### **RESEARCH ARTICLE**

**Open Access** 

# Epidemiology of hepatitis B virus infection in people living in poverty in the central-west region of Brazil



Lara Cristina da Cunha Guimarães<sup>1</sup>, Sandra Brunini<sup>1</sup>, Rafael Alves Guimarães<sup>2</sup>, Hélio Galdino-Júnior<sup>1</sup>, Ruth Minamisava<sup>1</sup>, Vanessa Elias da Cunha<sup>1</sup>, Jordana Rúbia Souza Santos<sup>1</sup>, Elisângela de Paula Silveira- Lacerda<sup>3</sup>, Christiane Moreira Souza<sup>1</sup>, Vera Lúcia Brandão de Oliveira<sup>4</sup>, Gabriela Cavalcante Albernaz<sup>1</sup>, Thiago Guida de Menezes<sup>5</sup> and Giovanni Rezza<sup>6\*</sup>

#### **Abstract**

**Background:** People living in poverty (PLP) are highly vulnerable to hepatitis B virus (HBV) infection. This study aimed to investigate the epidemiology of HBV infection in PLP in the metropolitan region of Goiânia, Goiás State, in the Central-West Region of Brazil.

Methods: A cross-sectional study was conducted from August to December 2016 in adults aged ≥12 years living in poverty. The following serological markers for HBV were investigated: hepatitis B surface antigen (HBsAg), antibody to HBV core antigen (total anti-HBc), IgM anti-HBc, and hepatitis B surface antibody (anti-HBs), which were detected by enzyme-linked immunosorbent assay (ELISA). Poisson regression analysis with robust variance was performed to verify the factors associated with HBV exposure.

**Results:** The study included 378 participants. The overall prevalence rate of HBV (any viral marker) was 9.8% (95% confidence interval [CI], 7.2–13.2). The prevalence rate of HBsAg in combination with total anti-HBc was 0.8% (95% CI, 0.3–2.4), total anti-HBc in combination with anti-HBs was 7.7% (95% CI, 5.4–10.9), and total anti-HBc alone was 1.3% (95% CI, 0.5–3.0) in the population. Furthermore, isolated positivity for anti-HBs was identified in only 25.4% (95% CI, 21.3–30.0) of the participants. Multiple regression analysis revealed that age (adjusted prevalence ratio [APR], 1.04; 95% CI, 1.01–1.07), female sex (APR, 2.18; 95% CI, 1.01–4.73), sexual intercourse under the influence of alcohol (APR, 2.49; 95% CI, 1.36–7.06), and exposure to *Treponema pallidum* (APR, 3.10; 95% CI, 1.36–7.06) were associated with HBV exposure.

**Conclusion:** There was a high prevalence of HBV exposure in PLP in the Central-West Region of Brazil, indicating significant viral spread of the infection. Additionally, there was low serological evidence of immunisation against hepatitis B, indicating that a large proportion of the participants in this study are susceptible to the infection. The results support the need for public health policies that facilitate access to the existing healthcare services in hard-to-reach groups with special regard to immunisation programmes against hepatitis B.

Keywords: Hepatitis B infection, Poverty, Epidemiology, immunisation

<sup>&</sup>lt;sup>6</sup>Department of Infectious Diseases, Istituto Superiore di Sanitá, Rome, Italy Full list of author information is available at the end of the article



<sup>\*</sup> Correspondence: giovanni.rezza@iss.it

Guimarães et al. BMC Public Health (2019) 19:443 Page 2 of 11

#### **Background**

Hepatitis B virus (HBV) infection is considered a serious public health problem worldwide, especially in less developed countries. It is estimated that 70% of new chronic HBV infections occur in low-income countries [1]. It is also estimated that 257 million people worldwide are chronic carriers of HBV and that there were 887,220 deaths as a result of the infection in 2015, most of which are related to complications such as cirrhosis and hepatocellular carcinoma [2].

Although Brazil has a low prevalence of HBV infection, it has the second largest population of individuals with positive hepatitis B surface antigen (HBsAg), accounting for 16.7% of all infections from the American Region [3]. In 2010, a population-based study conducted in adults aged 20–69 years in all macro-regions of Brazil estimated the prevalence of HBV exposure (11.6% positivity for total antibody to HBV core antigen [anti-HBc]) and infection (0.6% positivity for HBsAg). The lowest and highest prevalence of exposure were found in the Southeast (7.9%) and North (14.7%) macro-regions, respectively [4].

In a modelling study, performed with a Delphi process that included a literature review of PubMed and Embase, followed by interviews with experts, it was found that the prevalence rate of HBV infection (HBsAg positive) in the population of Brazil was 0.6%, where only 28% individuals were diagnosed (212,000) and only 12% of the population was treated. In children up to 5 years of age, the prevalence rate of infection is < 0.1%, which may be related to the high rates of vaccination coverage; 86% of children living in Brazil complete the vaccination schedule before reaching 1 year of age [5].

HBV can be transmitted by parenteral, sexual, vertical, and horizontal routes. Thus, individuals who engage in risky behaviours, such as the use of psychoactive substances, unprotected sex, multiple sexual partners, early initiation of sexual activity, and sharing of personal objects, are more vulnerable to HBV infection [6]. In addition to behavioural risk factors, social determinants such as low income and low level of schooling may contribute to increased exposure to HBV [6, 7].

People living in poverty (PLP) are significantly vulnerable to infectious diseases in general, and hepatitis B in particular, due to poor living conditions and/or difficulties in accessing healthcare. The concentration of poverty is associated with scarcity of basic services [8], resources [9], education [10], and employment opportunities, resulting in deterioration of quality of life of the residents [11, 12]. Moreover, the absence of primary healthcare services makes it difficult to access preventive services [13]. Lack of educational opportunities may limit the access to sexually transmitted infection (STI) prevention courses that are generally offered in schools.

Furthermore, economic factors such as unemployment and poverty can promote risky behaviours such as inconsistent condom use and multiple sexual partnerships [14, 15], thus increasing an individual's risk of acquiring an STI, such as hepatitis B [8].

