration)  
134 for better comparison.

135 Comparison between groups was done using independent sample t-test for parametric quantitative data or  
136 Mann Whitney for non-parametric quantitative data, and Chi square test for qualitative variables. The logistic  
137 regression model was conducted to explore the significant predictors of fatty liver as well as sever form of fatty  
138 liver infiltration. Receiver Operating Characteristics (ROC) curve analysis was performed to explore the  
139 discriminant ability of the developed model. P values equal to or less than 0.05 were considered statistically  
140 significant. Graphs were used to illustrate some information.

#### 141 **Results:**

142 The basic characteristics of the studied group were demonstrated in table 1, they include socio-demographic  
143 profile of both cases and controls. Age, sex and residence matching were obvious with no significant difference  
144 between cases and controls (  $p= 0.3, 0.9$  and  $0.3$  respectively. An anthropometric assessment showed that  
145 cases exhibited significantly higher BMI and waist circumference ( $p<0.001$ ). Hypertension was more prevalent  
146 among cases than controls (11.3% vs. 8.3%); however no significant difference was detected. On the other  
147 hand, blood pressure measurement demonstrated a significant difference in both systolic and diastolic  
148 measurements being highest in cases but still within normal values. Diabetes mellitus was significantly more  
149 prevalent among cases than controls (22% vs. 9.2%) that was reflected on significantly elevated fasting blood  
150 sugar among them ( $p=0.03$ ). Despite of that, the mean FBS among cases was in the normal range. Nearly one  
151 third of the cases had hepatomegaly, three cases suffered from splenomegaly and nine cases had calcular gall  
152 bladder. More than one third of cases had a moderate form of fatty infiltration (36%) as shown in figure 1.  
153 Studying lipid profile of recruited population revealed a significantly higher level of total cholesterol, triglyceride,  
154 and LDL (  $p<0.001$ ) as well as a significantly lower level of HDL (  $p<0.001$ ). Although a significant difference  
155 was detected as regards ALT level being highest among cases, it is still within normal range with a median and  
156 interquartile ranges of 27 (21 - 35) vs. 24 (18 - 31.5).Normal level of AST was observed among both groups.

157 The backward stepwise logistic regression model was demonstrated in table 2. The last step revealed that only  
158 BMI, systolic blood pressure, total cholesterol, triglycerides, LDL and HDL were the actual significant predictors  
159 for severe fatty liver ( $X^2 = 534.5, df = 6, P = <0.001, R^2 = 0.527$ ). The model equation will be

160  $\text{Logit (P of fatty liver)} = -30.818 + 0.679 (\text{BMI}) + 0.044 (\text{SBP}) + 0.014 (\text{T. cholesterol}) + 0.023 (\text{Triglycerides}) +$   
161  $0.047 (\text{LDL}) - 0.110 (\text{HDL})$

162 ROC (receiver operating characteristics) curve analysis was performed to explore the discriminant  
163 ability of the predicted probability in differentiating fatty liver; it revealed that area under the curve (AUC) was  
164 0.979 with 95% CI (0.967 – 0.990). This means that the model equation expresses good discrimination.

165 The most suitable cut-off point in the predicted probability was 0.212 or more with sensitivity 92.7%  
166 (87.3-96.3), Specificity 94.0% (91.7-95.8), PPV 80.4% (74.6-85.0), and NPV 98.0% (96.5-98.8).

167 Comparing severe form of fatty liver versus other forms was presented in table 3. Among the socio-demographic  
168 characteristics only age was significantly different. Severe infiltration was more obvious with older age. The  
169 anthropometric assessment showed that severe cases exhibited significantly higher BMI and waist  
170 circumference than other forms ( $p < 0.001$  &  $0.007$  respectively). Hypertension was more prevalent among  
171 severe cases than other forms (14.9% vs. 9.7%); however, this difference was not statistically significant.  
172 Also, blood pressure measurement demonstrated no significant difference in both systolic and diastolic  
173 measurements which were within normal values. Diabetes mellitus was significantly more prevalent among  
174 severe cases than other forms (31.9% vs. 17.5%). Nearly half of the severe cases had hepatomegaly, 2 cases  
175 suffered from splenomegaly and all patients who had calculi gall bladder were belonging to non severe degree  
176 groups. Lipid profile analysis of patients revealed a significantly higher level of total cholesterol, triglyceride, and  
177 LDL ( $p < 0.001$ ), with lower, but not significant HDL level ( $p < 0.08$ ). Also, no significant difference was  
178 detected as regards ALT level and AST level in both groups ( $p = 0.9$  and  $0.7$  respectively).

