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Prevalence of Anti-HBcore Total and HBsAg among Health Care Workers in Public Hospitals, White Nile State, Sudan; 2013

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Authors' contributions

This work was carried out in collaboration between all authors. Authors MAEAE and TAE designed the study. Authors AAA, SAB and ZS revised the tools for data collection. Authors MAEAE and AAA maintained the quality of data collection in the field work. Author TAE drafted the first draft of the manuscript. Authors SAB and AAB revised the manuscript for grammar and spelling. All authors read and approved the final manuscript.

Article Information

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ABSTRACT

Background: HBV infection is an occupational disease where health care workers (HCW) are at high risk.

Aim: To measure the sero-prevalence of Anti-HBcore Total and HBsAg among HCWs in Public Hospitals, White Nile State, Sudan; 2013.

Methods: A cross-sectional, hospital- based study was conducted among health care workers in

Public Hospitals in White Nile State, Sudan; 2013. A sample of 385 HCWs was selected using two stage cluster sampling. A pre-tested structured questionnaire was used. The HCWs signed the informed consent to fill the questionnaire and to draw 5 ml venous blood sample for HBV tests. Blood samples were investigated for Anti-HBcore Total. Positive blood specimens for Anti-HBcore Total were further investigated for HBsAg. Data was processed using statistical package for social sciences (SPSS), version 16. Descriptive statistics and non-parametric Z test for single proportion was used at 95% CL.

Results: Out of 385 HCWs, 230 (60%) were positive for Anti-HBcore Total. Out of 230 HCWs, 62 (27%) were positive for HBsAg. Prevalence for Anti-HBcore Total and HBsAg is significantly different from the expected values, *P*=0.001

Conclusion: Sero-prevalence of Anti-HBcore Total and HBsAg is high among HCWs in Public Hospitals in White Nile State, Sudan.

Keywords: Anti-HBcore total; HBsAg; HCWs; public hospitals; White Nile State; Sudan.

1. INTRODUCTION

HBV infection is defined as the presence of Anti-HBcore in the serum of an individual whether he/she is HBsAg negative or positive. So, he/she may or may not be shedding virus to others. Carrier state: It is the presence of HBsAg in the serum of an individual, whether he/she has symptoms and signs of HBV infection or not. Thus, he/she is shedding virus to others [1].

Hepatitis B virus (HBV) is a major cause of cirrhosis of the liver and hepatocellular carcinoma (HCC). About half of hepatocellular carcinoma cases and one third of liver cirrhosis are due to chronic HBV infection. Yearly, about 500000 – 700000 deaths were estimated to be due to HBV infection. Across the world, two billion individuals were infected with HBV; among whom 360 million were chronically infected [2,3].

There is a variation in the prevalence of HBV infection worldwide; regarding different areas and population in the same area. The world is divided into: (i) Hyper-endemic area with a prevalence of 70% - 90% of Anti-HBcore and 8% of HBsAg; where 45% of the population lives (South-Eastern Asia and sub-Saharan Africa). (ii) Moderate endemic area with a prevalence of 2% – 7% of HBsAg (Southern countries of Central and Eastern Europe, Mediterranean basin, the Amazon's sink, Middle East, and Northern Africa) (iii) Low endemic area with a prevalence less than 2% of HBsAg (North-Western Europe and North America) [4,5].

A study was carried in Tamil Nadu, Southern State of India, it showed HBV carrier rate of 5.7% (CI 4.6- 6.8) among 1981 respondents [6].

Sudan belongs to Sub-Saharan countries with high HBV sero-prevalence. Among the general population; infection rate (positive Anti-HBcore) varied from 47% to 78%, while carrier rate (positive HBsAg) prevalence ranged from 6.8% in Central Sudan to 26% in Southern Sudan [7,8].

The spectrum of clinical manifestations of HBV infection varies in both acute and chronic status of the disease. During the acute phase, manifestations range from subclinical or anicteric hepatitis to icteric hepatitis and, in some cases, fulminant hepatitis. During the chronic phase, manifestations range from an asymptomatic carrier state to chronic hepatitis, cirrhosis, and hepatocellular carcinoma. Extra hepatic manifestations can occur with both acute and chronic infection [9].