In 2016, Brazil was ranked 96 in Healthcare Access and Quality Index by the Global Burden of Disease [16], setting up a country with great social inequalities and with a high number of PLP. According to the Brazilian Institute of Geography and Statistics (in Portuguese Instituto Brasileiro de Geografia e Estatística - IBGE) data, 16.27 million people were living in poverty in 2010, representing 8.5% of the country's total population. Of these, 53.3% lived in urban areas, where the majority of the Brazilian population resides (84.4%) [17]. In Latin America, Brazil is one of the countries with the highest prevalence of HBV associated with the lower socioeconomic class [18]. Investigations among PLP have shown variations in the prevalence rate of HBV exposure in Brazil, ranging from 5.9% in adolescents in Goiânia (Goiás State, Central-West Region) [19] to 40.7% in the population of Buriticupu (Maranhão State, Northeast Region) [20].

However, studies that address exposure to HBV and factors associated with HBV in the general PLP population are still lacking, especially in the Central-West Region of Brazil. Most of the studies published to date address specific population groups (waste collectors, homeless, and immigrants) or do not present analytical approaches to the determinants associated with HBV. Thus, there is a gap in the literature on the epidemiology of hepatitis B in PLP using home-based and populationbased surveys. This study intended to contribute to the monitoring of the prevalence of hepatitis B and to the understanding of the exposure profile and associated risk factors in Brazilian PLP. This is necessary to implement actions, services, and policies that meet the specific need of this population. The present study aimed to investigate the epidemiology of hepatitis B infection in PLP of the metropolitan region of Goiânia, Goiás State, Central-West Region of Brazil.

#### Methods

#### Design, population, and study sites

This is a cross-sectional study conducted among PLP in the metropolitan region of Goiânia, the capital of Goiás State, Central-West Region of Brazil, which has a population of 1,466,105 inhabitants.

In the metropolitan region of Goiânia, there is an area where, until 2013, all the wastes generated in Goiânia were destined and accumulated without any separation of common or infectious wastes. Over the years, a population group settled in favelas in the adjacent areas, living in precarious conditions and working with the collection of

Guimarães et al. BMC Public Health (2019) 19:443 Page 3 of 11

these wastes. Since the closure of the zone and the creation of the sanitary landfill, some families received masonry housing from the government and were transferred to a new living space (Neighbourhood 1, Fig. 1 a, b). Other families remained in the area (Neighbourhood 2, Fig. 1 c). Still, families who moved to neighbourhood 1, despite living in masonry homes and geographically closer to public health and education facilities, remained without access to basic sanitation and paved streets where sewage runs in the open. Therefore, these two neighbourhoods were selected in this study because they reflect signs of extreme poverty within the metropolitan region and have in common, beyond the social precariousness, the history of cohabitation with the waste.

According to Kageyama and Hoffman [21], income is an unreliable indicator of living conditions that characterise poverty due to inaccurate information provided by the people. Thus, it is necessary to resort to variables that are related to consumption expenditures as a more advantageous and specific method of identifying PLP [21].

In the neighbourhoods, there are a large number of canvas, cardboard, or wood residences without access to pipe-borne water, basic sanitation, and public lighting. There are no pavements in the streets. There are no family healthcare units or other healthcare services for the population. Under this scenario, reports of exploitation and child labour, abuse and domestic violence against women, the use of illicit drugs, and sex workers are frequent.

In this context, the criteria of the Brazilian Economic Classification (in Portuguese Critério da Classificação Econômica do Brasil - CCEB) were adopted for the selection of the participants for the present study [22, 23]. The CCEB is a system of proxy punctuation of the consumption capacity of a household, expressing the wealth of the residences and the purchasing power of the families. In this investigation, the CCEB 2015 version was used. This takes into account the possessions and quantity of household comfort goods (presence of automobiles, microcomputer, and digital versatile disc player, degree of schooling of the head of the family (illiterate to complete higher education), and access to public services (pipe-borne water and paved streets), which together represent a permanent family income construct [22, 23]. For each CCEB variable, a specific score ranging from 0 to 100 is assigned, where the higher the score, the higher the socioeconomic classification of the family. The CCEB system generates a score that stratifies families into the following socioeconomic classes: A (scores from 45 to 100), B1 (38-44), B2 (29-37), C1 (23-28), C2 (17-22), and D-E (0-16) [23]. Families who belong to D–E classification are families living in poverty.

In the present study, people aged 12 years or older who, according to the CCEB, belonged to classes D–E, that is, who had a score in the CCEB equal to or less than 16 points, were included. Participants with apparent cognitive impairment or major incapacitating psychic disorders were considered ineligible.



Fig. 1 Areas of study sites. a-b, Neighbourhood 1; c, Neighbourhood 2

Guimarães et al. BMC Public Health (2019) 19:443 Page 4 of 11

The sample size was calculated for the finite population, and a statistical power of 80% ( $\beta$  = 20%) with a significance level of 95% ( $\alpha$  = 0.05) and a global prevalence rate of HBV of 12.8%, according to Marinho et al. [24], with design effect correction of 2.0 were considered. Therefore, the minimum sample size needed to estimate the prevalence rate in the study population was 325 participants, which was increased by 10% to correct for loss and refusal, totalling a sample of 358 participants. This study included 378 participants.

#### Data collection

Data collection occurred between August and December 2016, and the recruitment occurred through home visits. All the houses in both neighbourhoods were visited. For each eligible household, a participant was drawn from the list of residents aged ≥12 years. Next, the selected resident was invited to participate in the study. In case of the absence of the resident at the time of the interview, at least three attempts to contact residents in each sampled home were made at different periods [25, 26]. There was no substitution in the event of an absence of the resident at the end of the third visit.

The data were collected through face-to-face interviews in the households, using a structured instrument to obtain socio-demographic data and other potential factors associated with HBV infection. After the interview, rapid testing for syphilis was performed locally, and 5 ml of blood was collected for serological tests for hepatitis B.

#### Laboratory analysis

HBV-related markers (HBsAg, total anti-HBc, IgM anti-HBc, and hepatitis B surface antibody [anti-HBs]) were detected by enzyme-linked immunosorbent assay (ELISA) according to each protocol. The results were delivered to the study participants during post-test counselling.

#### Study variables

#### Dependent variable

The presence or combination of the following HBV exposure markers was considered to be the endpoint variable: (i) HBsAg and total anti-HBc, (ii) total anti-HBc and anti-HBs, or (iii) total anti-HBc isolated.

