Study the Effect of Weight Loss in Improvement of Hepatic Fibrosis in Patients with Chronic HCV

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Abstract

Background: Liver fibrosis considered as a reversible disease, but curing the primary hepatic disease, sometimes, doesn’t mean fibrosis improvement. Steatosis could be one of the reasons responsible for this failure. Weight loss has been shown to improve fibrosis, but few data is available from Egypt.

Objective: to test the effect of weight reduction on the degree of fibrosis improvement in patients with chronic hepatitis C (CHC). Methods: Adult patients (> 18 years) were recruited from hepatitis C patient’s clinic, National Liver Institute (NLI), University of Menoufia, from 2016 to 2018.

Results: Thirty CHC patients with moderate and severe steatosis, who were initially assessed by liver biopsy and Transient elastography (TE) for their fibrosis stage, were followed up by Transient elastography after losing >10% of their weight. Median age was 44.7 years (29 – 59 years) and 50% were men. The degree of fibrosis and its values improved with the significant weight loss (p < 0.01 for both).

Conclusion: Weight reduction is associated with improvement of hepatic fibrosis in patient with CHC.

Keywords: Fibrosis; Weight loss; Transient elastography; HCV; Steatosis

Abbreviations: CHC: Chronic Hepatitis C; NLI: National Liver Institute; TE: Transient Elastography; NAFLD: Nonalcoholic Fatty Liver Disease; N: Number, SD: Standard Deviation, BMI: Body Mass Index; ALT: Alanine Aminotransferase, AST: Aspartate Aminotransferase, RNA: Ribonucleic Acid, INR: International Normalized Ratio, kPa: Kilopascal

Introduction

Obesity and its direct consequence, nonalcoholic fatty liver disease (NAFLD), rates are increasing dramatically worldwide, and steatosis becomes one of the most common hepatic histological findings [1]. It was found to be associated with liver fibrosis especially in chronic hepatitis C (CHC) with positive correlation between its grade and the fibrosis stage [2,3]. Liver fibrosis is a dynamic progressive process as long as the hepatic injurious stimulus continues unopposed. It was considered irreversible in the past, but the current studies showed that fibrosis [4-6], and even cirrhosis [7,8] in some series, could be reversible diseases. Our study aimed to compare the degree of hepatic fibrosis in overweight patients with CHC before and after weight loss achieved by diet and exercise, as it is considered the standard treatment for steatosis [9,10].

Methods

Adult patients (> 18 years) were recruited from hepatitis C clinic, National Liver Institute (NLI), University of Menoufia, from January 2016 to May 2018. Standard available treatment at that time was pegylated interferon with ribavirin. The assessment process necessitates doing liver biopsy and transient elastography (TE). The NLI protocol asked the patient with moderate or severe steatosis to lose at least 10% of his weight, based on food diet and practicing exercises, to improve the response rate to treatment. FibroScan was done again to reassess the patient’s hepatic fibrosis stage after losing weight without the need for re-biopsy. Our retrospective study was conducted on those patients who succeeded to lose the desired weight. We reviewed patients’ records and we had thirty patients to be involved in the study after excluding files with incomplete data and patients with BMI > 40 kg/m², as that could affect the FibroScan result.

Our study was approved from the ethical committee, NLI, University of Menoufia. Height and weight were recorded while BMI was calculated as defined by WHO [11]: bodyweight in kilograms divided by height in meters squared. Liver function evaluation was done by the available data of clinical examinations and laboratory tests.

All liver biopsy specimens were analyzed by one experienced pathologist blinded to the clinical data according to METAVIR scoring system for assessment of fibrosis stage [12]. Steatosis was categorized by visual assessment as: if steatosis presents...
in 1-10% of hepatocytes [mild]; in 11-30% [moderate]; and 31-100% [severe].