HCWs are more prone to acquire blood-borne diseases as occupational hazard and the degree of their exposure determines the rate of HBV infection [10,4].

A sero-epidemiologic survey of HBV markers among health care workers (HCWs) in Public Teaching Hospitals in Khartoum State, Sudan; showed that HBVs infection and carrier rates were 57% ($\text{Cl}_{95\%}$: 53%–60%) and 6.0% ($\text{Cl}_{95\%}$: 4.0%–8.0%) respectively, P < 0.05 [11].

Aim of the study: To measure the prevalence of Anti-HBcore (infection rate) and HBsAg (carrier rate) among health care workers (HCWs) in Public Hospitals in White Nile State, Sudan; 2013.

2. METHODS

2.1 Study Design

This is a cross-sectional, hospital-based study.

2.2 Study Area

White Nile State lies south to Khartoum City and it is traversed by White Nile River and composed of eight localities with seventeen public hospitals.

2.3 Study Population

HCWs that working in the Public Hospitals in White Nile State for more than 45 days. The total number was 1808 health care workers.

2.4 Sample Size and Selection Procedure

The overall sample size was determined by the formula:

$$n = \frac{Z^2 PQ}{d^2}$$

n = the desired sample size.

z = confidence coefficient = 1.96

p = prevalence rate. p = 50% or 0.5

q = 1-p = 1-0.5=0.5, d = the degree of accuracy, was set at 0, 05

Accordingly

$$n = \frac{1.96x1.96 \times 0.5 \times 0.5}{0.05 \times 0.05} = 384.6 = 385$$

A cluster sampling was used. The hospitals were divided into groups according to the number of specialties in them. It was selected proportionally; every hospital was given a proportion of the sample HCWs according to the total number of health workers. The target sample size was 385; it was distributed as follow:

Group A: Hospitals with all specialties; with 1182 heath care workers (HCWs) (sample size = 252).

Group B: Hospitals with one specialty; with 157 heath care workers (HCWs) (sample size = 33)

Group C: hospital with no specialty; with 469 heath care workers (HCWs) (sample size = 100)

2.5 Data Collection, Analysis and Processing

Data was collected using pre-tested structured questionnaire. Structured questionnaire is a quantitative method of research, which includes the low level of involvement of the researcher and high number of respondents (the individuals who answer the questions). Pre-testing used to

be carried out in a situation similar to that of the study, in order to identify difficulties and problems that related to the questions and also to train the data collectors. The questionnaire was composed of socio-demographic variables. Five ml venous blood was drawn after the signature of the informed consent and before filling the questionnaire. Blood sera was separated and stored at −20℃, until testing. Using ELISA tests with 99.64% sensitivity and 99.64% specificity; all specimens were tested for anti-HBcore total were tested for HBsAg.

Data was processed using statistical package for social sciences (SPSS), version 16. Descriptive statistics and non parametric Z-test for single proportion was used. The P-value ≤ 0.05 was considered statistically significant for the results.

2.6 Ethical Issue

The study was approved by the ethical committee of Sudan Medical Specialization Board.

3. RESULTS AND DISCUSSION

3.1 Health Care Workers (HCWs) Distribution and Prevalence of antiHBcore total and HBsAg among Them is Presented in Fig. 1

Fig. 1 indicates that the most representative localities in the study were Kostee, with 94 HCWs (24.4%) followed by Algeteena and Rabak, with 80 HCWs (20.8%) and the least localities were Alsalam and Kenanna with 11 HCWs (2.9%) for each. From the same Figure we noticed that the highest percentage of HBV infection (24.4%) is among HCWs in Kostee locality; the least one (2.9%) is in Alsalam and Kenana localities. For the carrier rate, Kostee and Aldweam localities have the highest percentage; while Tandalti and Alsalam localities have the least.