179 The backward stepwise logistic regression model was presented in table 4. The significant predictors  
180 of severe fatty liver were only BMI, Total cholesterol and LDL ( $X^2 = 117.5, df = 3, P = <0.001, R^2 = 0.543$ ). The  
181 model equation will be

182  $\text{Logit (P of severe)} = -25.717 + 0.440 (\text{BMI}) + 0.031 (\text{T. cholesterol}) + 0.023 (\text{LDL})$

183 ROC (receiver operating characteristics) curve analysis was performed to explore the discriminant  
184 ability of the predicted probability in differentiating severe fatty liver, it revealed that area under the curve (AUC)  
185 was 0.966 with 95% CI (0.941 – 0.990). This means that the model equation expresses good discrimination.

186 The most suitable cut-off point in the predicted probability was 0.236 or more with sensitivity 95.7%  
187 (85.5-99.5), Specificity 88.3% (80.5-93.8), PPV 78.9% (66.1-88.6), and NPV 97.8% (92.4-99.7).

188

## 189 **Discussion:**

In the current study, age was a significant risk factor for higher grades of fatty liver. This may be due to 190  
long duration of exposure to unhealthy dietary and life style factors. Similarly findings were reported by other  
studies [18,19, 20,21]. Contrary to that, other studies concluded that age was a non-significant predictor for fatty liver  
[22,23]. The discrepancy of age association with high prevalence of NAFLD as well as its complications may be  
attributed to the duration of disease rather than age. 194

195 Metabolic syndrome components are strongly associated NAFLD is [2]. This was noticed in the current  
196 study. BMI and WC showed significantly higher mean values among cases than controls. It is noteworthy to  
197 mention that the mean BMI among all cases of NAFLD in the current study was in the obesity category ( $33.3 \pm$   
198  $4.3 \text{ Kg/m}^2$ ). Similar findings were reported by other studies such as the Egyptian study conducted by Hegazy  
199 and Mostafa, where the BMI in NAFLD and NASH patients were in the obese category [24]. Similarly, Fu and  
200 colleagues concluded that overweight and obese persons had a high probability to develop fatty liver than  
201 subjects with normal BMI [25]. On the contrary, a Japanese study found lower BMI among fatty liver patients [22].  
202 This discrepancy may be due to demographic and dietary differences between Egyptian and Japanese  
203 population.

204 Additionally, this study's participants with severe fatty liver showed significantly higher WC, weight and  
205 BMI than those with non severe forms of the disease. This coincides with a study performed by Lin and  
206 colleagues, where BMI was found to be a significant independent predictor for different grades of fatty liver [18].

207 Furthermore, it was noticed that cases of NAFLD in the current study demonstrated statistically significant  
208 higher levels of lipid profile parameters compared to their matching controls. However, triglycerides and LDL  
209 levels among cases were the only two parameters in lipid profile that exceeded the cut off limits of Mayo clinic  
210 recommendations [17].

211 Comparing the lipid profile parameters among the different grades of fatty liver, it was noticed that  
212 participants with severe fatty liver demonstrated elevated and statistically significant higher levels of total  
213 cholesterol, triglycerides and LDL compared to participants with non-severe fatty liver. However, LDL levels

214 were above the recommendations in both groups. Also, HDL level was below the recommendations and lower  
215 among cases with severe fatty liver than those with non-severe forms. Similar findings were reported in another  
216 study; where elevated total cholesterol level, triglycerides and LDL, and decreased HDL were significantly  
217 associated with higher degree of fatty liver, but only total cholesterol and triglycerides were the independent  
218 predictors <sup>[18]</sup>. Also, high total cholesterol and triglycerides were associated with the development of NAFLD <sup>[22]</sup>.