#### Independent variables

The following independent variables were analysed:

- (i) Sociodemographic: age (years), gender (male or female) as gender proxy, schooling (full years of formal study), and marital status (married, separated/divorced/widower, or single).
- (ii) Non-sexual risk factors: tattoo (no or yes), body piercing (no or yes), sharing of personal objects (toothbrushes, pliers, razor, pliers, or nail clippers) (no or yes),

history of hepatitis in the family (no or yes), history of incarceration (no or yes), marijuana use (no or yes), use of intranasal cocaine (no or yes), crack use (no or yes), injectable drug use (no or yes), alcohol use (no or yes), and alcohol dependence (no or yes).

Alcohol dependence was assessed using the CAGE (an acronym for the following: 'Cut down, Annoyedby criticism, Guilty, and Eye-opener') questionnaire [27], which includes these four questions: (1) Have you ever felt you needed to cut down on your drinking? (2) Have people annoyed you by criticising your drinking? (3) Have you ever felt guilty about drinking? (4) Have you ever felt you needed a drink first thing in the morning (eye-opener) to steady your nerves or to get rid of a hangover? This questionnaire was used with a cut-off of two affirmative responses suggesting positive screening for alcohol abuse or dependence [27, 28]. In Brazil, CAGE assessment was validated in 1983 by Masur and Monteiro, who found a sensitivity of 88% and a specificity of 83% for the detection of alcohol dependence [29].

(iii) Sexual risk factors: use of condoms during sexual intercourse (always or never/sometimes), condom use at the last sexual intercourse (no or yes), practice of anal sex (no or yes), homosexual relationship (no or yes), sexual relations with sex workers (no or yes), sexual intercourse with illicit drug users (no or yes), sexual intercourse under illicit drug use (no or yes), sexual intercourse under the influence of alcohol (always or never/sometimes), number of sexual partners in life, number of sexual partners last year, age of first sexual intercourse, STI history (no or yes), and exposure to *Treponema pallidum* (negative or positive).

The exposure to *Treponema pallidum* (*T. pallidum*), the etiologic agent of syphilis, was evaluated by the rapid test, performed after the interview.

#### Statistical analysis

The data were analysed using the Stata software, version 14.0 [30]. Initially, the Anderson-Darling (AD) test was performed to verify the normality of the quantitative variables [31]. Three levels of analysis were then conducted: (i) descriptive, (ii) bivariate regression, and (iii) multiple regression.

In the descriptive analysis, quantitative variables were presented as median and interquartile range (IQR) due to the absence of normality in the AD test and qualitative variables in absolute and relative frequencies. The prevalence of HBV serological markers was estimated with 95% confidence interval (CI).

Subsequently, bivariate Poisson analysis was performed to verify the potential factors associated with HBV exposure. Variables that were included in the Poisson regression model were those with robust variance [14, 32, 33] with

Guimarães et al. BMC Public Health (2019) 19:443 Page 5 of 11

**Table 1** Sociodemographic characteristics and potential risk factors for hepatitis B among the study population (Goiânia, Goiás, Central-West Region of Brazil, 2016)

Variables N = 378% Age (years), median (IQR) 31.0 (21.0-42.0) Education (years), median (IQR) 7.0 (5.0-9.0) Sex Male 188 49.7 Female 190 50.3 Marital status Married 224 59.3 8.5 Divorced/separated/widowed 32 Single 122 32.3 Sharing of personal objects<sup>a</sup> 30.0 No 113 70.0 Yes 264 Tattoo No 257 68.0 Yes 121 32.0 Body piercing 427 88.9 No 11.1 Yes 41 History of hepatitis in the family No 281 87.8 39 Yes 12.2 History of imprisonment<sup>b</sup> No 290 81.0 19.0 Yes 68 Injecting drug useb No 374 98.9 4 Yes 1.1 Alcohol use<sup>c</sup> No 191 50.5 Yes 187 49.5 Alcoholic dependence No 246 69.3 Yes 109 30.7 Marijuana use<sup>c</sup> No 286 75.7 Yes 92 24.3 Intranasal cocaine use<sup>c</sup> No 331 87.6 12.4 Yes 47 Crack use<sup>c</sup> No 347 91.8 Yes 32 8.2

**Table 1** Sociodemographic characteristics and potential risk factors for hepatitis B among the study population (Goiânia, Goiás Central-West Benjon of Brazil, 2016) (Continued)

Variables	N = 378	%
STIs history <sup>d</sup>		
No	337	89
Yes	41	10.8
Treponema pallidum exposure		
Negative	342	90.
Positive	36	9.5
Condom use <sup>a</sup>		
Never	65	18.
Sometimes/always	280	81.
Condom use at last sexual intercourse		
No	261	76.
Yes	82	23.
Anal sex <sup>a</sup>		
No	215	63.
Yes	122	36.
Sexual relation homosexual <sup>a</sup>		
No	316	92.
Yes	26	7.6
Sexual intercourse with illicit drug users <sup>a</sup>		
No	171	49.
Yes	174	50.
Sexual intercourse with sex workers <sup>a</sup>		
No	265	77.
Yes	78	22.
Sexual intercourse with STIs carriers <sup>a</sup>		
No	313	91.
Yes	30	8.7
Sexual intercourse under the influence of alcohol	3	
Never	201	58.
Sometimes/always	144	41.
Age of first sexual intercourse, median (IQR)	15.0 (13.0–17.0)	
Number of sexual partners <sup>b</sup> , median (IQR)	5.0 (3.0–15.0)	
Number of sexual partners <sup>a</sup> , median (IQR)	1.0 (1.0–1.0)	

<sup>c</sup>Pevious month

dPevious 12 months

p-value < 0.20 in the bivariate analysis. The final modelling of factors associated with HBV infection was adjusted for age, sex, education, history of hepatitis in the family, injecting drug use, sexual intercourse under the influence of alcohol, the age at the time of sexual intercourse, STI history, and exposure to T. pallidum. Variables with

Guimarães et al. BMC Public Health (2019) 19:443 Page 6 of 11

*p*-value < 0.05 in multivariate analysis were considered to be statistically significant.

#### **Ethical aspects**

This study was approved by the Ethics Committee of the Hospital das Clínicas of the Federal University of Goiás, with protocol number 1.669.956/2016. Informed written consent was obtained from all participants. For minors, consent was also obtained from their legal guardians. Individuals with positive test results were referred to specialist services for clinical evaluation and treatment.

