Estimation of HCV viral load and liver enzymes among different patients groups of District Gujrat, Pakistan

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ABSTRACT
Hepatitis C has a 3% of the global disease burden that remains endemic in many regions of the world. According to a general statistical survey it has approximately 5.3% seroprevalence in Pakistan. HCV is a persistent and silent disease thus making the primary diagnosis complicated. Occasionally HCV positive population could not be diagnosed by routine HCV antibody testing therefore requires molecular diagnosis. This study is aimed to determine the prevalence of HCV and to estimate the HCV viral load by quantitative analysis among different patient groups of District Gujrat, Pakistan. A total of 597 samples were collected from clinically diagnosed liver patients that were categorized into three age groups: 1) up to 25 years; 2) 26 - 50 years 3) above 50 years. All samples were subjected to real time PCR for determination and quantification of HCV RNA. Activity of liver aminotransferases was measured. The overall prevalence of HCV-RNA was 73.87%. Females had slightly higher HCV prevalence which is 74.06% while 73.45% in males. Highest prevalence of active HCV infection was found in age group 26 - 50. In addition, liver function tests showed that 28.12% HCV-positive patients do not have elevated ALT level whereas 32.65% did not show elevated AST levels. It may be assumed that there is not a significant relationship between increased viral load and liver amino transferases. The study concluded a significantly higher rate of HCV infection in young population. Moreover screening with antibody and liver function tests alone does not exclude the possibility of HCV infection.

Keywords: Hepatitis C; Viral Load; Alanine Aminotransferase; Aspartate Aminotransferase; HCV RNA; Gujrat

1. INTRODUCTION
Hepatitis C virus (HCV) is a widespread infection that causes significant morbidity and mortality all over the world [1]. HCV infection is often asymptomatic and almost in 70% cases leads to chronic hepatitis. This may results in cirrhosis, liver failure, hepatocellular carcinoma or progressive liver disease within 20 to 30 years. Viral load is one of the factors associated with progression of disease in HCV infection [2]. Hepatitis C was first discovered in 1989, at that time it was thought of minor importance because of its difference from both hepatitis A and B. At present it has become one of the major global health issue that needs to be actively controlled and prevented [3].

Around the globe, approximately 130 - 170 million persons are living with chronic hepatitis C virus infection [4]. Egypt has the largest burden of HCV in the world, with 10% prevalence of chronic HCV infection among the persons aged 15 - 59 years [5]. It is also a common chronic blood-borne infection in USA, causing an estimated 8000 - 10,000 death annually which will be increased substantially in the next 20 years [6]. There are almost 10 million people with HCV infection in Pakistan [7].

Chronic HCV is usually treated with interferon (IFN) alpha (a glycoprotein) that is considered as a backbone of HCV treatment because of its efficient role to enhance the immune response against virus [8]. At present, peglated interferon (PEG-IFN) in combination with ribavirin is regularly used for HCV treatment. This therapy achieves
about 80% SVR for genotype 2 & 3 and 50% for genotype 1 [9]. Pegelated interferons are of two types, PEG-IFN-alpha-2a and PEG-IFN-alpha-2b. They are different in their size and configuration of the polyethylene glycol that contains binding sites for interferon. The two formulated interferons are equally good for HCV treatment but their functioning is not compared [10]. Pre viral loads are useful determinants of response to antiviral therapy [11], hence initial viral load and decrease of viral load at initial stage of antiviral therapy are useful parameters to check efficacy and to optimize antiviral therapy [12].

There are different parameters for HCV diagnosis. In most of the cases HCV prognosis is based on liver function tests. Abnormal level of serum amino transaminases may lead to the prognosis of HCV infection. ALT is most intense in liver and released into the bloodstream as the result of liver damage that fairly indicates liver status [13]. In fact, before the detection of HCV and the exploitation of molecular based essays for HCV RNA, the non-A, non-B hepatitis was based on records of steadily abnormal ALT levels [14]. On the other hand, almost 25% of patients with active HCV infection have persistently normal ALT values [15]. It is considered that occult-HCV (characterized by presence of liver disease and absence or very low levels of detectable HCV-RNA in serum) can be indicated by abnormal liver function tests although virus may persist for years after spontaneous recovery or after sustained viral response [16]. In this scenario, Viral Load is a good parameter for the presence of hepatitis C virus and it also measures the amount of HCV in the blood. If viral load (RNA) is found significantly in an individual’s blood, it confirms active infection [17]. If viral load is less than 800,000 IU/mL, it is considered as low viral load while more than 800,000 IU/mL, is called high viral load [18].