3.2 Socio-demographic Characteristics of the Respondents

The sample composed of 154 males (40%) and 231 females (60%). Most of them were in the age group 27-36 years (30.9%), followed by 47-56 (20.0%) and the least one was the age group of 57+ (13.2%).

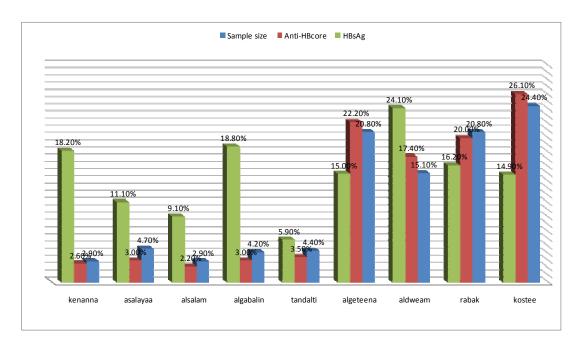


Fig. 1. Health Care Workers (HCWs) distribution for the survey of anti-HBcore total and HBsAg, by localities (n = 385); and the prevalence of anti-HBcore total (n = 385) and HBsAg (n = 62) among them in Public Hospitals, White Nile State, Sudan, 2013

For marital status, 215 HCWs (55.8%) were married, 150 HCWs (39%) were single, and 11 HCWs (2.9%) were widowed, while 9 HCWs (2.3%) were divorced.

The level of education was as follows: 149 HCWs, (38.8%) were university, 95 HCWs (24.7%) were high secondary and 6 HCWs (1.6%) were Quranic school.

Regarding occupation of the sample population; 121 (31.4%) were labour, 107 (27.8%) were nurses, 60 (15.6%) were doctors, 49 (12.7%) were technicians in labs and blood banks, 15 (3.9%) were nurse midwives, 12 (3.1%) were pharmacists, 11 (2.9%) were theatre attendants and 10 (2.6%) were Village midwives.

3.3 Testing the Prevalence of Infection Rate of HBV (Anti-HBcore total) and that of Carrier of HBV (HBsAg) against Values of test Probability of 0.5 among Health Care Workers (HCWs) is Presented in Table 1

As Table 1 shows, 230 (60%) of the tested HCWs showed positive Anti-Bcore total marker, while 155 (40%) were negative for Anti-HBcore

total marker. The *P*-value of the Z- test was 0.001, which indicates a significant statistical difference between the prevalence of 60% and the tested rate of 50% (i.e. 0.5). So, there is a high infection rate of HBV (measured by Anti-HBcore total) among HCWs in Public Hospitals, White Nile State, Sudan. The outcome of the test was that the prevalence of past or current infection with HBV among HCWs in Public Hospitals, White Nile State, Sudan, was 60%. The lower and the upper bound of Anti-Bcore total prevalence at 95% confidence level was 56% and 62% respectively, *P*-value = 0.001.

Regarding carrier rate; Table 1 shows, 62 (27%) of the tested HCWs showed positive HBsAg, while 168 (73%) were negative for HBsAg marker. The *P*-value of Z- test was 0.001, which indicates a significant statistical difference between the prevalence of 27% and the tested rate of 50% (i.e. 0.5). So, there is a high carrier rate of HBV (measured by HBsAg) among HCWs in Public Hospitals, White Nile State, Sudan. The outcome of the test was that the carrier rate (measured by HBsAg) among the respondents was 27%. The lower and upper bounds of the prevalence of HBsAg was 26% and 31%, respectively; *P*-value = 0.001.