#### Results

Between August and December 2016, 455 households were visited. Of these, 13 residences were excluded from the study because none of their residents were found even after three visits on different days and times. In five households, all residents refused to participate in the survey. Thus, the study applied the CCEB questionnaire in 437 residences. Of these, 33 residents who did not meet the poverty criteria were excluded from the study; thus, overall data from 404 residences were included.

Thus, only one resident was randomly selected from all residents in the eligible households (N = 404) and invited to participate in the study. Of these, 26 refused to participate in the study, resulting in a study population of 378 participants.

The socio-demographic and behavioural characteristics of the participants are shown in Table 1. The median age was 31 (IQR, 21–42) years, and for schooling was 7 (IQR, 5–9) years. The majority of participants were married (59.3%).

The main known risk factors for HBV infection found in our study were the following: sharing of personal objects (70.0%), tattoo (32.0%), body piercing (11.1%), imprisonment (19.0%), alcohol use (49.5%), alcohol dependence (30.7%), marijuana use (24.3%), intranasal cocaine use (12.4%) and crack use (8.2%), STI history (10.8%), inconsistent condom use (81.2%), sexual intercourse under the influence of alcohol (41.7%), and exposure to *T. pallidum* (9.5%). The median age at first sexual intercourse and the number of lifetime sexual partners were 15 years and five partners, respectively.

Markers of exposure to HBV were detected in 40 participants, representing an overall prevalence rate of 9.8% (95% CI, 7.2–13.2). Three participants (0.8%) tested positive for HBsAg, 29 for anti-HBc and anti-HBs (7.7%), and five for isolated anti-HBc (1.3%). Isolated positivity for anti-HBs was observed in 25.4% (95% CI, 21.3–30.0) of the participants, suggesting that they were vaccinated against HBV. All of the participants with total anti-HBc positivity were tested for IgM anti-HBc, but none of them tested positive. Finally, the majority of the studied

population (64.8%) were considered to be susceptible to HBV infection (Table 2).

In the Poisson bivariate analysis, exposure to HBV was associated with higher age, history of hepatitis in the family, injecting drug use, alcohol use, exposure to T. pallidum, sexual intercourse under the influence of alcohol, and age at first sexual intercourse (p-value < 0.05). These variables and sex (p-value = 0.112) were included in the Poisson regression model with robust variance (Table 3).

Table 4 shows the multiple regression analysis of factors associated with HBV exposure. After adjusting for the model, age (adjusted prevalence ratio [APR], 1.04; 95% CI, 1.01–1.07), female sex (APR, 2.18; 95% CI, 1.01–4.73), sexual intercourse under the influence of alcohol (APR, 2.49; 95% CI, 1.09–5.67), and *T. pallidum* exposure (APR, 3.10; 95% CI, 1.36–7.06) remained as the independent risk factors associated with HBV exposure (Table 4).

#### Discussion

This study investigated the prevalence and risk factors of HBV infection in PLP in the Central-West Region of Brazil. The research findings provide important data on the epidemiology of HBV infection, which may intensify the prevention and control actions in this population. The observed prevalence was similar to that found in the general population. However, approximately 3/5 of the population showed absence of any marker of infection, indicating viral susceptibility. Age, female sex, sexual intercourse under the influence of alcohol, and *T. pallidum* exposure were the independent risk factors associated with HBV exposure.

In the present study, the prevalence rate of HBV exposure was 9.8% (95% CI, 7.2–13.2%), similar to that found in the general population aged 20–69 years in Brazil (11.6%; 95% CI, 10.7–12.4%) and the Central-West Region (12.7%; 95% CI, 10.9–14.5%) [4]. In Brazil, few population-based

**Table 2** Prevalence of serological markers of hepatitis B virus in people living in poverty. Goiânia, Goiás, Central-West Region of Brazil, 2016

Serological markers	N = 378	%	95.0% CI
HbsAg + Total anti-HBc	3	0.8	0.3-2.3
Total Anti-HBc + anti-HBs	29	7.7	5.4-10.8
Total Anti-HBc isolated	5	1.3	0.5-3.0
IgM Anti-HBc	0	0	0
Some exposure markers	37	9.8	7.2-13.2
Anti-Hbs isolated	96	25.4	21.3-30.0
Absence of marker (susceptible)	245	64.8	59.9-69.5

Abbreviations: HbsAg: Hepatitis B surface antigen; Total anti-HBc: antibody to HBV core antigen; anti-HBs: hepatitis B surface antibody; IgM Anti-HBc: IgM antibody to hepatitis B core antigen; 95.0% CI: 95.0% Confidence Interval

Guimarães et al. BMC Public Health (2019) 19:443 Page 7 of 11

**Table 3** Bivariate analysis of the potential factors associated with HBV exposure in people living in poverty. Goiânia, Goiás, Central-West Region of Brazil, 2016