In the present study patient data were collected from densely populated District Gujrat of Pakistan and tested for the presence of HCV infection with molecular analysis. Quantification of viral load was done to estimate the disease progression and compared with liver enzymes activity. The data showed high HCV prevalence in young population particularly in females. Also a discrepancy was found between viral load and liver function tests.

2. MATERIAL AND METHODS

2.1. Sample Collection

A total of 597 blood samples were collected in a well known diagnostic lab and its collection centres located in different remote areas of Gujrat (a district of province Punjab, Pakistan). Informed consent was added in laboratory software from each patient including name, sex, age, address and contact number. Blood samples were collected in 5 mL chilled, sterilized disposable syringes. Serum separation was done in 1.5 mL eppendorf tubes at 13,000 rpm for 2 min and used fresh or stored at −20°C. All analysis was done in accordance with the ethical standards mentioned in the declaration of Helsinki.

2.2. RNA Extraction and Quantification

Total viral RNA was extracted from 150 μL serum samples by using Nucleospin RNA virus extraction kit according to the manufacturer’s instructions. Finally each RNA sample was dissolved in 50 μL TE buffer.

HCV RNA was amplified by using real time amplification and detection kit HCV Real-TM Quant (Sacace Biotechnologies, Italy). Fluorescently labelled HCV specificprobes were used (reporter dye: cy3). FAM dye was used to amplify HCV internal control (IC). Internal control was provided as an amplification control for each sample being processed to detect the possible contamination risks. Two positive controls for different quantities (quantities of standard 125 IU/mL and 12,500,000 IU/mL) and two negative controls were carried out using cDNA as substrate and spiked with mimic target in each run. From the total RNA extracted, 12.5 μL was used as template and mixed with equal volume of super mix containing 300 μL of RT-PCR-mix-1, 200 μL of RT-PCR-mix-2, 20 μL of Hot Start Taq Polymerase and 10 μL of M-MLV Revertase. Amplification was carried out by standard method (Stage 1: 50°C for 1800 sec, stage 2: 95°C for 900 sec and stage 3: 95°C for 20 sec and 60°C for 40 sec for 42 repeats. PCR reaction was performed in Smart cycler 11 (Cepheid, Germany). Results were analyzed by using smart cycler software v2.0d. Viral load was calculated with the help of Cy3 and FamCt value (Cy3 STD/Fam STD × 510,000 = No. of HCV IU/mL).

2.3. Assessment of ALT and AST

ALT and AST levels were measured using Ecoline reagents according to manufacturer’s instructions. The threshold of 42 IU/L was used as upper normal range for ALT and 40 IU/L for AST. Both enzyme activities were measured at the same time when PCR for HCV quantification was run.

2.4. Statistical Analysis

Relationship of viral load with ALT and AST in both males and females was analysed by determining Correlation Coefficient. Correlation was significant at 0.01 level (2-tailed). Correlation coefficient was performed by using commercially available computer program SPSS (SPSS16.0).

3. RESULTS

A total of 597 proposed anti HCV samples (171 detected
by ELISA and 426 by ICT) were collected in Dr. Muja- hid’s lab and FNA clinic district Gujrat and from its different collection centres. Out of 597 individuals 374 were females and 223 were males. All the individuals were categorized in three age groups (1st group: up to 25 years, 2nd group: 26 - 50 years, 3rd group: Above 50 years).

Since anti HCV antibodies are unable to show active HCV infection in some cases therefore, all the samples were subjected to RNA extraction and RT PCR to check active infection. The PCR results detected HCV RNA in the serum of 441 (73.87%) individuals. In males, 164 (73.45%) of 221 showed active HCV infection due to presence of HCV RNA in their serum while in females 277 (74.06%) of 374 had active infection. Quantification analysis further demonstrated that that population in 3rd age group is at higher risk then the other two groups where more than 70% of people have viral titre higher than 800,000 IU/mL (Table 1). Distribution of active HCV infection according to the sex and age is given in Figures 1(a) and (b).

