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Original Research Article

Seroprevalence of HIV & HCV coinfection among patients in a rural tertiary care centre in North India

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Abstract

Objective: Viral hepatitis C infection is associated with high morbidity and mortality rates. Chronic HCV infection can cause a wide spectrum of liver disease, potentially leading to severe liver damage, including cirrhosis, organ failure and hepatocellular carcinoma. It accounts for nearly 12–32% of all cases of liver cancer and 10-20% cases of cirrhosis of liver, both of which have high treatment costs and poor outcomes. As route of infection of Human Immunodeficiency Virus, Hepatitis B Virus (HBV) and the Hepatitis C Virus (HCV) is same, People with HIV infection are at more risk of acquiring Hepatitis B Virus (HBV) and the Hepatitis C Virus (HCV) infection. The co-infection of Hepatitis C virus with HIV accelerates disease progression and also has an effect on the management of patients infected with HIV. The prevalence of HIV co-infection with Hepatitis C virus varies widely. This study is planned to evaluate the prevalence of HIV co-infection with Hepatitis C viruses in North India. **Materials and Methods:** A total of 25443 patients enrolled in the microbiology lab were retrospectively analyzed for the presence of HCV and HIV infection on the basis of the presence of anti-HCV and anti HIV markers. **Results:** In patients infected with HIV, the prevalence of co-infection with HCV was 0.07%. The mean age of the study group was 28.7 years. **Discussion:** The prevalence rate of HCV are increasing in patients infected with HIV. Having acquired the knowledge about the importance of such a co-infection, it is essential that all the patients infected with HIV be screened for HCV co-infection.

Keywords: HIV, HCV, Seroprevalence, Co-infection

Introduction

Hanatitis C virus (HC

Hepatitis C virus (HCV) is a single-stranded RNA virus. Viral hepatitis C infection is associated with high morbidity and mortality rates. Chronic HCV infection can cause a wide spectrum of liver disease, potentially leading to severe liver damage, including cirrhosis, organ failure and hepatocellular carcinoma. It accounts for nearly 12–32% of all cases of liver cancer and 10-20% cases of cirrhosis of liver, both of which have high treatment costs and poor outcomes [1].

As route of infection of Human Immunodeficiency Virus, Hepatitis B Virus (HBV) and the Hepatitis C Virus (HCV) is same, People with HIV infection are at more risk of acquiring Hepatitis B Virus (HBV) and the Hepatitis C Virus (HCV) infection. The epidemiological parameters of these viruses are quite similar, like routes of transmission, there associated risk factors and the

Manuscript received: 14th November 2018 Reviewed: 24th November 2018 Author Corrected: 30th November 2018 Accepted for Publication: 5th December 2018 presence of these viruses in various body fluids [2]. With increased availability of antibiotics and antifungal drugs, HBV and HCV infection are becoming a cause for significant concern for individuals infected with HIV. The co-infection of HCV with HIV is associated with a loss of immunological control of HCV thus leading to rapid progression of HCV disease [3]. In coinfection, one virus affects the life cycle of the other virus which is present simultaneously in the same host.

Persons who inject drugs (PWID) are at highest risk of acquiring HIV and HCV coinfection as these viruses have same route of infection. HCV is 10 times more infectious than HIV through percutaneous blood exposures. Other high-risk factors of HIV and HCV coinfection are people who take injections, people who are immunocompromised, vertical transmission from infected mother to child. People with HCV infected sexual partners. People with HIV infection, in particular MSM, are at higher risk of HCV infection trough

unprotected sex. Tattoo recipients are also at higher risk of acquiring HCV infection. Out of total population of PWID, 67% are infected with HCV worldwide. [4] Transmission through blood transfusions is rare now a days due to rapid and regular screening of blood and blood products Of the estimated 36.7 million people living with HIV globally, around 2.3 million have past or present HCV infection, and of those, around 1.36 million are people who inject drugs [4].

In HIV coinfection with HCV, HIV accelerates the natural course of HCV infection, leading to faster progression of liver disease to cirrhosis finally causing hepatocellular carcinoma.

Anti-retroviral therapy has led to decline in death rate due to AIDS but liver diseases caused by HCV coinfection are one of the major cause of non AIDS related deaths of HIV patients.[5]

Universal screening of Hepatitis C in HIV infected patients is highly recommended for the treatment purpose. First the patient should be checked for HCV antibodies. If antibodies are positive then confirm the presence of HCV in patients' blood. Positive result confirms the presence of HCV. [fact sheet]

The present study was conducted to detect the current seroprevalence of HCV co-infection in patients infected with HIV in a Rural Tertiary Care hospital in North India.

Material and Methods:

Study design: Prospective study

Study site: Microbiology laboratory of SGRDIMSR, Amritsar.

Duration of study: November 2017 to October 2018

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Inclusion criteria

Patients with clinical history and signs and symptoms suggestive of an immuno-compromised condition pre-operative patients

Antenatal patients

Exclusion criteria: Patients who were seronegative for HIV or HCV

Sample collection: 5ml of blood sample collected using a sterile plain vacutainer, under all aseptic precautions.

Sample processing: 5 ml blood was collected in a clean dry sterile vial and allowed to clot or serum separated by centrifugation at room temperature. If serum isnot to be assayed immediately then stored at 2-8°C or frozen at minus 20°C (-20°C).

The marker used for screening of HIV was antibodies to HIV-1 & HIV-2 and was detected by one rapid test and a solid phase enzyme linked immunosorbant assay (ELISA), which was based on direct sandwich principle (HEPALISA- manufactured by J. Mitra and co Pvt Ltd).