Variables	Total <sup>a</sup>	HBV exposure				PR (95.0% CI)	<i>p</i> -value <sup>b</sup>
		Negative		Positive			
Age (years)	33.0 (23.0–46)	32.0 (21.	0–42.5)	47.0 (34.5–54.0) 7.0 (2.0–10.0)		1.03 (1.01–1.06)	<0.001
Education (years)	7.0 (4.0–9.0)	7.0 (4.0-	9.0)			0.97 (0.89–1.07)	0.657
Sex							
Male	144	130	90.3	14	9.7	1.00	
Female	138	115	83.3	23	16.7	1.71 (0.88–3.33)	0.112
Marital status							
Married	167	147	88.0	20	12.0	1.00	
Divorced/separated/widowed	28	23	82.1	5	17.9	1.49 (0.55–3.97)	0.424
Single	87	75	86.2	12	13.8	1.15 (0.56–2.35)	0.699
Tattoo							
No	195	171	87.7	23	12.3	1,00	
Yes	87	74	85.1	13	14.9	1.21 (0.61–2.38)	0.573
Body piercing							
No	219	187	85.4	32	14.6	1,00	
Yes	63	58	92.1	5	7.9	0.54 (0.21–1.39)	0.204
Sharing of personal objects							
No	83	73	88.0	10	12.0	1,00	
Yes	198	172	86.9	26	13.1	1.08 (0.52–2.26)	0.817
History of hepatitis in the family							
No	204	178	87.3	26	12.7	1,00	
Yes	32	23	71.9	9	28.1	2.20 (1.03-4.70)	0.041
Blood transfusion							
No	245	216	88.2	29	11.8	1,00	
Yes	30	24	80.0	6	20.0	1.68 (0.70-4.06)	0.242
History of imprisonment							
No	220	192	87.3	28	12.7	1,00	
Yes	50	42	84.0	8	16.0	1,11 (0,54-2,26)	0,772
Marijuana use							
No	212	182	85.8	30	14.2	1,00	
Yes	70	63	90.0	7	10.0	0.70 (0.31–1.60)	0.408
Crack use							
No	253	221	87.4	32	12.6	1,00	
Yes	29	24	82.8	5	17.2	1.36 (0.53–3.49)	0.519
ntranasal cocaine use							
No	244	211	86.5	33	13.5	1,00	
Yes	38	34	89.5	4	10.5	0.77 (0.27–2.19)	0.636
njecting drug use							
No	278	243	87.4	35	12.6	1,00	
Yes	4	2	50.0	2	50.0	3.97 (0.95–16.51)	0.058
Alcohol use							
No	139	128	92.1	11	7.9	1,00	
Yes	143	117	81.8	26	18.2	2.29 (1.13-4.64)	0.021

Guimarães et al. BMC Public Health (2019) 19:443 Page 8 of 11

**Table 3** Bivariate analysis of the potential factors associated with HBV exposure in people living in poverty. Goiânia, Goiás, Central-West Region of Brazil, 2016 (Continued)

Variables Total <sup>a</sup> HBV e			V exposure			PR (95.0% CI)	<i>p</i> -value <sup>b</sup>
	Negative			Positive			
Alcoholic dependence							
No	181	160	88.4	21	11.6	1,00	
Yes	84	69	82.1	15	17.9	1.53 (0.79–2.98)	0.202
STIs history							
No	249	219	88.0	30	12.0	1,00	
Yes	33	26	78.8	7	21.2	1.76 (0.77-4.00)	0.178
Treponema pallidum exposure							
Negative	252	226	89.7	26	10.3	1,00	
Positive	30	19	63.3	11	36.7	3.55 (1.75–7.19)	< 0.001
Condom use at last sexual interco	urse						
Yes	60	52	86.7	8	13.3	1,00	
No	195	166	85.1	29	14.9	1.11 (0.50–2.43)	0.785
Anal sex							
No	153	129	84.3	24	15.7	1,00	
Yes	98	85	86.7	13	13.3	0.84 (0.43-1.66)	0.626
Sexual relation homosexual							
No	233	201	86.3	32	13.7	1,00	
Yes	21	16	76.2	5	23.8	1.73 (0.67-4.44)	0.253
Sexual intercourse with illicit drug	users						
No	230	197	85.7	33	14.3	1,00	
Yes	26	22	84.6	4	15.4	0.86 (0.45-1.64)	0.653
Sexual intercourse with sex worke	rs						
No	195	166	85.1	29	14.9	1,00	
Yes	60	52	86.7	8	13.3	00.89 (0.40-1.96)	0.785
Sexual intercourse with STIs carried	rs						
No	230	196	85.2	34	14.8	1,00	
Yes	25	22	88.0	3	12.0	0.81 (0.24-2.64)	0.729
Sexual intercourse under the influ	ence of alcohol						
Never	139	126	90.6	13	9.4	1,00	
Sometimes/always	117	117	79.5	24	20.5	1.19 (1.11–4.30)	0.027
Age of first sexual intercourse	15.0 (13.5–17.0)	15.0 (14.0	0–16.0)	15.0 (13	.0–17.5)	1.04 (0.99–1.11)	0.100
Number of sexual partners	6.0 (3.0–15.0)	6.0 (3.0-	15.0)	9.0 (2.2-	-28.7)	1.00 (0.99–1.00)	0.792
Number of sexual partners	1.0 (1.0-1.0)	1.0 (1.0-	1.0)	1.0 (1.0-	-1.0)	1.02 (0.90-1.15)	0.716

Abbreviations: HBV: hepatitis B virus; PR: Prevalence ratio; STIs: Sexually transmitted infections; 95.0% CI: 95.0% Confidence interval Note: quantitative variables presented as medians and IQR

studies investigated the epidemiology of hepatitis B in PLP [34]. Considering the studies conducted in other PLP or proxy populations, the results were similar to low-income adolescents (5.9%; 95% CI, 4.2–8.0%) [19] and recyclable material pickers (12.8%; 95% CI, 9.8–16.2%) [35] of Goiânia, Central-West Region. These results indicated that the Central-West Region has a low endemicity for hepatitis B.

On the other hand, the results of this investigation were lower than those estimated in recyclable waste pickers in the City of Santos (São Paulo State, South-eastern Brazil) (34.4%; 95% CI, 28.5–40.2%) [24] and in the low-income population of Buriticupu (Maranhão State, Northeast Region) (40.7; 95% CI, 34.7–47.0%) [20]. In addition to the methodological differences and variations in risk behaviours,

<sup>&</sup>lt;sup>a</sup>Number of valid answers

<sup>&</sup>lt;sup>b</sup>Wald statistic

Guimarães et al. BMC Public Health (2019) 19:443 Page 9 of 11

**Table 4** Multiple regression analysis of risk factors associated with HBV exposure in individuals living in poverty in the metropolitan region of Goiânia, Central Brazil, 2016

Risk factors	APR (95.0% CI) <sup>a</sup>	<i>p</i> -value <sup>b</sup>
Age (years)	1.04 (1.01–1.07)	0.002
Education (years)	1.06 (0.96–1.17)	0.243
Female sex	2.18 (1.01–4.73)	0.048
History of hepatitis in the family	1.66 (0.73–3.77)	0.219
Injecting drug use	1.61 (0.29–8.78)	0.578
Sexual intercourse under the influence of alcohol	2.49 (1.09–5.67)	0.029
Age of first sexual intercourse	1.04 (0.98–1.10)	0.168
STIs history	0.64 (0.21–1.95)	0.439
Treponema pallidum exposure	3.10 (1.36–7.06)	0.007

Abbreviations: HBV: hepatitis B virus; APR: Adjusted prevalence ratio; STIs: Sexually transmitted infections; 95.0% CI: 95.0% Confidence interval

the first study included an older population than that of the present study. Moreover, in the case of the second study, the North Region had an intermediate endemicity when compared to the Central-West Region. These factors may explain the differences of prevalence found between these studies.