Liver function tests (ALT and AST) were performed for the individuals with active infection. Among 441 confirmed patients of active HCV, 317 (71.88%) had elevated ALT level and 297 (67.35%) had elevated AST level (Table 2).

The correlation between viral load and ALT is more significant than the correlation between viral load and AST in females. Similarly, in case of males viral load and ALT had more significant relation than viral load and AST. Generally, males have strong relationship of ALT and AST with viral load than females (Figures 2(a) and (b)).

Table 1. HCV RNA viral load distribution in males and females.

<table>
<thead>
<tr>
<th>Age</th>
<th>&lt;800,000 IU/mL</th>
<th>&gt;800,000 IU/mL</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>≤25</td>
<td>≥25</td>
</tr>
<tr>
<td>Females</td>
<td></td>
<td></td>
</tr>
<tr>
<td>26 - 50</td>
<td>34.78% (8/23)</td>
<td>65.21% (15/23)</td>
</tr>
<tr>
<td>&gt;50</td>
<td>41.63% (92/221)</td>
<td>58.37% (129/221)</td>
</tr>
<tr>
<td>Males</td>
<td></td>
<td></td>
</tr>
<tr>
<td>26 - 50</td>
<td>31.58% (36/114)</td>
<td>68.42% (78/114)</td>
</tr>
<tr>
<td>&gt;50</td>
<td>28.57% (8/28)</td>
<td>71.43% (20/28)</td>
</tr>
</tbody>
</table>

Table 2. Comparison of liver function tests with HCV viral load.

<table>
<thead>
<tr>
<th>HCV RNA titer (IU/mL)</th>
<th>No. of patients</th>
<th>Serum ALT level (mean ± SD)</th>
<th>Serum AST level (mean ± SD)</th>
<th>Serum abnormal ALT (%age)</th>
<th>Serum abnormal ALT (%age)</th>
</tr>
</thead>
<tbody>
<tr>
<td>≤10^4</td>
<td>89</td>
<td>47.30 ± 27.69</td>
<td>44.02 ± 23.44</td>
<td>37.08 (33/89)</td>
<td>35.95 (32/89)</td>
</tr>
<tr>
<td>10^5</td>
<td>96</td>
<td>74.22 ± 53.57</td>
<td>59.52 ± 36.41</td>
<td>76.04 (73/96)</td>
<td>65.62 (65.62)</td>
</tr>
<tr>
<td>10^6</td>
<td>161</td>
<td>87.34 ± 73.35</td>
<td>70.21 ± 47.91</td>
<td>74.53 (120/161)</td>
<td>68.32</td>
</tr>
<tr>
<td>10^7</td>
<td>79</td>
<td>113.40 ± 81.43</td>
<td>87.49 ± 47.57</td>
<td>88.61 (67/79)</td>
<td>75.95 (60/79)</td>
</tr>
<tr>
<td>10^8</td>
<td>15</td>
<td>144.80 ± 101.21</td>
<td>120.13 ± 67.01</td>
<td>93.33 (14/15)</td>
<td>86.66 (13/15)</td>
</tr>
<tr>
<td>10^9</td>
<td>1</td>
<td>128</td>
<td>128 ± 0</td>
<td>100 (1/1)</td>
<td>100 (1/1)</td>
</tr>
</tbody>
</table>

Figure 1. Distribution of active HCV Infection in (a) Males; (b) Females.
4. DISCUSSION

According to world health organization, there is about 3% world’s population infected with hepatitis C in the world. Pakistan is one of the many developing countries where Hepatitis C is budding as a major health issue [19, 20]. Among the major causes of this dilemma are the absence of good preventive measures e.g. extensive reuse of non-sterilized syringes and razors, fragile health structure, unscreened blood transfusion, general poverty and poor education and high burden of immigrants have all affected epidemiology of HCV in Pakistan. Punjab is second largest province of Pakistan and has largest population. Gujrat is a densely populated district of Punjab located between two well-known rivers, the Jhelum and the Chenab. It is encircled on the northwest by the Jhelum River, on the northeast by Jammu and Kashmir and southeast by Chenab River and on the west by district Mandi Bahauddin [21].