For HCV, one rapid test and anti HCV (IgG) ELISA was done using 3rd generation ELISA kit (HCV Microelisa by J. Mitra and co pvt Ltd). For HIV tri-dot rapid visual test for the qualitative detection of antibodies to HIV-1 & HIV-2 in human serum/plasma used. (Diagnostic enterprises) HCV tri-dot rapid visual test for the qualitative detection of antibodies to Hepatitis C virus in human serum/plasma was used. (Diagnostic enterprises).

The tests were performed as per the manufacturer's instructions. ELISA test was done as per the manufacturer's instructions along with validity check and incorporation of internal controls.

Results

The present study received total 25443 samples in the microbiology lab for HIV and HCV detection together. Out of these 9136 presented for HIV detection. Total HIV positive patients were 0.73% (67). Total patients presented for HCV detection were 16307. Out of these 3.23% (527) patients were reactive for HCV. The prevalence of HCV coinfection with HIV was 0.07% (19)

Table No-1: Prevalence of HCV in HIV reactive patients.

Samples received	Tested	Reactive
FOR HIV	9136	67 (0.73%)
FOR HCV	16307	527 (3.23%)
Total (HIV+HCV)	25433	19 (0.07%)

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Table No-2: The distribution of HIV+HCV co-infection by age and gender in the study group

Age	Male	Female
<10	1	0
10-20	0	0
20-30	10	0
30-40	01	1
40-50	02	1
50-60	02	0
60-75	01	0
Total	17	2

The mean age of the study group was 28.7 years.

Discussion

India has the third largest number of individuals with HIV after South Africa and Nigeria [6]. Coinfection of HIV with the HCV is a common event due to similar route of transmission [6]. India which contributes significantly to the global HCV burden, is believed to harbour 12–18 million HCV-infected people. The estimated prevalence of HCV in India is 0.5%–1.5% [7].

All the HIV positive patients should be screened for HCV coinfection. HCV antibodies appear between six weeks to six months of infection. Screening is done with anti HCV elisa test.

It is noted in the present study that the age of the study group is 20-30 years, which is less as compared to previous studies from India [7] while in accordance with the study by Kalyani et al [3]. The mean age of the study group is 28.7 years which is slightly lower than that in the study by Ahuja et al. Thus, the younger population in the economically productive age group is being increasingly affected by HIV, leading to a loss to the economy. This could be because of the increased exposure of this population to the risk factors like promiscuity, parenteral drug abuse, etc.

Among the patients presenting for treatment of either HCV or HIV infection, HIV & HCV coinfection is emerging as important and more frequent finding. Similar route of transmission appears to be the most important factor for higher prevalence rate of coinfection. Coinfection with HCV leads to chronic liver damage in HIV patients and more than 15% developed severe liver damage or cirrhosis [8]. The present study shows that mainly male patients are infected, this is in accordance with previous study that shows male patients are at a higher risk of acquiring infection.

Present study reports HIV and HCV coinfection in 0.07% which is lesser than the study by Jain etal [1]. The HCV co infection in the study by Kalyani et al was 2%, while Oslanisun, Olefemi, Adewole et al [9] reported 2.3%. A study by Baveja et al has shown prevalence rate of 9.64% [10]. However, another study by Ponamagi et al shows that HIV infected patients are at a higher risk of coinfection as shown by the prevalence of HCV (3.02%) [8]. Prevalence rate among various cities in India is 2.3% in Chennai [11], 1.6% in Lucknow, whereas higher in Nagpur 7.2%, 8% from Mumbai and 8.3% from Hyderabad [3]. In another study from Nigeria 0% rate of coinfection was observed, as their study comprised mainly of blood donors [12].

Study from Iran showed very high 72% rate of coinfection [13]. However, in India, study by Mahajan.A et al showed 0% coinfection of HCV with HIV [14] while another study from North India by Tripathi AK etal observed 1.6% coinfection rate of HCV and HIV [15]. Seroprevalence in the present study is much lower than the result obtained in study performed by Ahuja et al (1.7%). Another study from Iran showed much higher (72%) coinfection rate of HIV & HCV [13] as these patients were mainly intravenous drug users.

Thus, various studies show that coinfection rates of HCV in HIV infected patients are variable worldwide depending on geographical region, risk groups, and also the type of exposure as seen in various studies. In India also HCV coinfection in HIV patients varies from area to area and varies from 0-8% [16]. The higher prevalence rate of HCV in HIV positive patients could be considered as noticeable and it could be attributed to diverse factors particularly lack of vaccine for HCV.

Also it is transmitted mostly via injection (especially in drug addiction) because of the increasing rate of addiction in certain countries [13]. It is found that HIV and HCV coinfection is higher in sexually active age group than other age groups. The present study reveals that male patients are mainly coinfected 89.4% (17/19), while the results by Kalyani et al differs which shows female predominance in their study [3]. The coinfection HCV with HIV is because of immunological suppression of control of HCV leading to more severe hepatitis C infection [15]. CTL & helper T cells are responsible for control of HCV infection. But loss of these cells in HIV infection can lead to HCV viremia

HCV antibodies testing alone is not adequate to diagnose chronic HCV infection as some patients spontaneously clear the virus without treatment but remain HCV positive. In such patients HCV viral load (HCV RNA) test is necessary. Also, it is noted that spontaneous viral clearance is less likely to occur among HIV positive patients.

Conclusion

[14].

HCV coinfection in HIV patients is of serious concern in India. The knowledge of HCV coinfection in patients with HIV is important as the life expectancy of these patients is prolonged with ART and they need to be managed for HCV coinfection simultaneously. Routine screening of these HCV should be mandatory for HIV infected patients as there is increased chance of coinfection with hepatitis viruses because of same route of transmission and immunosuppression of such patients.

What the study adds to the existing knowledge?

The present study underlines the necessity of a uniform guideline in which all the patients infected with HIV should be screened for HCV to help the management of co infection.

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