The vaccine card is considered to be the gold standard for the evaluation of vaccination coverage. However, when unavailable, isolated detection of the anti-HBs marker can be used as an indicator of the vaccine status of populations. In this study, only 25.4% of the participants presented a serological profile consistent with previous immunisation, and 64.8% did not present hepatitis B serologic markers, indicating viral susceptibility. Low serological evidence of HBV immunisation has also been found in other studies in PLP or in those with extreme social vulnerability in Brazil [20, 35, 36].

The Global Health Sector Strategy on Viral Hepatitis by the World Health Organization (WHO) proposed to eliminate hepatitis B by 2030 treating 80% of patients with HBV and HCV and achieving 90% vaccination coverage of the three doses in childhood by 2020. Although several member states of the WHO, including Brazil, are working on strategies to control hepatitis and eliminate the epidemic, the challenges are still immense, especially when it comes to financial issues [37]. Globally, only 11% of HBV and HCV cases are diagnosed. There is a dire need to hasten hepatitis diagnosis and search the missing millions living with hepatitis who remain undiagnosed [38]. The diagnosis may be even more underestimated in PLP, because of inaccessibility to healthcare services.

Currently, in Brazil, hepatitis B vaccine is widely available at most public health network to the populace regardless of age and/or level of vulnerability [39]. However, the low immunisation rate and the high susceptibility rate shown in

this study indicate that extremely poor populations have difficulties or fail to access the immunisation programme. Consequently, this will increase the susceptibility to more advanced age groups, which could constitute clusters of maintenance of the virus transmission chain. This is often related to the absence of primary healthcare services in impoverished regions. The presence of susceptible population groups may negatively impact the extent of elimination of hepatitis B transmission by 2030.

Multiple regression analysis, as well as the other studies, revealed an association between exposure to HBV and age increase [14, 20, 36, 40]. The displacement of the epidemiological curve of HBV in populations with higher age groups can be explained by the Brazilian policy of increasing the age for eligibility of hepatitis B vaccine over the years [34]. This result supports the importance of offering and intensifying vaccination against hepatitis B in elderly [36]. The association between age and HBV exposure may also reflect the cumulative risk of HBV infection associated with lifelong sexual and parenteral exposure [14, 41].

Alcohol consumption has a complex association with health status, and the determination of the damages caused by it is even more complicated because it is influenced by the volume and consumption pattern of the individual. In 2016, 60 to 79.9% of Brazilian men and 40 to 59.9% of Brazilian women reported alcohol consumption in the last 12 months, with a daily intake of 3 to 4 doses and 1 to 2 doses, respectively, accounting for a total of 47.3 deaths per 100,000 inhabitants [42]. Alcohol consumption during intercourse was also associated with exposure to HBV, which is a well-documented association in the national and international literature [34, 43]. Alcohol consumption during intercourse decreases the perception of risk and increases the chance of risky behaviours, such as inconsistent condom uses and multiple sexual partners. These behaviours potentiate the risk for HBV infection [43, 44].

<sup>&</sup>lt;sup>a</sup>Model adjusted for age, sex, education, history of hepatitis in the family, injecting drug use, sexual intercourse under the influence of alcohol, age at first sexual intercourse, STIs history and exposure to *Treponema pallidum*; Wald chi-squared test; McFadden's R<sup>2</sup>: 0.154

bWald statistic

Guimarães et al. BMC Public Health (2019) 19:443 Page 10 of 11

In the present study, there was an association between exposure to *T. pallidum*, the etiological agent of syphilis, and exposure to HBV. Of the total number of participants exposed to HBV, 29.7% were also exposed to *T. pallidum*. The presence of ulcerative lesions caused by syphilis significantly increases the risk of infection and transmission of HBV and other STI such as human immunodeficiency virus (HIV) [45, 46]. This result highlights the need for testing for multiple STI, such as hepatitis B and C, syphilis, and HIV, in PLP.

This study had some limitations. The cross-sectional nature of this study does not allow the establishment of causality between HBV exposure and predictor variables. The frequencies of risk behaviours may have been underestimated due to recall bias and socially desirable responses to particular issues, such as number of partners, use of condoms, and consumption of licit and illicit substances. The generalisability of our findings is limited because our study included PLP in only one Brazilian region. However, this study represented one of the first investigations on the epidemiology of HBV infection, which may contribute to the implementation of control policies in this population.

#### **Conclusion**

In conclusion, there was a high prevalence of HBV exposure in PLP in the Central-West Region of Brazil, indicating significant viral spread of the infection. High rates of risk behaviour were found, suggesting that this population is at a high risk of acquiring HBV and other STI. Moreover, there was low serological evidence of immunisation against hepatitis B, indicating that a large proportion of the participants in this study are susceptible to this infection. The results support the need for public health policies to promote the access of hard-to-reach groups by the existing healthcare services, especially the immunisation programme against hepatitis B.

#### Abbreviations

95% Cl: 95% confidence interval; AD: Anderson-Darling; Anti-HBc total: Total antibody to HBV core antigen; Anti-HBs: Hepatitis B surface antibody; APR: Adjusted prevalence ratio; HBsAg: Surface antigen of the hepatitis B virus; HBV: Hepatitis B virus; IgM anti-HBc: IgM antibody to hepatitis B core antigen; IIQ: Interquartile range; PLP: People living in poverty; PR: Prevalence ratio

#### Acknowledgements

We would like to thank all the volunteers and researchers involved in this study.

#### **Funding**

Not applicable.

#### Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

#### Authors' contributions

SB & LCCG contributed to the conception and design of this project. RAG conduct the analyses. GR substantively reviewed the work. SB, LCCG, RAG, RM & GR have full access to the data and take responsibility for the integrity and accuracy of the data. VEC, JRS, CMS & TGM conduzed to the collection of data and contributions to the interpretation of data. VLBO & EPSL

conduzed the processing of ELISA tests and have drafted the work. HG-J contributed to the construction of the English contents. All authors (LCCG, SB, RAG, HG-J, RM, VEC, JRS, EPSL, CMS, VLBO, GCA, TGM & GR) contributed to interpreting the data and revising the manuscript for intellectual content. All authors approved the final manuscript.