Table 1. Testing the prevalence of infection rate of HBV (Anti-HBcore total) (n = 385) and that of carrier of HBV (HBsAg) (n = 230) against values of test probability of 0.5 among health care workers (HCWs) in Public Hospitals, White Nile State, Sudan, 2013

Markers*	Category	N	Observed prob.	Test prob.	<i>P</i> -value	Conclusion
Anti-HBcore	+ ve	230	0.60	0.5	0.001	Significant difference
Total	- ve	155	0.40			-
	Total	385	1.00			
	+ ve	62	0.27	0.5	0.001	Significant difference
HBsAg	- ve	168	0.73			_
	Total	230	1.00			

^{*}Test used was Z test for single proportion

3.4 The Relation between Sero-positivity of (Anti-HBcore Total and HBsAg) and the Various Demographic Factors among Health Care Workers (HCWs) are Presented in Table 2

Looking at Table 2 there is no statistical association between the various demographic factors and the prevalence of Anti-HBcore total and HBsAg among HCWs in Public Hospitals in White Nile State; Sudan.

4. DISCUSSION

The study was an observational hospital base study. Three hundred and eighty five HCWs in Public Hospitals, White Nile State, Sudan, were enrolled. As shown by Z-test for single proportion there was a statistical difference between the expected (50%) and actual (60%) prevalence, p-value = 0.001, indicating that the difference was statistically significant. Sudan is one of the high endemic countries with HBV [12-15]. There is a high rate of HBV infection among HCWs in

Kostee and Aldweam localities in comparison to other localities; while carrier rate measured by HBsAg is high among HCWs in Kostee locality. These are the heavily populated localities in White Nile State and they have the more established hospitals in this State with regard to other localities. So, the high prevalence of both Anti-HBcore and HBsAg among HCWs may be due to their exposure to blood and body fluids of patients in this high endemic area. The result was consistent with many national studies as reported in Public Teaching Hospitals in Khartoum State, Sudan, where the infection and carrier rates are high [1,16,17]; the Gezira State of Central Sudan, a community base study indicates a high carrier rate among the general population [18]; and international studies as that of Hepatitis B and E viral infections among Nigerian healthcare workers [7,1,15,19-21]; and Southern State of India [6] There is no statistical association between the different demographic variables and the studied markers. This may be due to high endemicity of the disease in this area.

Table 2. Relation between sero-positivity of (Anti-HBcore total and HBsAg) and the various demographic factors among health care workers (HCWs) in Public Hospitals in White Nile State, Sudan; 2014; (n= 385)

Demographic factors	Test	P- value	Conclusion
Localities	Anti-HBcore total	.228	
	HBsAg	.569	
Gender	Anti-HBcore total	.832	
	HBsAg	.390	
Education level	Anti-HBcore total	.279	
	HBsAg	.193	Insignificant
Marital status	Anti-HBcore total	.092	
	HBsAg	.174	
Occupation	Anti-HBcore total	.373	
	HBsAg	.463	

5. CONCLUSION AND RECOMMENDA-TION

The outcome of this study concluded that the infection and carrier rates of HBV were high among HCWs in Public Hospitals, White Nile State, Sudan. Further study to address the possible risk factors is highly recommended.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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ANEX 1

Questionnaire

Serological and molecular characterizations of HBV, HDV and HCV among Health Care Workers in Public Hospitals, in White Nile State, Sudan; the year 2013

0 QUESTIONNAIRE IDENTIFICATION DATA		
001 QUESTIONNAIRE IDENTIFICATION NUMB	ER _	_
002 CITY		
003 HOSPITAL		
004 DEPARMENT		
005 TELEPHONE NO		
Introduction: "My name is	Medicine, Universor department BV, HDV and HO a been interviewe ESPONDENT HA EW THIS PERSO	sity of Khartoum. We're interviewing] in order to find out the CV among Health Care Workers in ed in the past few weeks [or other AS BEEN INTERVIEWED BEFORE ON AGAIN. WE tell them we cannot
Confidentiality and consent: "I'm going to ask written on this form. You do not have to answer you may end this interview at any time you we questions will help us better understand the situated Civil Teaching Hospitals. We would greatly approximately will take about XX minutes to ask the questions."	any questions the want to. However ation of HBV amoreciate your help	nat you do not want to answer, and er, your honest answers to these ng HCWs in Khartoum State Public in responding to this survey. The
(Signature of interviewer certifying that informed	consent has been	n given verbally by respondent)
Interviewer visit		
	Date	
	Interviewer	
	Result	
Result codes:		
Completed 1; Respondent not available 2; Refus	sed 3; partially cor	mpleted 4.
005 INTERVIEWERS: Code [Name	e	
006 DATE OF INTERVIEW:\ \		
007 CHECKED BY SUPERVISOR: Signature _		Date
The HCWs questionnaire includes the following	ng sections:	

Section 0 – Questionnaire identification data ((7) codes)

Section 1 – Background characteristics

(9) Questions

Section 2 - Knowledge, believes, and attitudes.

