Hepatitis C, Delta and Human Immunodeficiency Virus Sero-Prevalence in Patients Chronically Infected with Hepatitis B Virus in Cotonou, Benin

Abstract
Infections with hepatitis viruses are known to be prevalent in sub-Saharan Africa but in Benin, data on these infections are still scarce. The study was carried out to assess the burden of these infections in Cotonou. From June to October 2016, we conducted a cross-sectional study on 156 Hepatitis B Virus (HBV) infected patients attending the University Teaching Hospital and a private hospital in Cotonou. For each patient, HBV viral load measurement and Hepatitis Delta Virus (HDV), Hepatitis C Virus (HCV) and Human Immunodeficiency Virus (HIV) serology tests were performed using standard methods. The median HBV viral load was 557.5 UI/mL and HDV, HCV and HIV sero-prevalence rates were 3.9%, 1.3% and 0.7% respectively. In conclusion, in HBV infected patients in Cotonou, the sero-prevalence rates of HDV, HCV and HIV were relatively low. A national prevalence survey is needed to assess the epidemiology of these infections in the country.

Keywords: HBV; HCV; HDV; HIV; Cotonou

Introduction
According to the World Health Organization (WHO), about 257 million people live with Hepatitis B virus (HBV) worldwide. Sub-Saharan Africa represents an area of high prevalence of this infection with more than 8% of chronic carriers [1]. In addition, HBV infection is a major cause of deaths in this region due to complications such as cirrhosis and hepatocellular carcinoma [1]. Hepatitis delta virus (HDV) is a defective virus that depends on the HBV surface antigen (HBsAg) for its existence. Therefore, infection with HDV requires necessarily HBV infection, either simultaneous infection with both viruses or HDV infection in a subject previously infected with HBV. Compared with infection with HBV alone, HDV infection leads to more rapid progression of liver disease to cirrhosis and hepatocellular carcinoma [2-4]. The number of people infected with HDV is estimated at 15 to 20 million worldwide [5]. In spite of this information, there is scarcity of data on the disease estimate in many endemic areas particularly in sub-Saharan Africa. HBV and HDV share with the hepatitis C virus (HCV) and human immunodeficiency virus (HIV) the same blood transmission route. In addition, co-infection with HIV and/or HCV worsens the liver disease in HBV and/or HDV infected patients. Data on HCV infection is rare in sub-Saharan Africa but a review from Riou et al. [6] ranged the prevalence in this region from 0.0% to 56.0% [6]. Since HIV infection is endemic in Africa region, the occurrence of HIV co-infection with hepatitis viruses may also be on a high side. In Benin, data on co-infection with HDV, HCV and HIV in HBV infected patients are rare. To our knowledge, the only two published studies on co-infection with these viruses were on a limited number of patients in one hospital in the Northern part of the country [7,8]. We aimed in the present study to assess the prevalence of HDV, HCV and HIV infections in patients chronically infected with HBV in Cotonou, the largest city in Benin.

Materials and methods
Setting
Benin is a country with a size of 114,763 square kilometers and an estimated population of 10.9 million [9]. Cotonou is the biggest city in the country with approximately 673,000 populations in 2013 [10].

Subjects
This cross-sectional study was conducted from June to October 2016 in two hospitals in Cotonou: the university teaching hospital, Hubert Koutoukou Maga, which is a public hospital and the Atinkamney polyclinic, the biggest private hospital in the city. In total, 156 subjects (15 years and above) were consecutively included in the study. These subjects were previously detected positive with HBs antigen and followed up in these hospitals. None of them was under treatment for HBV infection.

Abbreviations: CI: Confidence Interval; EDTA: Ethylene Diamine Tetra Acetic Acid; HBV: Hepatitis B Virus; HCV: Hepatitis C Virus; HDV: Hepatitis Delta Virus; HIV: Human Immunodeficiency Virus; WHO: World Health Organization
Sample

Venous blood was collected from each subject into two tubes. Ethylene diamine tetra acetic acid (EDTA) tubes were used to collect blood for plasma separation (viral load measurement) while plain tubes were used for serum separation (serology). All tests were performed and interpreted according to manufacturer’s instructions. Anti-HDV antibody was detected using ETI-AB-DeltaK-2® kit (Diasorin, France) while in positive samples, anti-HDV IgM were detected with ETI-AB-DeltaGA-MC-2® kit (Diasorin, France). HCV testing was performed using immunochemiluminescence anti-HCV® II test (Roche Diagnostics, USA) or VIDAS Anti-HCV® Assay (bio Merieux, France). HIV serology was performed using Alere Determine HIV-1/2® test (Alere Medical, Japan) for screening while samples that were reactive were confirmed by Immuno Comb HIV 1&2 BiSpot® (Orgenics, France). HBV viral load measurement was carried out using Cobas TaqMan® 48 kit (Roche Diagnostics, USA).

