

**INTERNATIONAL JOURNAL OF ADVANCES IN
PHARMACY, BIOLOGY AND CHEMISTRY**

Research Article

**Sero-prevalence of hepatitis C virus among Patients
attending dental clinics in Khartoum- Sudan**

Khadeejah Mohammed Mohammed Othman*, Wafa Ibrahim Elhag*.

Microbiology Department, Faculty of Medical Laboratories Sciences,
Al- Neelain University, Sudan.

ABSTRACT

This study was carried out to detect frequency of Hepatitis C virus (HCV) among patients attended dental clinics of Khartoum-Sudan, and to determine the relationship between the presence of HCV and certain factors such as gender, age, social status, history of haemodialysis and blood transfusion. It was a descriptive cross-sectional study conducted between January-March 2015. A total of 90 apparently healthy individuals who visited dental clinics, (43 males and 47 females) were enrolled. Serum specimens were tested by ELISA, sero prevalence of HCV was 0.00%.

Keywords : frequency, hepatitis C, ELISA, dental clinics, Khartoum -Sudan.

INTRODUCTION

Hepatitis continues to have an impact on the practice of dentistry, Possible transmission in the dental setting, management of the chronically ill and the legal issues related to the treatment of infectious patients who may act as source of spreading of the infection among dental health care workers and other patients in dental clinics, such patients are unaware of their status because of long incubation period and post infection window period during which antibodies cannot be detected^{1,2}.

Hepatitis C is recognized as an important global infectious disease affecting primarily the liver, caused by hepatitis C virus (HCV), with more than 170 million people are chronically infected³. Hepatitis C is generally asymptomatic with up to 80% of infected cases which will progress to persistent infection. About 15-20% of chronic HCV infection progressed to cirrhosis and 1 to 4% are found to be an annual risk of developing hepatocellular carcinoma within 20-30 years^{3,4}. Blood, blood-products and intravenous drug abuse (IV) are related with the transmission of this virus, Probabilities due to sexual and interfamilial transmission are somehow low⁵. Surprisingly, Nokhodian and his colleagues (2012) stated that HCV transmission may also be acquired through non-

parenteral and non- sexual routes⁶. The presence of viral hepatitis C particles in oral fluids has been demonstrated by many authors and this may indicate that transmission via saliva and gingival cervical fluid might occur^{7,8}. Up to 40% of patients infected with HCV may have non identifiable routes of viral acquisition. Dental extraction may be one of these risk factors⁹. The non-existence of a good patient's history in as well as in private dental clinics and hospitals, unsterilized dental and medical equipment, used syringes and unsterilized instruments are common causes of the spread of HCV¹⁰.

Hepatitis C is very important for dentists because of its transmission route. The dentists are particularly at risk because of exposure to the oral secretions and blood of potentially infectious patients¹¹. There is no published data available regarding the frequency of HCV among the dental patients in Sudan. The aim of this study to detect frequency of Hepatitis C virus (HCV) among patient who had attended dental clinics of Khartoum-Sudan, and to determine the relationship between the presence of HCV and certain factors such as gender, age, social status, past history of haemodialysis and blood transfusion.

MATERIALS AND METHODS

This was descriptive- cross sectional study which had been conducted in Khartoum state during period from January to March 2015, 90 patients who attended dental clinics for extraction and other surgical procedures such as diagnostic biopsy prior to oral and maxillofacial surgical procedures were enrolled, Data was collected by using direct interviewing questionnaire; ethical clearance was obtained from research ethical committee of faculty of graduate studies Al-Neelain University and Ministry of Health Khartoum state, written consent also was obtained from each patient.

Experimental work

Samples collection:

blood samples were collected from patients, under direct medical supervision by medial vein puncture using 5 ml syringe into plain tube to obtain serum by centrifugation at

5000 rpm for 10 min. serum was kept in -20°C till serological study was performed.

Specimens were processed by Enzyme linked immune sorbent assay (ELISA) (4th generation ELISA) (fortress diagnostics, UK) for detection HCV antibodies.