(28) Questions

Section 3 - Knowledge about vaccination against HBV and HDV (6) Questions

Section 4 - Past Medical History

(5) Questions

Section 5- Occupational Exposure.

(5) Questions

Section 6: infection control measures

(10) Questions

Total number of questions:

(63) Questions

Section 1: Background characteristics

No.	Questions and filters	Coding categories
Q1	Gender	1. Male 2. Female
Q2	Age/ years	
Q3	Original Residency	 South North East West Central
Q4	Marital Status	1. Married 2. Not married 99. No Response
Q5	Educational Level	 Illiterate Khalwa Primary Secondary Higher Graduated Postgraduate No Response
Q6	Occupation	 Doctor Nurse Midwife Lab +blood bank technicians Labor Theatre attendants Manager No Response
Q7	If you are a doctor, what is your classification?	1. Consultant2. Registrar3. Medical officer4. House officer
Q8	What is your department?	1. Surgery2. Obs. & Gyn.3. Dentistry.4. Others
Q9	Your duration of work in this hospital?	 Less than 1year 1 - 2 years More than 2 - 5 years More than 5 - 10 years More than 10 - 20 years More than 20 year
		88. Don't know 99. no response

Section 2: Knowledge, believe and attitude

No.	Questions and filter	Co	ding categories		
Q10	Have you ever hear	d about:			
	1. HBV	1. Yes	2. No	88. Don't know	99. No respons
	2. HCV	1. Yes	2. No	88. Don't know	99. No response
	3. HDV	1. Yes	2. No	88. Don't know	99. No response
Q11	Is HBV transmitted	by blood t	ransfusior	or blood produc	ts
	1. Yes 2. No 88. Don't		know 99.		
	Is HCV transmitted	by blood t	ransfusior	or blood produc	ts
	1. Yes 2. No	88. Don't	know 99.	No response	

	Is HDV to	ransmitted	by blood transfe	usion or blood products			
	1. Yes	2. No	88. Don't know	99. No response			
Q12	Can a pr	egnant wo	man infected wit	h HBV transmit the virus to her unborn child?			
	1. Yes	2. No	88. Don't know	99. No response			
	Can a pr	egnant wo	man infected wit	h HCV transmit the virus to her unborn child?			
	1. Yes	2. No	88. Don't know	99. No response			
	Can a pr	egnant wo	man infected wit	h HDV transmit the virus to her unborn child?			
	1. Yes	2. No	88. Don't know	99. No response			
Q13	Can infe	cted woma	an with HBV trans	smit the virus to her newborn during labor?			
	1. Yes	2. No	88. Don't know	99. No response			
	Can infe	cted woma	an with HCV trans	smit the virus to her newborn during labor?			
	1. Yes	2. No	88. Don't know	99. No response			
	Can infected woman with HDV transmit the virus to her newborn during labor?						
	1. Yes	2. No	88. Don't know	99. No response			
Q14	Can infected woman with HBV transmit the virus to her newborn by breastfeeding?						
	1. Yes	2. No	88. Don't know	99. No response			
	Can infected woman with HCV transmit the virus to her newborn by breastfeeding?						
	1. Yes	2. No	88. Don't know	99. No response			
	Can infected woman with HDV transmit the virus to her newborn by breastfeeding?						
	1. Yes	2. No	88. Don't know	99. No response			
Q15	Can HB\	/ transmitt	ed sexually?				
	1. Yes	2. No	88. Don't know	99. No response			
	Can HC\	/ transmitt	ed sexually?				
	1. Yes	2. No	88. Don't know	99. No response			
	Can HD\	/ transmitt	ed sexually?				
	1. Yes	2. No	88. Don't know	99. No response			