219 Liver biopsy is the gold standard for the diagnosis of NAFLD, but it is an invasive technique that can  
220 cause complications <sup>[26]</sup>. Ultrasound is an available accurate technique <sup>[27]</sup>. In addition; laboratory and clinical  
221 parameters are not always consistent. About 70% of patients with NASH and significant fibrosis show normal  
222 liver enzymes <sup>[28]</sup>. Furthermore, NAFLD is not necessarily accompanied by obesity and metabolic syndrome <sup>[29]</sup>.

223 A reliable, non invasive tool for screening and staging of NAFLD is thus urgently needed. The current  
224 study utilizes an algorithm that can quickly and easily address patients with severe degrees of fatty liver. It  
225 would be useful as a primary screening tool for severe fatty liver that is fast and inexpensive; especially in  
226 clinics where ultrasound or a specialist are not available

227 Since bright liver is considered a silent precursor for a wide variety of non-communicable diseases like  
228 metabolic syndrome, liver cirrhosis and cancer liver, it is better to pick up those at risk as early as possible with  
229 a simple, fast and reliable tool to be adjusted for prompt treatment before permanent disorders occur. The  
230 current study provides an easy, simple and quick algorithm to predict higher degrees of fatty liver without the  
231 need for any trained personnel or advanced techniques. It is of a high predictive power with a coefficient of  
232 determination ( $R^2 = 0.543$ ) despite using only three variables (BMI, total cholesterol and LDL) i.e. about fifty four  
233 percent of variability of occurrence of bright liver was explained by these three variables. In addition to that, the  
234 algorithm also reported high validity parameters in predicting bright liver (sensitivity 78.7%, specificity 94.2%,  
235 PPV 86.0%, NPV 90.7% and accuracy 89.3%). The area under the ROC curve was also so high (0.966 95% CI  
236 0.941-0.990) with most suitable cut-off point  $\geq 0.236$  with sensitivity 95.7%, specificity 88.3%, PPV 78.9%, NPV  
237 97.8% and accuracy 90.7%

238 Four other algorithms using biochemical and demographic parameters to assess liver steatosis are the  
239 SteatoTest <sup>[30]</sup>, the Fatty Liver Index <sup>[31]</sup>, Lin, et al Index <sup>[18]</sup> and Bedogni, et al Index <sup>[32]</sup>.

240 In contrast to the Steato Test and Fatty Liver Index, the algorithm used in the current study was  
241 developed with data from fatty liver of apparently healthy participants and was intended for prediction of severe



242 stages of hepatic steatosis. Although its defect in predicting mild and moderate steatosis, it had a reasonable  
243 predictive power for the presence of severe steatosis. Lin et al Index was developed for predicting moderate to  
244 severe degrees of fatty liver, but had a sensitivity of 70.8%, a specificity of 85.2%, a PPV of 63.2%, and a NPV  
245 of 88.8% [18].

246 **Conclusion:** The current study provides an easy, simple and quick algorithm to predict higher degrees of fatty  
247 liver. It is of a high predictive power with a coefficient of determination ( $R^2 = 0.543$ ) despite using only three  
248 variables (BMI, total cholesterol and LDL) i.e. about fifty four percent of variability of occurrence of bright liver  
249 was explained by these three variables. In addition to that, the algorithm also reported high validity parameters  
250 in predicting bright liver (sensitivity 78.7%, specificity 94.2%, PPV 86.0%, NPV 90.7% and accuracy 89.3%).  
251 The area under the ROC curve was also so high (0.966 95% CI 0.941-0.990) with most suitable cut-off point  $\geq$   
252 0.236 with sensitivity 95.7%, specificity 88.3%, PPV 78.9%, NPV 97.8% and accuracy 90.7%.The method  
253 described in the current study utilizes an algorithm that can quickly and easily address patients with higher  
254 degrees of fatty liver.