**Transient elstography**

It was performed using the FibroScan® apparatus (Echosens, Paris, France), with 5-MHz transducer. The fibrosis grading score used was: F0= 0-2.9 kPa; F1= 3-5.9 kPa; F2= 6-8.9 kPa; F3= 9-16.9 kPa; and F4= 17-75 kPa (results were expressed in kilopascals) [13-15].

**Statistical analysis**

Data was statistically analyzed using SPSS version 21 for windows and for all the analysis a p value< 0.05 was considered statistically significant. Data are shown as mean, range or value. As appropriate, stages of fibrosis were tested against continuously-distributed variables using the Anova test. Chi-squared test was used to test two ordinal variables, while paired t test was used for pre and post values of the same variable of the same group of patients.

**Results**

Table 1: Descriptive data and patients’ characteristics at the beginning of assessment.

| Studied Variables | (N=30) Mean ± SD | Range |
|-------------------|------------------|-------|
| Age               | 44.7 ± 8.9       | 29 – 59 years |
| Male              | 15               |       |
| Female            | 15               |       |
| BMI               | 33.2 ± 2.5       | 27 – 36.9 kg/m² |
| ALT               | 49.8 ± 15.8      | 22 – 88 U/L |
| AST               | 49.9 ± 16.1      | 19 – 79 U/L |
| Albumin           | 4.2 ± 0.4        | 3.6 – 4.9 gm/dl |
| Total bilirubin   | 0.7 ± 0.2        | 0.3 – 1.1 mg/dl |
| Direct bilirubin  | 0.2 ± 0.1        | 0.03 – 0.5mg/dl |
| INR               | 1.1 ± 0.1        | 1 – 1.4 |
| HCV RNA level     | 779.6 ± 1385.5   | 17.5-5639.9 IU/ml× 10³ |
| TE                | 13.8 ± 7.5       | 6.1 – 30.8kPa |

N: Number; SD: Standard Deviation; BMI 1 or 2: Body Mass Index at the beginning of assessment or after weight reduction; ALT: Alanine Aminotransferase; AST: Aspartate Aminotransferase; RNA: Ribonucleic Acid; INR: International Normalized Ratio; kPa: kilopascal

The present study included 30 patients with CHC with moderate and severe steatosis who were initially assessed by liver biopsy and TE for their fibrosis stage, activity grade and degree of steatosis. After losing 10% or more of their weight, they were followed up by TE only. Our patients’ median age was 44.7 years (29 – 59 years) and 50% were men, Table 1 shows the characteristics of patients. According to the histological examination there were 20 patients with moderate and 10 with severe steatosis. Table 2 shows the frequency of different grades of fibrosis between patients according to Metavir classification. Our results showed great improvement of liver stiffness measured by TE after body weight loss. Table 3 shows the frequency of different grades of fibrosis between patients before and after weight reduction and the significant reversibility of fibrosis after weight reduction (p < 0.01).

Table 2: Characteristics of patients’ liver biopsies.

| Fibrosis stage | Frequency | % |
|----------------|-----------|---|
| F1             | 11        | 36.7 |
| F2             | 9         | 30  |
| F3             | 8         | 26.6 |
| F4             | 2         | 6.7  |

| Activity grade | Frequency | % |
|----------------|-----------|---|
| A1             | 16        | 53.3 |
| A2             | 13        | 43.3 |
| A3             | 1         | 3.4  |

| Steatosis degree | Frequency | % |
|------------------|-----------|---|
| No               | 0         | 0  |
| Mild             | 0         | 0  |
| Moderate         | 20        | 66.7 |
| Severe           | 10        | 33.3 |

| Total            | 30        | 100 |

Liver stiffness 1: at the beginning of assessment; Liver stiffness 2: after weight reduction

Table 3: Classification according to liver stiffness measured by TE.

| Stage of Fibrosis | Liver Stiffness 1 Number (%) | Liver Stiffness 2 Number (%) |
|-------------------|------------------------------|------------------------------|
| F1                | 0 (0%)                       | 5 (16.70%)                   |
| F2                | 9 (36.70%)                   | 9 (30%)                      |
| F3                | 10 (33.30%)                  | 10 (33.30%)                  |
| F4                | 9 (30%)                      | 6 (20%)                      |