Q16	Could sh	narp conta	minated instrume	ent transmit HBV infection?			
	1. Yes	2. No	88. Don't know	99. No response			
	Could sh	narp conta	minated instrume	ent transmit HCV infection?			
	1. Yes	2. No	88. Don't know	99. No response			
	Could sh	narp conta	minated instrume	ent transmit HDV infection?			
	1. Yes	2. No	88. Don't know	99. No response			
Q17	Is HBV ti	ransmitted	by eating or drin	king contaminated food and water?			
	1. Yes	2. No	88. Don't know	99. No response			
	Is HCV to	ransmitted	by eating or drin	king contaminated food and water?			
	1. Yes	2. No	88. Don't know	99. No response			
	Is HDV to	ransmitted	by eating or drin	king contaminated food and water?			
	1. Yes	2. No	88. Don't know	99. No response			
Q18	Is HBV transmitted by cough?						
	1. Yes	2. No	88. Don't know	99. No response			
	Is HCV to	ransmitted	by cough?				
	1. Yes	2. No	88. Don't know	99. No response			
	Is HDV transmitted by cough?						
	1. Yes	2. No	88. Don't know	99. No response			
Q19	Is HBV ti	ransmitted	by sharing razor	s?			
	1. Yes	2. No	88. Don't know	99. No response			
	Is HCV transmitted by sharing razors?						
	1. Yes	2. No		99. No response			
	IS HDV ti	ransmitted	by sharing razor				
	1. Yes	2. No	88. Don't know	99. No response			
Q20	Do you t	hink that h	ealthy-looking H	CWs could be a carrier of HBV?			
	1. Yes	2. No	88. Don't know	99. No response			
	Do you t	hink that h	ealthy-looking H	CWs could be a carrier of HCV?			
	1. Yes	2. No	88. Don't know	99. No response			

	Do you t	hink that h	ealthy-looking H	CWs could I	be a carrier of HDV?			
	1. Yes	2. No	88. Don't know					
Q21	Do you t	hink that h	ealthy-looking pe	erson could	be a carrier of HBV?			
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Do you t	hink that h	ealthy-looking pe	erson could	be a carrier of HCV?			
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Do you t	hink that h	ealthy-looking pe	erson could	be a carrier of HDV?			
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
Q22	Can peo	ple protect	themselves from	າ:				
	HBV?	1. Yes			99. No response			
	HCV?				99. No response			
	HDV?	1. Yes	2. No 88.	Don't know	99. No response			
Q23	Is protec	tion from I	HBV can be by us	sing gloves	?			
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Is protection from HCV can be by using gloves?							
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Is protection from HDV can be by using gloves?							
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
Q24	Is using sterile instrument and syringes protects from HBV transmission?							
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Is using sterile instrument and syringes protects from HCV transmission?							
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Is using sterile instrument and syringes protects from HDV transmission?							
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
Q25	Is screening of donated blood protects from HBV transmission?							
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Is screen	ning of don	ated blood prote	cts from HC	CV transmission?			
	1. Yes	2. No	88. Don't know	99. No resp	ponse			
	Is screen	ning of don	ated blood prote	cts from HD	OV transmission?			
	1. Yes	2. No	88. Don't know	99. No resp	ponse			