255 **Recommendations:** Enhancing "Health Literacy" of the public is recommended as well as periodic screening  
256 of at risk groups for early detection of modifiable risk factors of fatty liver disease. Additionally, people with  
257 diabetes are advised to properly control their metabolic parameters .Further research is recommended in order  
258 to validate the algorithm developed in the current study on a large scale before dissemination to the outpatient  
259 clinics as an easy, non-invasive, applicable and accessible screening tool, especially when abdominal  
260 ultrasonography and or experts are not available.

#### 261 **Ethical consideration:**

262 **Administrative issues:** This study was approved from both Public Health and Tropical Medicine Departments  
263 through Department Council meeting on July and August 2013 respectively. Approval from the ethical  
264 committee of Public Health Department was obtained as well.

265 **Informed consent:** The study was conducted after explaining the study objectives to the patients. Only those  
266 who agreed were included in the study. Verbal consents were obtained from all the study participants before  
267 starting to collect data. Confidentiality of obtained information was ensured. All subjects were treated  
268 according to the Helsinki Declaration of biomedical ethics [33].

#### 269 **Limitations of the study:**

270 Inability to perform liver biopsy due to ethical consideration as this invasive maneuver needs strict indications  
271 and certain precautions.

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351 **Subjects, as adopted by the 52nd WMA General Assembly, Edinburgh, October 2013.**

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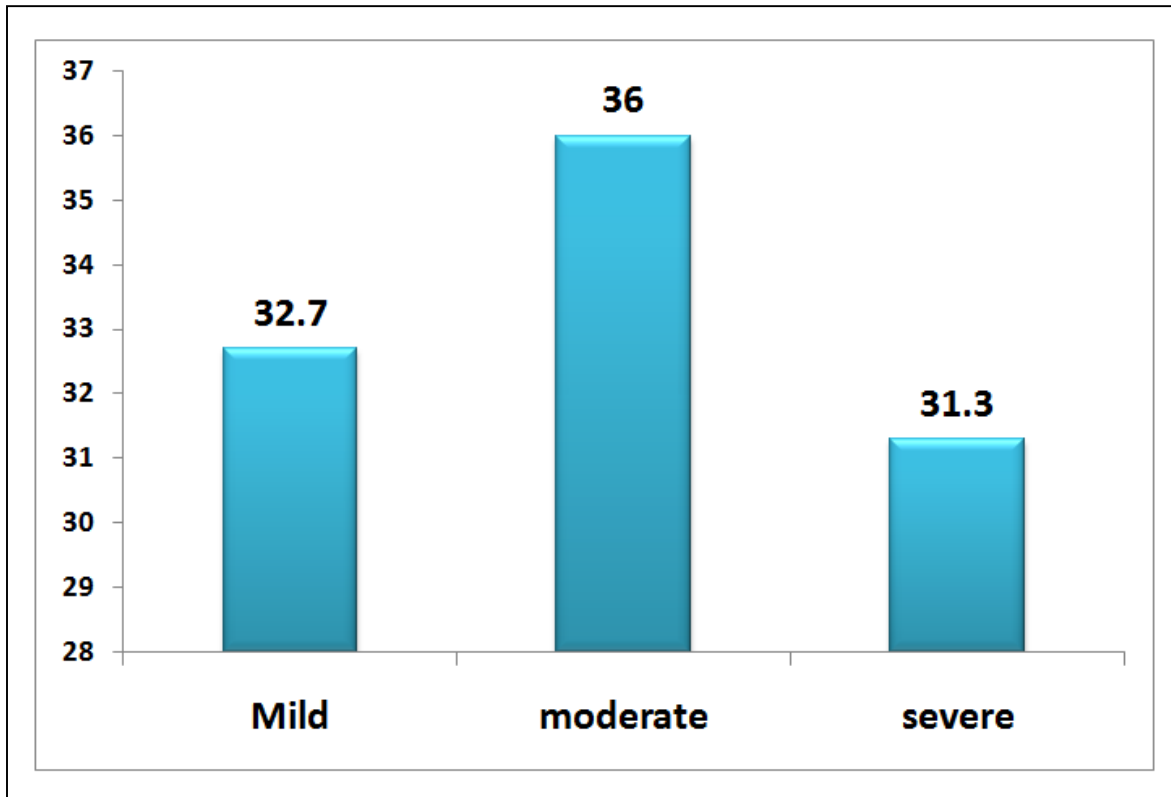
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354 Tables and graphs:

355 Table (1): Basic characteristics of the studied population

Characteristics	Cases (150)	Control(564)	P value
<b>Socio-demographic</b>			
Age (years)	46.8 ± 9.1	46 ± 8.9	0.306*
Sex			
Male	58 (38.7)	219 (38.8)	0.971#
Female	92 (61.3)	345 (61.2)	
Residence			
Urban	131 (87.3)	472 (83.7)	0.273#
Rural	19 (12.7)	92 (16.3)	
<b>Anthropometric measurement</b>			
BMI (Kg/m <sup>2</sup> )	33.3 ± 4.3	26.4 ± 2.8	<0.001*
Waist circumference (cm)	109.6 ± 9.1	92.7 ± 12.2	<0.001*
<b>Co-morbidities</b>			
Smokers	27 (18)	139 (24.6)	0.087#
Hypertension	17 (11.3)	47 (8.3)	0.253#
Systolic blood pressure (mmHg)	129.1 ± 18.1	118.4 ± 12.1	<0.001*
Diastolic blood pressure(mmHg)	81.3 ± 12.3	79 ± 8.1	0.026*
Diabetes mellitus	33 (22)	52 (9.2)	<0.001#
<b>Sonographic findings</b>			
Hepatomegaly	47 (31.3)	0 (0)	<0.001#
<b>Laboratory investigation</b>			
Total cholesterol (mg%)	199 ± 69.8	143.1 ± 30.1	<0.001*
Triglycerides (mg%)	154.7 ± 55.8	102.7 ± 26.3	<0.001*
LDL (mg%)	139 ± 40	89.3 ± 19.6	<0.001*
HDL (mg%)	45.2 ± 9.8	56.7 ± 12.8	<0.001*
Fasting blood sugar (mg%)	103.4 ± 34	96.8 ± 22.7	0.027*
ALT (IU/L)	27 (21 - 35)	24 (18 - 31.5)	0.011@
AST (IU/L)	29 (22 - 37)	28 (22 - 36)	0.127@

356 \*independent sample t-test, @Mann Whitney test, #Chi square test, qualitative variables described as n (%),  
 357 quantitative variables described as mean ± standard deviation or median(interquartile range), LDL= Low-density  
 358 lipoprotein, HDL= High-density lipoproteins, BMI= body mass index,ALT=Alanine Aminotransferase, and AST=  
 359 Aspartate Aminotransferase



360

361 **Figure (1): Percent distribution of fatty liver degrees among cases**

362

363

**Table (2): significant predictors of fatty liver**

	OR	95% CI for OR	P value
<b>BMI (Kg/m<sup>2</sup>)</b>	1.972	1.603-2.425	<0.001
<b>Systolic blood pressure (mmHg)</b>	1.045	1.021-1.070	<0.001
<b>Total cholesterol (mg%)</b>	1.014	1.002-1.026	0.018
<b>Triglycerides (mg%)</b>	1.023	1.008-1.038	0.003
<b>LDL (mg%)</b>	1.048	1.029-1.068	<0.001
<b>HDL (mg%)</b>	0.896	0.856-0.938	<0.001