Enzyme linked immune sorbent assay for detection HCV antibodies:

All reagents and samples were allowed to reach room temperature for 15 minutes before use washing buffer was prepared 1:20 from buffer concentrate with distilled water. 100µl of sample diluents was added into appropriate wells except the blank well and negative well. 20µl from each sample was added to the appropriate wells and mixed by pipette repeatedly until liquids turn blue. 20µl from negative and positive control was dispense and added to the negative and positive wells separately without dispensing

liquid into the blank control well. Microtiter wells was flicked for 30 seconds and mixed well, then plate was covered and incubated for 30 minutes at 37° C .plate was taken out and wash buffer was added to each well (washing 1) and aspirated off after 20 seconds. This step was repeated for 5 times until each well become dry, and 50µl of HRP-Conjugate Reagent was added in to each well except the blank, the plate was mixed well and covered with the plate cover and incubated for 30 min at 37°C.

The plate cover was removed and discarded. The liquid was aspirated and each well was rinsed in wash buffer . This step was repeated for 5 times until each well become dry (washing 2).

50µl of substrate A and 50µl substrate B solution were added in to each well including the Blank and mixed by tapping the plate gently. The plate was incubated at 37°C for 15 min. 50 µl Stop solution was added into each well and mixed gently.

Measuring the absorbance: The plate reader was calibrated with blank well and the absorbance was read at 450nm. The results were calculated by relating each sample optical density (OD) value to the Cut off value of plate. Calculation of cut off (C.O) value.

$$C.O = *Nc * 2.1$$

*Nc= the mean absorbance value for the three negative controls.

The absorbance was read with micro well reader at 450nm.

Interpretation of Results:

Negative results: samples giving absorbance less than Cut-off value are negative for this assay. Positive result: sample giving absorbance equal to or greater than Cut-off considered initially reactive. Borderline: sample with absorbance to Cut-off value are considered borderline and retesting of these samples in duplicate is recommended.

Data analysis: Data was analyzed by SPSS (Statistical Package of Social Science) software program version 16.

RESULT

A total of 90 patients who attended dental clinics during the period from January-March 2015, consented to the study were included, study subjects were 43(47.8%) males and 47(52.2%) females .The average age of patients was 42.27 years (range from 12 to 80 years) ,most of patients 33 (36.7%) were belonged to the age group (31-50) fig(1). The overall result showed that no one had HCV antibodies (0.00%). Study population were divided into 2 groups 59 (65.6%) were married and 31(34.4%) were single, most of study population where from Khartoum locality (28(31.1%)), table (1). fig(2) summarized the demographic data of study population, regarding their occupation, most of them were housewife 26(28.9%). Regarding clinical data,16 (17.8%) had a history of hepatitis, 87.5% were did not know type of hepatitis they got , most of study population had no history of surgical operation (57 (63.3%)) , 69(76.7%) had no history of blood transfusion, 69(76.7%) had no history of tattooing. table(2) . all study population had neither history of haemodialysis nor organ transplantation.

DISCUSSION

More than 170 million people are infected with HCV, causing over 350,000 deaths annually³. Some health care interventions may act as risk factors for HCV infection and dental procedures may be one of these¹². Dental procedure is one of the major source of exposure for HCV transmission (39.7%)¹³. The present study result revealed 0.00% seropositivity, when compared with other studies in Sudan, it is slightly similar to the result obtained by Isam-*et al* (2001) a 0.4% of HCV antibodies was detected among blood donors in Khartoum state¹⁴ and 0.6% among pregnant women in Khartoum –state in study conducted by Elsheikh- *et al* (2007)¹⁵. However, the obtained seropositivity was lower than the result of 4.5% obtained by Mudawi- *et al* (2007) among patients with hepatosplenic schistosomiasis in Khartoum state¹⁶ and 1.3% among pregnant women in central Sudan in study conducted by Osman- *et al*(2014)¹⁷. Other reports among the general population from Western and Southern Sudan showed a prevalence of 2-3%^{18,19}. the few studies on HCV infections in Sudan demonstrated a low seroprevalence ranging from 2.2% in Gezira²⁰ To 4.8% in patients with schistosomal periportal fibroses¹⁶.the present findings (subjects had no history of haemodialysis) was extremely lower than a study conducted by El-Amin – *et al* (2007)among haemodialysis patients in Khartoum –state in which a prevalence of 23.7% was reported²¹ and study conducted by Nalam- *et al*(2014) in which a 39.8% of HCV antibodies was detected among patients undergoing renal dialysis, this higher incidence was significantly associated with higher duration of