Q26	Is screening of organ donor's blood protects from HBV transmission?						
	1. Yes 2. No 88. Don't know 99. No response						
	Is screening of organ donor's blood protects from HCV transmission?						
	1. Yes 2. No 88. Don't know 99. No response						
	Is screening of organ donor's blood protects from HDV transmission?						
	1. Yes 2. No 88. Don't know 99. No response						
Q27	Can people protect themselves by using HBV vaccine?						
	1. Yes 2. No 88. Don't know 99. No response						
	Can people protect themselves by using HCV vaccine?						
	1. Yes 2. No 88. Don't know 99. No response						
	Can people protect themselves by using HDV vaccine?						
	1. Yes 2. No 88. Don't know 99. No response						
Q28	Can people protect themselves by drinking clean water?						
	1. Yes 2. No 88. Don't know 99. No response						
Q29	Can they protect themselves by cauterization?						
	1. Yes 2. No 88. Don't know 99. No response						
Q30	How could pregnant woman reduce the risk of transmission of HBV to her unborn child?						
	Take Immunoglobulin						
	2. Take Vaccination 88. Don't know						
	99. no response						
Q31	How could pregnant woman reduce the risk of transmission of HCV to her unborn child?						
	1. Take Immunoglobulin						
	2. Take Vaccination 88. Don't know						
	99. no response						
Q32	How could pregnant woman reduce the risk of transmission of HDV to her unborn child?						
	1. Take Immunoglobulin						
	2. Take Vaccination 88. Don't know						
	99. no response						

Q33	Did you voluntarily do:							
	HBV test? 1. Yes 2. No 88. Don't know 99. No response							
	if no go to question 35							
	HCV test? 1. Yes 2. No 88. Don't know 99. No response							
	if no go to question 35							
	HDV test? 1. Yes 2. No 88. Don't know 99. No response							
	if no go to question 35							
Q34	When did you have your most recent,							
	HBV test?							
Q35	Do you required having the test once in the past and you did not have it?							
	1. Yes 2. No 88. Don't know 99. No response							
Q36	Can people protect themselves by using personal protection? (e.g. Gloves, Goggles and Gowns)							
	1. Yes 2. No 88. Don't know 99. No response							
Q37	Are unsafe injection practices one of sources of transmission of HBV?							
	1. Yes 2. No 88. Don't know 99. No response							
	Are unsafe injection practices one of sources of transmission of HCV?							
	1. Yes 2. No 88. Don't know 99. No response							
	Are unsafe injection practices one of sources of transmission of HDV?							
	1. Yes 2. No 88. Don't know 99. No response							

Section 3: Knowledge about vaccination against HBV, HCV, HDV

No.	Question	ns and filte	ers	Coding categories
Q38	Is HBV i	nfection pr	evented by a vac	ccine?
	1. Yes	2. No	88. Don't know	99. No response
	Is HCV i	nfection pr	evented by a vac	ccine?
	1. Yes	2. No	88. Don't know	99. No response
	Is HDV i	nfection pr	evented by a vac	ccine?
	1. Yes	2. No	88. Don't know	99. No response

A. Number of vaccine doses of HBV 1. One dose										
B. Number of vaccine doses of HCV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response C. Number of vaccine doses of HDV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _ if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _ if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _ if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _ if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
B. Number of vaccine doses of HCV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response C. Number of vaccine doses of HDV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response C. Number of vaccine doses of HDV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response C. Number of vaccine doses of HDV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
88. Don't know 99. No response C. Number of vaccine doses of HDV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
C. Number of vaccine doses of HDV 1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
1. One dose 2. Two doses 3. Three doses 4. Others() 88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
88. Don't know 99. No response Q40 Have you ever been vaccinated against HBV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
All										
1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HCV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_ _] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
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Have you ever been vaccinated against HDV? 1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country	1. Yes 2. No 88. Don't know 99. No response									
1. Yes 2. No 88. Don't know 99. No response If yes, when? Date [_] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
If yes, when? Date [_] if no go to Q 42 Q41 Source of vaccine 1. Inside the country 2. Outside the country										
Q41 Source of vaccine 1. Inside the country 2. Outside the country										
1. Inside the country 2. Outside the country										
1. Inside the country 2. Outside the country Q42 What do you know about the vaccine										
Q42 What do you know about the vaccine										
□HBV □ HCV □HDV										
1. I do not know about the vaccine										
2. I do not know where to find it										
3. Expensive4. Not interested to be immunized.										
5. Others										