Ethical considerations

All patients gave informed consents and the study was approved by the institutional review board.

Data analysis

Data were collected using Epi Data version 3.1 and statistical analyses were performed using Stata software version 12.0.

Results

In total, 156 HBV infected patients were included in the study. Their characteristics are presented in (Table 1). Median age of patients was 36.0 years with a male to female ratio of 2.0:1.0. The median HBV viral load was 557.5 UI/mL while in two thirds of the patients (67.3%), the HBV viral load was less than 2,000 UI/mL; among them, three had undetectable HBV viral load (Table 1). Of the 156 HBV infected patients, six (3.9%) were screened positive for anti-HDV antibodies of which, two (33.3%) were positive for anti-HDV IgM antibodies. In HBV infected patients, HCV and HIV sero-prevalence rates were 1.3%, 95% CI = 0.5% - 3.1% and 0.7%, 95% CI = 0.6% - 2.0% respectively (Table 2). No HBV infected patient was co-infected with the other three viruses.

Table 1: Characteristics of patients included in the study.

| Patients | Number | Percentage |
|----------|--------|------------|
| Gender   |        |            |
| Male     | 104    | 66.7       |
| Female   | 52     | 33.3       |
| Age (years) |    |            |
| 15-24    | 21     | 13.5       |
| 25-34    | 46     | 29.5       |
| 35-44    | 49     | 31.4       |
| 45-54    | 23     | 14.7       |
| ≥55      | 17     | 10.9       |
| HBV viral load (UI/mL) |    |            |
| <2,000   | 105    | 67.3       |
| ≥2,000   | 51     | 32.7       |

Discussion

Information on HDV infection is rare in Africa. However, the magnitude of the problem needs to be assessed in a particular setting in order to set up appropriate measures to tackle the disease. In this study, the sero-prevalence of anti-HDV antibody was 3.9%. This finding was in agreement with 3.0% and 3.2% found in Senegal and Uganda respectively [18,19]. This observation showed that the risk of having an active HDV infection. However, one cannot exclude active HDV infection in patients with negative anti-HDV IgM since a more sensitive test (HDV PCR) was not performed. This is the main limitation of this study. In our study, HIV sero-prevalence was 0.7% similar to what was observed in the general population (1.1%) [17].

Several studies in sub-Saharan Africa had investigated HBV infection in HIV infected patients and showed a higher co-infection rates of 16.8%, 13.6% and 11.1% in Senegal, Ghana and Benin respectively [18,19]. This observation showed that the risk of HBV infected patients to be infected with HIV was lower than that of HIV infected patients to be infected with HBV. Possible explanations for this are the facts that most HBV infected patients

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in our setting are infected during infancy and their characteristics regarding HIV infection are similar to those of the general population. In contrary however, HIV infected patients represent a specific group with an increased risk of sexual transmission for both HIV and HBV. HCV sero-prevalence was 1.3% in our study. Previous studies within the country showed a prevalence ranging from 0.7% in Cotonou to 7.4% in pregnant women in Tangueta [B, 20]. In other countries in Africa, reports showed that HCV prevalence were generally relatively low [6]. This is fortunate since both viruses cause liver damages leading to cirrhosis and hepatocellular carcinoma. Common effects of both viruses may accelerate progression to these complications. In contrast to HIV/AIDS, there are no national programs to address policies on viral hepatitis in most sub-Saharan Africa countries including Benin despite the high prevalence of these infections in the region [1,5,6,17]. As a result, epidemiological data available on hepatitis infections are either scarce or not representative of the population of patients infected. The present study contributes in getting a picture of the situation and may help in setting up measures to fight against these viruses [21,22].

Conclusion

In HBV infected patients in Cotonou, sero-prevalence rates of HDV, HCV and HIV infections were 3.9%, 1.3% and 0.7% respectively. These rates seem lower than what was previously reported in the northern part of the country. A national prevalence survey is urgently needed to determine the actual picture of the epidemiology of these infections in the country.

Conflict of interest

Authors declare no conflict of interest.

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