364 LDL= Low-density lipoprotein, HDL= High-density lipoproteins, BMI= body mass index, OR= odds ratio, CI=

365 confidence interval

366 Table (3): Comparison between severe degree of fatty liver versus other degrees

	Severe Fatty liver (n=47)	Non-Severe Fatty liver (n=101)	P value
<b>Socio-demographic characteristics</b>			
Age (years)	49.5 ± 8.4	45.6 ± 9.2	<b>0.016*</b>
Sex			
Male	18 (38.3)	40 (38.8)	0.950#
Female	29 (61.7)	63 (61.2)	
Residence			
Urban	41 (87.2)	90 (87.4)	0.980#
Rural	6 (12.8)	13 (12.6)	
<b>Anthropometric measurement</b>			
BMI (Kg/m <sup>2</sup> )	37.2 ± 3	31.5 ± 3.6	<b>&lt;0.001*</b>
Waist circumference (cm)	114.6 ± 7.9	107.7 ± 8.9	<b>0.007*</b>
<b>Co-morbidities</b>			
Smokers	6 (12.8)	21 (20.4)	0.260#
Hypertension	7 (14.9)	10 (9.7)	0.353#
Systolic blood pressure (mmHg)	129.7 ± 18.7	128.8 ± 18	0.780*
Diastolic blood pressure (mmHg)	81.2 ± 11.9	81.4 ± 12.5	0.935*
Diabetes mellitus	15 (31.9)	18 (17.5)	<b>0.048#</b>
<b>Sonographic findings</b>			
Hepatomegaly	23 (48.9)	24 (23.3)	<b>0.002#</b>
Calcular Gall bladder	0 (0)	9 (8.7)	0.057#
Splenomegaly	2 (4.3)	1 (1)	0.231#
<b>Laboratory investigation</b>			
Total cholesterol (mg%)	265.4 ± 76.4	168.7 ± 39.1	<b>&lt;0.001*</b>
Triglycerides (mg%)	212.9 ± 66.1	128.2 ± 17.3	<b>&lt;0.001*</b>
LDL (mg%)	167.1 ± 34.5	126.1 ± 35.7	<b>&lt;0.001*</b>
HDL (mg%)	43.2 ± 9.3	46.2 ± 9.9	0.081*
Fasting blood sugar (mg%)	107.6 ± 36.2	101.5 ± 33	0.311*

<b>ALT (IU/L)</b>	27 (18 - 35)	27 (21 - 34)	0.913@
<b>AST (IU/L)</b>	28 (21 - 38)	29 (23 - 37)	0.703@

367 \*independent sample t-test, @Mann Whitney test, #Chi square test, qualitative variables described as n (%),  
368 quantitative variables described as mean ± standard deviation or median(interquartile range), LDL= Low-density  
369 lipoprotein, HDL= High-density lipoproteins, BMI= body mass index, ALT=Alanine Aminotransferase, and AST=  
370 Aspartate Aminotransferase

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371 **Table (4): significant predictors of severe form of fatty liver**

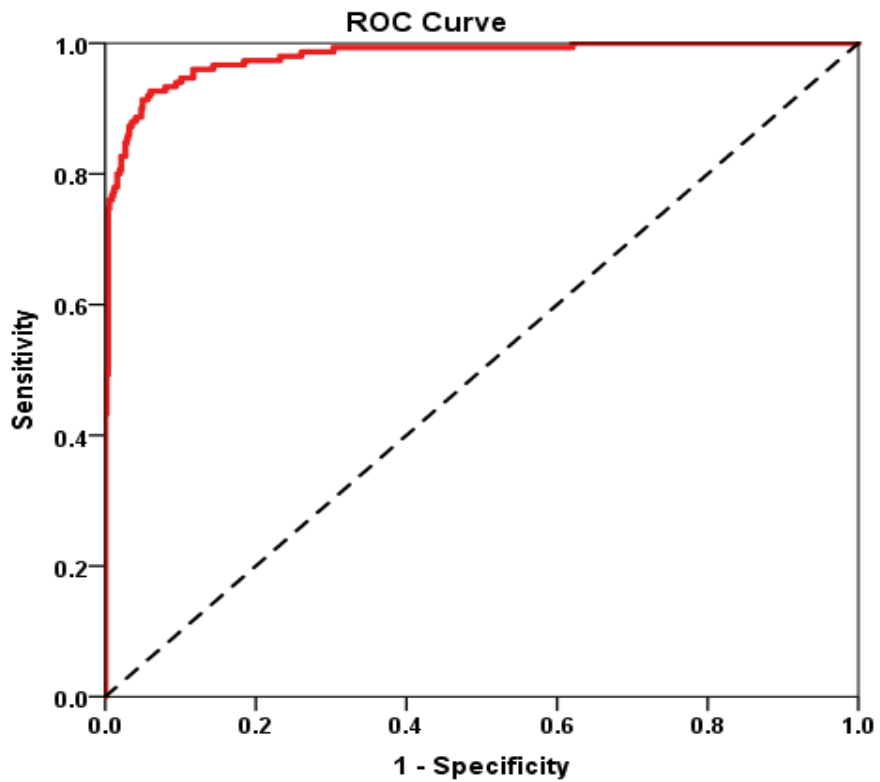
	OR	95% CI for OR	P value
<b>BMI (Kg/m2)</b>	1.553	1.269-1.899	<0.001
<b>Total cholesterol (mg%)</b>	1.032	1.016-1.048	<0.001
<b>LDL (mg%)</b>	1.023	1.007-1.040	0.006