Section 4: Past medical history of exposure to risk factors:

No.	Questions and File	ters	Coding categories			
Q43	Have you ever been subjected to blood transfusion					
	1. Yes	2. Never	88. Don't know	99. No response		
	If yes, when?					
	1. Less than 45 days 2. More than 45 days					

Q44	Have you ever been subjected to surgical procedure								
	1. Yes 2. Never 88. Don't know 99. No response								
	If yes, when?								
	1. Less than 45 days 2. More than 45 days								
Q45	Have you ever been subjected to tattooing								
	1. Yes 2. Never 88. Don't know 99. No response								
	If yes, when?								
	1. Less than 45 days 2. More than 45 days								
Q46	Have you ever been subjected to cautery 1. Yes 2. Never 88. Don't know 99. No response If yes, when?								
	1. Less than 45 days 2. More than 45 days								
Q47	Have you ever been having jaundice?								
	1. Yes 2. Never 88. Don't know 99. No response If yes, at what age?								
	1. Below 30 years 2. Between 30-49 years 3. 50 years or more								

Section 5: Exposure to occupational risk factors

No.	Questions and filters Coding categories			
Q48	Have you ever been experienced needle stick injury?			
	1. Yes 2. Never 88. Don't know 99. No response. If yes, when?			
	1. Less than 45 days 2. More than 45 days			
Q49	Have you ever been exposed to injury with contaminated sharp instruments?.			
	1. Yes 2. Never 88. Don't know 99. No response. If yes, when?			
	1. Less than 45 days 2. More than 45 days			
Q50	Have you ever been dealing with blood?.			
	1. Yes 2. Never 88. Don't know 99. No response. If yes, when?			
	1. Less than 45 days 2. More than 45 days			
Q51	Have you ever been dealing with amniotic fluid during delivery event?.			
	1. Yes 2. Never 88. Don't know 99. No response. If yes, when?			
	1. Less than 45 days 2. More than 45 days			
Q52	Have you ever been dealing with jaundiced patient?			
	1. Yes 2. Never 88. Don't know 99. No response. If yes, when?			
	1. Less than 45 days 2. More than 45 days			

Section 6: Infection control measures

Q53	Had you received formal training on infection control precautions						
	1. Yes	2. No	88. Don't know	99. No response			
Q54	Do you wash or decontaminate your hands before and after direct contact with patients?						
	1. Yes	2. No	88. Don't know	99. No response			
Q55	Do you wash or decontaminate your hands after contact with blood, body fluid, secretions or execration?						
	1. Yes	2. No	88. Don't know	99. No response			
Q56	Do you w	ash or deco	ontaminate your ha	inds before handling an invasive devise (± gloves)?			
	1. Yes	2. No	88. Don't know	99. No response			
Q57	Do you w	ash your ha	ands before gloving	g and before leaving the examination room?.			
	1. Yes	2. No	88. Don't know	99. No response			
Q58	Do you w	ear gloves	when Touching b	lood and body fluids?			
	1. Yes	2. No	88. Don't know	99. No response			
Q59 Do you change gloves between tasks and procedures on the		and procedures on the same patients?					
	1. Yes	2. No	88. Don't know	99. No response			
Q60	Do you remove gloves immediately after use before attending to another patients						
	1. Yes	2. No	88. Don't know	99. No response			
Q61	Do you wear surgical masks during procedures that are likely to generate splashes or sprays of blood or other body fluids?						
	1. Yes	2. No	88. Don't know	99. No response			
Q62	Do you wear Gowns during procedures that are likely to generate splashes or sprays of blood or other body fluids?						
	1. Yes	2. No	88. Don't know	99. No response			
Q63	Do you wear goggles during procedures that are likely to generate splashes or spray blood or other body fluids?						
	1. Yes	2. No	88. Don't know	99. No response			

That is the end of our questionnaire. Thank you very much for taking time to answer these questions. We appreciate your help.

Peer-review history:
The peer review history for this paper can be accessed here:
http://sciencedomain.org/review-history/13842

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