dialysis²², also the present finding was lower than seroprevalences of HCV noted in other African countries such as Ethiopia (2%)²³, central African republic (5%)²⁴and Libya (7.9%)²⁵.Generally the serofrequency of HCV in Sudan is low compared with neighboring countries ,Egypt the northern neighboring country to Sudan reported the highest HCV seroprevalence in the world , 12 to 31 %^{26,27,28,29}, UK have high rate of HCV which is 214,000 individuals have long-term (chronic) infection with hepatitis C in study conducted by Annastella- *et al*(2014)³⁰,also 4.1% and 2.1% of HCV antibodies was reported among transfused patients and untransfused blood donors respectively in study conducted by Balogun-*et al* (2014)³¹.

CONCLUSION

This study revealed none HCV seropositivity among dental patients, the discrepancies of this result may be due to small sample size and differences in the used techniques, for this large scale screening is recommended.

ACKNOWLEDGEMENT:

We would like to thanks all participants of this study, laboratory staff at Khartoum teaching dental hospital specially Miss. Fatima in blood bank and all doctors and staff at dental clinic of dental collage Al-Neelain University, Sudan. We also grateful to Mr. Badawi and staff at national public Health laboratory-Khartoum for technical assistance and all Microbiologydepartment Al-Neelain University.

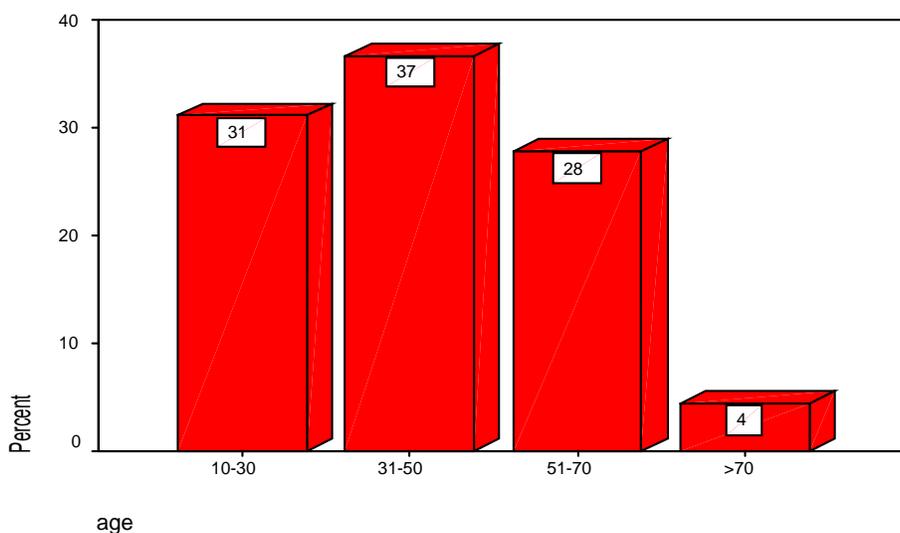


Fig 1
Distribution of study population(n=90) according to their age.