#### Ethics approval and consent to participate

This study was approved by the Human Research Ethics Committee of the Federal University of Goiás. Written consent was obtained from all participants. Written informed consent for the inclusion of minors was obtained from the parents or legal guardians to participate to study according to rules of the Brazilian Ethics Committee.

#### Consent for publication

Written informed consent was obtained from the people for publication of the accompanying images. A copy of the written consent is available for review by the Editor of this journal.

#### Competing interests

The authors declare that they have no competing interests.

#### **Publisher's Note**

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

#### **Author details**

<sup>1</sup>Faculty of Nursing, Federal University of Goiás, Goiânia, Goiás, Brazil. <sup>2</sup>Instituto de Patologia Tropical e Saúde Pública, Federal University of Goiás, Goiânia, Goiás, Brazil. <sup>3</sup>Instituto de Ciências Biológicas, Federal University of Goiás, Goiânia, Goiás, Brazil. <sup>4</sup>Laboratório de Análises Clínicas e Estudo em Saúde (LACES), Federal University of Goiás, Goiânia, Goiás, Brazil. <sup>5</sup>Secretaria de Estado da Saúde de Goiás/Coordenação Estadual de Controle das Hepatites Virais – CECHV, Goiânia, Goiás, Brazil. <sup>6</sup>Department of Infectious Diseases, Istituto Superiore di Sanitá, Rome, Italy.

## Received: 26 November 2018 Accepted: 15 April 2019 Published online: 29 April 2019

#### References

- Nayagam S, Thursz M, Sicuri E, Conteh L, Wiktor S, Low-Beer D, et al. Requirements for global elimination of hepatitis B: a modelling study. Lancet Infect Dis. 2016;16:1399–408.
- World Health Organization. Global Hepatitis Report, 2017. 2017. http://apps. who.int/iris/bitstream/10665/255016/1/9789241565455-enq.pdf.
- Schweitzer A, Horn J, Mikolajczyk RT, Krause G, Ott JJ. Estimations of worldwide prevalence of chronic hepatitis B virus infection: a systematic review of data published between 1965 and 2013. Lancet. 2015;386:1546–55.
- Pereira LM, Martelli CM, Hamann EM, Merchán-Hamann E, Montarroyos UR, Braga MC, et al. Population-based multicentric survey of hepatitis B infection and risk factor differences among three regions in Brazil. Am J Trop Med Hyg. 2009;81:240–7.
- Razavi-Shearer D, Gamkrelidze I, Nguyen MH, Chen DS, Van Damme P, Abbas Z, et al. Global prevalence, treatment, and prevention of hepatitis B virus infection in 2016: a modelling study. Lancet Gastroenterology & Hepatology. 2018;3(6):383–403.
- Franco E, Bagnato B, Marino MG, Meleleo C, Serino L, Zaratti L. Hepatitis B: Epidemiology and prevention in developing countries. World J Hepatol. 2012;4:74–80.
- Rachiotis L, Papagiannis D, Markas D, Thanasias E, Dounias L. Hepatitis B virus infection and waste collection: prevalence, risk factors, and infection pathway. Am J Ind Med. 2012;55:650–7.
- Lutfi K, Trepka MJ, Fennie KP, Ibanez G, Gladwin H. Racial residential segregation and risky sexual behavior among non-Hispanic blacks, National Survey of family growth. HHS Public Access Soc Sci Med. 2015;140:95–103. https://doi.org/10.1016/j.socscimed.2015.07.004.
- Walker KO, Leng M, Liang LJ, Forge N, Morales L, Jones L, et al. Increased patient delays in care after the closure of Martin Luther king hospital: implications for monitoring health system changes. Ethn Dis. 2011;21:356–60.
- Acevedo-Garcia D, Osypuk TL, McArdle N, Williams DR. Toward a policyrelevant analysis of geographic and racial/ethnic disparities in child health. Health Aff. 2008;27:321–33.