372 LDL= Low-density lipoprotein, OR= odds ratio, CI= confidence interval

373 The model equation will be

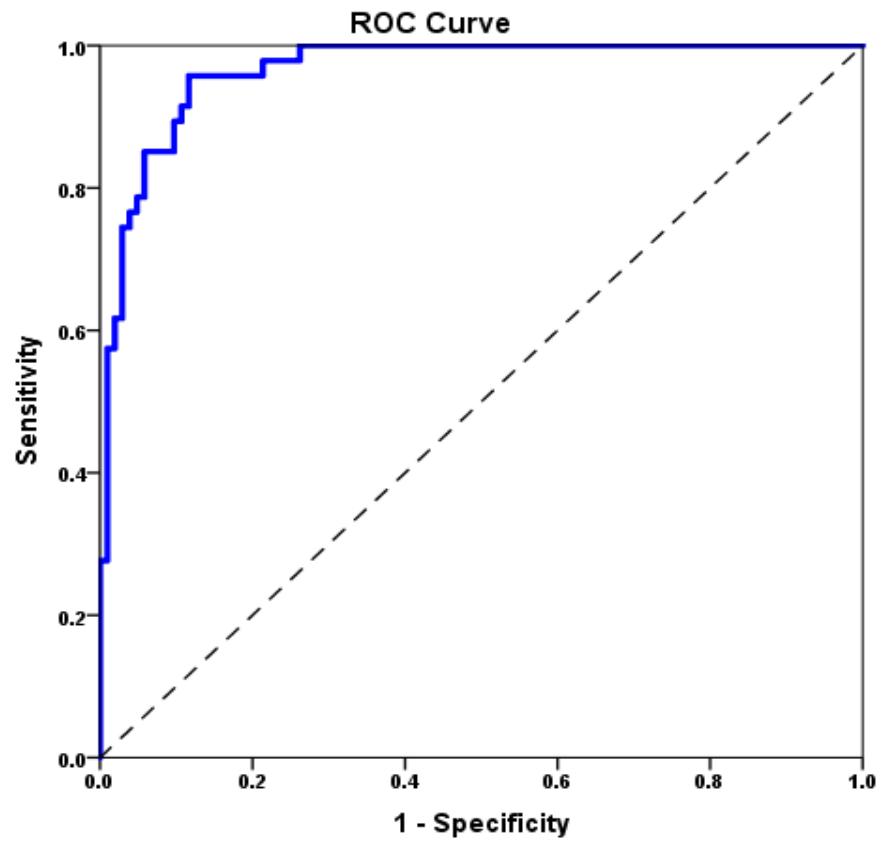
374  $\text{Logit (P of severe)} = -25.717 + 0.440 (\text{BMI}) + 0.031 (\text{T. cholesterol}) + 0.023 (\text{LDL})$

375



376

377 **Figure (2a): ROC curve for the predicted probability to discriminate fatty liver**



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379 **Figure (2b): ROC curve for the predicted probability to discriminate severe fatty liver**

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