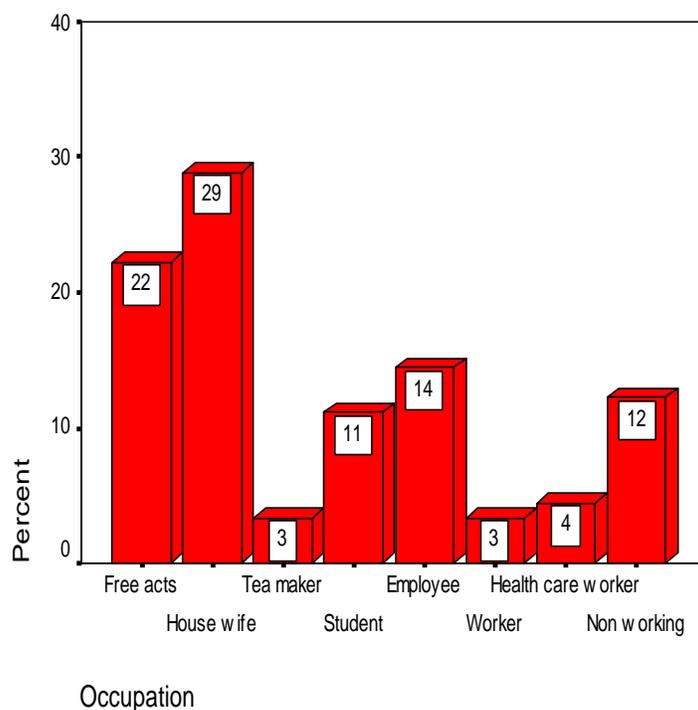


Fig 2
Distribution of study population (n=90) according to their occupation.

Table 1
Distribution of study population (n=90) according to their residence.

		Frequency	Percent
Valid	Khartoum	28	31.1
	Omdurman	16	17.8
	Bahri	25	27.8
	Out of Khartoum	21	23.3
	Total	90	100.0

Table 2
Clinical data of study population (n=90).

Clinical data	yes	No	Total
History of surgical operation	33(36.7%)	57(63.3%)	90
Family history of hepatitis	14(15.6%)	76(84.4%)	90
History of blood transfusion	21(23.3%)	69(76.7%)	90
History of tattooing	21(23.3%)	69(76.7%)	90