- Poundstone KE, Strathdee SA, Celentano DD. The social epidemiology of human immunodeficiency virus/acquired immunodeficiency syndrome. Epidemiol Rev. 2004;26:22–35.
- Oka T, Enoki H, Tokimoto Y, Kawanishi T, Minami M, Okuizumi T, et al. Employment-related difficulties and distressed living condition in patients with hepatitis B virus: a qualitative and quantitative study. BMC Public Health. 2017;17:1–12.
- Kim TH, Samson LF, Lu N. Racial/ethnic disparities in the utilization of hightechnology hospitals. J Natl Med Assoc. 2010;102:803–10.
- Ferreira PM, Guimarães RA, Souza CM, Guimarães LCDC, Barros CVDL, Caetano KAA, et al. Exposure to hepatitis C virus in homeless men in Central Brazil: a cross-sectional study. BMC Public Health. 2017;17:90.
- Adimora AA, Schoebach VJ. Contextual factors and the black-white disparity in heterosexual HIV transmission. Epidemiology. 2002;13:707–12.
- Fullman N, Yearwood J, Abay SM, Abbafati C, Abd-Allah F, Abdela J, et al. Measuring performance on the healthcare access and quality index for 195 countries and territories and selected subnational locations: a systematic analysis from the global burden of disease study 2016. Lancet. 2018; 391(10136):2236–71.
- 17. Estatística IB de G e. Uma análise dos resultados do universo do Censo Demográfico 2010. 2011.
- Tanaka J. Hepatitis B epidemiology in Latin America. Vaccine. 2000;18(Suppl 1):517–9.
- Oliveira MDDS, Martins RMB, Matos MA, Ferreira RC, Dias MA, Carneiro MAS, et al. Seroepidemiology of hepatitis B virus infection and high rate of response to hepatitis B virus Butang vaccine in adolescents from low income families in Central Brazil. Mem Inst Oswaldo Cruz. 2006;101:251–6 http://www.scielo.br/scielo.php?script=sci\_arttext&pid=S0074-02762006000300004&Ing=en&nrm=iso&tlng=en.
- El Khouri M, Cordeiro Q, Da Luz DABP, Duarte LS, Gama MEA, Corbett CEP. Endemic hepatitis B and C virus infection in a Brazilian eastern Amazon region. Arq Gastroenterol. 2010;47:35–41.
- 21. Kageyama A, Hoffmann R. Pobreza no Brasil: uma perspectiva multidimensional. Econ e Soc. 2006;15:79–112.
- Estratifica RDE, Socioecon DEC, Brasil MNO. Critérios de estratificação e comparação de classificadores socioeconômicos no brasil. 2016;:55–70.
- Associação Brasileira de Empresas de Pesquisa. Critério Brasil 2015 e atualização da distribuição de classes para 2016. 2016. http://www.abep.org/ Servicos/Download.aspx?id=12.
- Marinho TA, Lopes CLR, Teles SA, de Matos MA, de Matos MAD, Kozlowski AG, et al. Epidemiology of hepatitis B virus infection among recyclable waste collectors in Central Brazil. Rev Soc Bras Med Trop. 2014;47:18–23.
- Szwarcwald CL, Malta DC, Pereira CA, Vieira MLFP, Conde WL, De Souza Junior PRB, et al. Pesquisa Nacional de Saude no Brasil: concepcao e metodologia de aplicacao. Cien Saude Colet. 2014;19:333–42.
- Damacena GN, Szwarcwald CL, Malta DC, de Souza Júnior PRB, Vieira MLFP, Pereira CA, et al. O processo de desenvolvimento da Pesquisa Nacional de Saúde no Brasil, 2013. Epidemiol e Serviços Saúde. 2015;24:197–206. https:// doi.org/10.5123/S1679-49742015000200002.
- 27. Mayfield D, McLeod G, Hall P. The CAGE questionnaire: validation of a new alcoholism screening instrument. Am J Psychiatry. 1974;131:1121–3.
- Derajew H, Tolessa D, Feyissa GT, Addisu F, Soboka M. Prevalence of depression and its associated factors among primary caregivers of patients with severe mental illness in southwest, Ethiopia. BMC Psychiatry. 2017;17:1–8.
- Masur J, Monteiro MG. Validation of the "CAGE" alcoholism screening test in a Brazilian psychiatric inpatient hospital setting. Braz J Med Biol Res. 1983;16:215–8.
- 30. StataCorp. Stata Statistical Software: Release 14. College Station, TX: StataCorp LP; 2015.
- Razali NM, Wah YB. Power comparisons of Shapiro-Wilk, Kolmogorov-Smirnov, Lilliefors and Anderson-Darling tests. J Stat Model Anal. 2011;2:21–33.
- Barros AJD, Hirakata VN. Alternatives for logistic regression in cross-sectional studies: an empirical comparison of models that directly estimate the prevalence ratio. BMC Med Res Methodol. 2003;3:21.
- 33. Coutinho LMS, Scazufca M, Menezes PR. Methods for estimating prevalence ratios in cross-sectional studies. Rev Saude Publica. 2008;42:1–6.
- 34. Ximenes RAA, Figueiredo GM, Cardoso MRA, Stein AT, Moreira RC, Coral G, et al. Population-based multicentric survey of hepatitis B infection and risk factors in the north, south, and southeast regions of Brazil, 10-20 years after the beginning of vaccination. Am J Trop Med Hyg. 2015;93:1341–8.
- Rozman MA, Alves IS, Porto MA, Gomes PO, RibeirO NM, Nogueira LAA. HIV infection and related risk behaviors in a community of recyclable waste collectors of Santos. Brazil Rev Saude Publica. 2008;42(5):838–43 Epub 2008 Jul 31.

- PMR C, Dos S, de Matos MA, Pinheiro RS, KAA C. Prevalence, risk factors and hepatitis B immunization: helping fill the gap on hepatitis B epidemiology among homeless people, Goiânia, Central Brazil. Cad Saúde Pública. 2017;33:7.
- World Health Organization. Draft global health sector strategies Viral hepatitis, 2016–2021. 2016. http://apps.who.int/gb/ebwha/pdf\_files/WHA69/ A69\_32-en.pdf?ua=1
- 38. Yasir W, Masood S, Zubia J, Muzammil HN. Hepatitis elimination by 2030: progress and challenges. World J of Gastroenterology. 2018;24(44):4959.
- Ministério da Sáude. Secretaria de Vigilância em Saúde., Departamento de Vigilância de Doenças e Agravos não Transmissíveis e Promoção da Saúde. Nota informativa n. 149, de 2015/CGPNI/DEVIT/MS. 2015. http://www.cevs.rs. gov.br/upload/arquivos/201705/18120135-nota-informativa-n-149-2015capni-devit-svs-ms.pdf.
- 40. Fontes AM, Ardisson JS, Souza MAA. Epidemiology of hepatitis B virus in the cities of the northern region of Espírito Santo. Brazil. 2016;88:1549–55.
- 41. Lopes CLR, Teles SA, Espírito-Santo MP, Lampe E, Rodrigues FP, Motta-Castro ARC, et al. Prevalence, risk factors and genotypes of hepatitis C virus infection among drug users, Central-Western Brazil. Rev Saude Publica. 2009; 43 Suppl. 1.
- Griswold MG, Fullman N, Hawley C, Arian N, Zimsen SRM, Tymeson HD, et al. Alcohol use and burden for 195 countries and territories, 1990-2016: a systematic analysis for the global burden of disease study 2016. Lancet. 2018;392(10152):1015–35.
- 43. Gitto S, Vitale G, Villa E, Andreone P. Update on alcohol and viral hepatitis. J Clin Transl Hepatol. 2014;2:228–33.
- Vagenas P, Lama JR, Ludford KT, Gonzales P, Sanchez J, Altice FL. A systematic review of alcohol use and sexual risk-taking in Latin America. Rev Panam Salud Publica. 2013;34:267–74.
- Salado-Rasmussen K. Syphilis and HIV co-infection. Epidemiology, treatment and molecular typing of Treponema pallidum. Dan Med J. 2015;62:B5176.
- 46. Lynn WA, Lightman S. Syphilis and HIV: a dangerous combination. Lancet Infect Dis. 2004;4:456–66.

#### Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

#### At BMC, research is always in progress.

Learn more biomedcentral.com/submissions