Although Pakistan is at a high risk by HCV infection, however, national level estimates of the prevalence of and risk factors for hepatitis C are currently not available. Many scientists tried to explore the rate of HCV infection and found that it varies among different population groups and sexes like it is moderate in the general population but very high in injecting drug users and multi-transfused populations [22]. Similarly Ali et al. [23] studied moderate to high prevalence and risk factors of hepatitis C in different areas of Pakistan. A weighted average of hepatitis C antibody was 3.0% (range 0.3% - 31.9%) that is even higher in the high-risk subgroups. Same is concluded by Umer et al. [24] who reviewed that the frequency of hepatitis C infection in Pakistan is high (4.7%), varying from 0.4% - 33.7%, indicating pockets of infections. In present study we found the viral loads in relation with liver aminotransferases in active HCV infected individuals of both sexes constituting different age groups. All the patients were anti HCV positive. The results clearly indicated that all the anti HCV individuals had not active HCV infection. Our results indicated that there is a relation between liver aminotransferases and viral load. One case study conducted in district Mansehra Pakistan revealed there was 7% population having HCV antibodies while only 3.5% population had active HCV infection [25].

We studied a total 597 anti HCV patients of which 441 (73.87%) had active infection. The purpose of our study was to check active HCV infection from anti HCV individuals by quantifying their viral load and to also verify the relationship of elevated liver aminotransferases with active HCV infection. We found that 150 of 171 anti HCV individuals detected by ELISA had active infection while 291 of 426 patients detected by ICT had active infection. This clearly illustrated that ELISA is better method for antibody detection than ICT because it gives less false positive results. In case of males, we found that 112 (68.29) had high viral load and 52 (31.70%) had low viral load in their serum, in females 168 (60.64%) had high viral load and 109 (39.35%). As if viral load is less than 800,000 IU/mL, it is considered as low viral load while more than 800,000 IU/mL, is called high viral load [15].

Abnormal level of serum transaminases may lead to
the prognosis of HCV infection. One study showed that serum ALT levels, especially the AST levels were coupled with liver damage [26]. Usually it is considered that patients with chronic HCV having high HCV RNA titers and abnormal ALT levels show active HCV reproduction in liver and have higher risk of liver injury. Moreover, ALT level is used for indication of histological liver damage and a factor for starting therapy or to check antiviral therapy response in chronic hepatitis C [27]. We found that 71.88% patients with active HCV infection had elevated ALT level in their sera. As there are almost 25% of patients with active HCV infection having persistently normal ALT values [18]. We found 67.35% individuals with active HCV infection having elevated AST levels.

Patients with low viral load have 2.7 times more incidence of SVR than individuals with high viral load [28]. It is also prominent that individuals with high viral load show decreased response to antiviral therapy than individuals with low viral load and early prognosis reveals low HCV RNA viral load. Patients with high viral load should be treated for 24 weeks and those with low viral load might be treated for 16 weeks if there HCV RNA PCR is undetectable at 4th week of treatment [29,30]. Our results indicate that 36.5% (161/441) have low viral load while 63.49% (280/441) have high viral load.

5. CONCLUSION

The prevalence of active HCV infection in anti HCV individuals increases with the increase in age. Individuals up to 25 years had less number of anti HCV antibodies and HCV RNA in their blood while with the increase in age the chances of active HCV infection increase and viral load also increases. Individuals with age >50 years have highest percentage of HCV RNA titer in anti HCV individuals. The viral load has a strong relationship with liver enzymes (ALT and AST). With the increase in viral load, the ALT and AST levels also increase in most cases but in some cases of active HCV infection ALT and AST levels remain normal.

6. ACKNOWLEDGEMENTS

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REFERENCES


ABBREVIATIONS

ALT: Alanine aminotransferase
AST: Aspartate aminotransferase
Ct: Cycle threshold
HCV: Hepatitis C virus

IC: Internal control
ICT: Immunochromatographic test
M-MLV RT: Moloney murine leukemia virus reverse transcriptase
PEG-IFN: Pegelated interferon