REFERENCES

1. Andres PL, Fabiano JA, THines TJ. Spec Care Dentists, 2000 Sep-Oct; 20(5):209-13.
2. Samarana Yake L, Rules of Infection Control, Int Dent J, 1993; 43:578-84.
3. Baha W, Foullous A, Dersi N, They-they TP, El alaoui K, Nourichafi N, Oukkache B, Lazar F, Benjelloun S, Ennaji MM, Elmalki A, Mifdal H and Bennani A. Prevalence and risk factors of hepatitis B and C virus infections among the general population and blood donors in Morocco. BMC Public Health, 2013; 13:50.
4. Alavian SM, Mahboobi N and Karayiannis P. Oral conditions associated with hepatitis C virus infection, Saudi J Gastroenterol, 2013; 19:245-51.
5. Yildirim B, Tahan V, Ozaras R, Aytakin H, Mert A, Tabak F and Senturk H. Hepatitis C virus risk factors in the Turkish community, Dig Dis Sci, 2005; 50:2352-5.
6. Nokhodian Z, Yazdani MR, Yaran M, Shoaie P, Mirian M, Ataie B, Babak A and Ataie M. Prevalence and risk factors of HIV, Syphilis, Hepatitis B and C among Female Prisoners in Isfahan, Iran. Hepat Mon, 2012; 12:442-7.
7. Mahboobi N, Porter SR, Karayiannis P and Alavian SM. Oral fluid and hepatitis A, B and C, 2012; 41:505-16.
8. Stewardson DA, Palenik CJ, McHugh ES and Burke FJ. Occupational exposures occurring in students in a UK dental school, 2002; 6:104-13.
9. Modi AA and Liang TJ. Hepatitis C: a clinical review, 2008; 14:10-4.
10. Rashid F, Dent M, But M. Pakistan Oral and Dental Journal, 2006; 26(1):51-54.
11. Sonis ST, Fazio RC, Fang L. editors, Principles and Practice of Oral Medicine, 1995; 131-45.
12. Mahboobi N, Porter SR, Karayiannis P and Alavian SM. Dental treatment as a risk factor for hepatitis B and C viral infection, 2013; 22:7986.
13. Ali I, Siddique L, Rehman LU, Khan NU, Iqbal A, Munir I, Rashid F, Khan SU, Attache S, Swati ZA and Aslam MS. Prevalence of HCV among the high risk groups in Khyber Pakhtunkhwa. Virol J, 2011; 8:296.
14. Isam KM, Hasabelgawi OA. Prevalence of hepatitis C virus antibodies in Khartoum State, A thesis submitted in the fulfillment of the requirements of master degree in immunology, University of Khartoum, Khartoum, Sudan, 2001.
15. Elsheikh RM, Daak AA, Elsheikh MA, Krsany MS and Adam I. Hepatitis B virus and Hepatitis C virus in pregnant Sudanese women, 2007; Virol J. 4:104.
16. Mudawi HM, Smith HM, Fletcher IA and Fedail SS. Prevalence and common genotypes of HCV infection in Sudanese patients with hepatosplenic schistosomiasis, 2007, J Med Virol; 79:1322-4.
17. Osman AMM, Mirghani OA, Gasim GI, Adam I. Hepatitis B virus, Hepatitis C virus and Human Immunodeficiency Virus infections among Pregnant Women in Central Sudan. Sudan journal of medical sciences, June 2014; Vol 9(2):91-96.
18. Omer RE, Verhoef L, Vant Veer P, et al, Peanut butter intake, GSTM-1 genotype and hepatocellular carcinoma: a case control study in Sudan. Cancer causes control, 2001; 312:23-32.
19. McCarthy MC, El-Tigani A, Khalid IO and Hyams KC. Hepatitis B and C in Juba Southern Sudan: Result of a serosurvey. Trans. R. Soc. Trop. Med. Hyg 1994; 88:534-536.
20. Mudawi HMY, Smith HM, Rahoud SA. Epidemiology of HCV infection in the Gezira state of central Sudan, 2007; 79:383-5.
21. El-Amin HH, Osman EM, Mekki MO, Abdelraheem MB, Ismail MO, Yousif ME, Abass AM, El-haj HS, Ammar HK, Hepatitis C virus infection in haemodialysis patients in Sudan :two centers report Saudi J. Kidney Dis, 2007; 18:101-106.
22. Nalam RG, Nalam SG, Thota HR, Ravichandra K, Rohit A and Madhavi A. Effect of end-stage renal disease on oral health in patients undergoing renal dialysis: A cross sectional study. Journal of International Society of Preventive and Community Dentistry, 2014; 4(3):164-169.
23. Formmel D, Tekle- Haimanot R, Berhe N et al. A survey of antibodies to hepatitis C virus in Ethiopia, 1993; 49:435-9.
24. Pawlotsky JM, Belec L, Gresenguet G et al, High prevalence of hepatitis B, C and E markers in young sexually active adults from the Central African Republic, 1995; 46:269-72.
25. Saleh MG, Pereira LM, Tibbs CJ et al. High prevalence of hepatitis C virus in the normal Libyan population, 1994; 88:292-4.
26. Frank C, Mohamed MK, Strickland GT, Lavanchy D, Arthur RR, Magder LS, El Khoby T, Abdel-Wahab Y, Alyohn ES, Anwar W. The role of parenteral antischistosomal therapy in the spread of hepatitis C virus in Egypt. Lancet, 2000; 355:887-891.
27. Leehman EM, Wilson ML. Epidemic hepatitis C virus infection in Egypt estimates of past

- incidence and future morbidity and mortality ,2009; 16:650-658.
28. Mohamed MK, Hussein MH, MassoudAA, Rakhaa MM, Shoeir S, Aoun AA, AboulNaser M. Study of the risk factors for viral hepatitis C infection among Egyptians applying for work abroad. J, 1996;71(1-2):113-47.
 29. Youssef A , Yano Y, Utsumi T, Abd El-alah EM, abd El-Hameed A el E, SerwahAel H, Hayashi Y. Molecular Epidemiological study of hepatitis viruses in Ismailia, Egypt. Intervirology, 2009; 52:123-131.
 30. Annastella C, David G, Helen H, *et al.* Hepatitis C in the UK..Public Health England, 2014:133-155.
 31. Balgun TM, Ehayieme TE and Aleshinloye CT. The Seroprevalence of hepatitis C virus antibodies among transfused patients with haematological disorders. US National Library of Medicine National Institutes of Health, 2014; 21(1):17-20.