Paired t-test

SD: Standard deviation; kPa: kilopascal, BMI 1: Body Mass Index at baseline; BMI 2: Body Mass Index after weight reduction

Table 4: Follow up of BMI and Fibro Scan after the decrease in BMI.

| Mean          | SD     | Range  | p-value  |
|---------------|--------|--------|----------|
| BMI 1 kg/m²   | 33.23  | 2.5    | 27 – 36.9| <0.01*   |
| BMI 2 kg/m²   | 29.05  | 1.92   | 24.2 – 32.5|         |
| Fibro Scan 1 kPa| 13.81  | 7.46   | 6.1 – 30.8| <0.01*   |
| Fibro Scan 2 kPa| 10.98  | 5.57   | 3.8 – 23.9|         |

Table 5: Correlation between TE stages of fibrosis before and after weight loss

| Fibrosis stage 1 | Fibrosis stage 2 |
|------------------|------------------|
| 0.7              | <0.01            |

For statistical reason we tested the FibroScan results of liver stiffness again as values in kPs, to avoid any statistical defect when grouping patients into grades. Table 4 shows descriptive data (mean, standard deviation and range) of BMI and liver stiffness measurements. It shows the highly significant improvement of liver fibrosis after weight loss (p < 0.01) (Table 5).
Discussion

Liver fibrosis reversibility is still a hot issue. Although liver fibrosis is agreed to be a dynamic process depending on the balance between degradation and remodeling of extracellular matrix deposition by metalloproteinase enzymes and their tissue inhibitors, yet the mechanism is not completely understood. For instance, treating the cause of liver disease was thought to improve the ability of liver tissue to reverse fibrosis, but studies showed incomplete or insignificant improvement in some patients [6,7]. This could mean that, there are other co-factors which if treated could improve the healing response.

Increased body weight is well known to increase fat deposition in hepatocytes i.e. steatosis and it was claimed to cause liver fibrosis. Studies showed that decreasing its grade by weight loss achieved through lifestyle modifications (diet and exercise) could improve hepatic fibrosis [16], but few or scarce data are available from Egypt, in spite of the high prevalence of liver disease in our population [17].

In our study we depended on Transient Elastography in the assessment of liver fibrosis as it was proved to be a reliable method to detect significant fibrosis compared to histological assessment of liver tissue [14,18], it measures liver stiffness based on the propagation speed of shear waves with overall diagnostic accuracy reaching 94% [19].

Our study goal was to test if losing significant weight could improve the degree of fibrosis in Egyptian patients suffering CHC. We included patients with moderate or severe steatosis and compared their TE results before and after losing weight. We used BMI11in the assessment of our patients' weight reduction, and we found a significant decrease in it with mean of 4.18 kg/m²; p < 0.01.

Our main result was that there is marked improvement in the degree of fibrosis in the second scan and before starting the treatment of CHC. This could mean that losing weight is an important factor for healing liver fibrosis and it could have an additive effect with treating the main cause of liver injury, hepatitis C in our case. Our results go with Hickman et al16 who investigated 19 patients using liver biopsy before and after weight reduction and found that it was associated with improvement of fibrosis.

Also, Dixon et al, [20] who investigated the effect of losing significant weight after laparoscopic adjustable gastric band placement on the degree of liver fibrosis and inflammation in patients suffering nonalcoholic fatty liver by doing two liver biopsies, before and after weight loss. They reported that significant weight loss was associated with marked improvement in hepatic necroinflammatory stage and stage of fibrosis.

Conclusion

Weight loss is important to achieve better improvement in hepatic fibrosis beside the treatment of the primary cause of the liver disease. More work is needed to find out other co-factors which, if treated could improve the response to the standard available treatments.

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