

Studies on the Prevalence of Hepatitis C Virus Infection in Diabetic Patients attending a Tertiary Health-care Facility South-west Nigeria.

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Abstract

Background: Hepatitis C virus (HCV) infection and type 2 diabetes mellitus (T2DM) are two major public health problems associated with increasing complications and mortality rates worldwide. The objective of this study to evaluate the prevalence of hepatitis C virus (HCV) infection in diabetic patients and to investigate the influence of several epidemiological and clinical factors on HCV infection.

Method: A total number of one hundred and eighty diabetic patients were recruited for this study. Consented subjects made up of 71(39.4%) males and 109(60.56%) females were recruited for the study. While one-Hundred (100) Non-Diabetics (Controls) were recruited for this study. Structured questionnaires were administered to the consented participants to obtain relevant data. Sera samples were assayed for antibodies to HCV using an enzyme linked immunosorbent assay [Inteco Diagnostic Limited]. ELISA technique.

Result: Overall prevalence of HCV infection among diabetes patients assayed was 13.3%. Out of which 8(11.3%) was obtained from the male subjects compared to 16 (14.7%) seropositivity recorded for the female subjects ($P = 0.511$; $P > 0.05$). Considering age distribution, Subjects aged 41-50 years recorded, 9 (22.5%) positivity ($P = 0.238$; $P > 0.05$). Considering educational status of subjects screened, 22 (14.9%) positivity was rescored among subjects who have attained tertiary status of education. ($P = 0.574$; $P > 0.05$). Risk factors considered showed that, 7 (18.9%) seropositive subject were alcoholic consumers (P value = 0.2621; $P > 0.05$) while 5 (8.9%) recorded history of sharing sharp objects $P = 0.2427$; $P > 0.05$).

Conclusion: Our study shows a slightly higher prevalence of hepatitis C infection in type 2 diabetics. This call for urgent routine screening exercise among diabetic patients for HCV infection. This study also emphasizes the need for public enlightenment on the association between HCV infection and T2DM, to avert possible complications among diabetic patients.

1. Background

Hepatitis is the inflammation of the liver as a result of viral infection which can either be acute or chronic, symptomatic or asymptomatic (1). This viral infection can lead to liver damage due to various causes such as heavy intake of alcohol, toxin ingestion and certain suppressive medical conditions(2). Hepatitis C virus (HCV) infection is a public health concern affecting more than 170 million people worldwide (3) (4). HCV is a positive, single-stranded RNA virus in the Flaviviridae family. HCV is most efficiently transmitted through transfusion of infected blood, transplantation of infected organs, mother-to-child transmission during childbirth and interaction with infected blood or body fluids of an infected person(5). Most cases of Liver transplant mainly occurs as a result of HCV infection progressing to chronic infections.(6).

Hepatitis C virus (HCV) has been identified as one of the leading causes of chronic liver disease with serious sequel as the end stage of cirrhosis and liver cancer(7). Hepatitis C has clearly been demonstrated

to be a precipitating factor for diabetes but only in patients with risk factors to develop such. People with hepatitis C virus (HCV) infection appear to be at increased risk of developing type 2 diabetes(8).Diabetes mellitus is a chronic disease of metabolism causing abnormal glucose homeostasis(9).

Diabetes Mellitus, commonly referred to as diabetes is an impairment in which the body is unable to process glucose leading to reduction in the blood sugar level. Diabetes is characterized by high blood glucose level(hyperglycemia) with problems of fat, protein, and carbohydrate metabolism due to the body's inability to secrete/act on insulin over a long period(10). Diabetic status of individuals are divided into two; namely Type 1 diabetes which occurs mostly in children and adolescents and Type 2 diabetes which occurs mostly in adults,(11).

Knobler, *et al.* (2000) reports an increase in DM type-2 before the development of advanced liver cirrhosis(12). Previous Research found that, after excluding chronic hepatitis C patients who received previous interferon treatment, higher fibrotic stages in liver histology and family history of DM were closely associated with higher prevalence of DM and impaired fasting glucose in patients with chronic hepatitis C(13).HCV infection and type 2 diabetes mellitus are two chronic conditions which contribute to a significant morbidity and mortality.A higher prevalence of type 2 diabetes mellitus has been observed in the developed world (2% to 9.4%) in patients with HCV infection than in those with other forms of chronic hepatitis(14,15)

Type 2 diabetes is a debilitating disease condition, while the co-infection of type 2 diabetes and HCV has been established to worsen this condition, hence it has become very necessary for a screening exercise to determine the prevalence rate of HCV among diabetic patients at our location of study, so as to increase awareness of the populace and health practitioners on the dangers of the co-infectious status of this virus with diabetics.

2. Method

Study area and population: The study was conducted at the Federal Teaching Hospital Ido-Ekiti which is a tertiary health institution. The study population comprised of randomly selected confirmed Diabetic patients attending the outpatient department of the Teaching Hospital.

Ethical permit and consent: A proposal of the project was submitted to the Ethical Review Committee of the Federal Teaching Hospital Ido-Ekiti. Where ethical permit was sought for and obtained with protocol number: ERC/2018/02/27/103B.

Inclusion and Exclusion criteria

Individuals confirmed for type 2 diabetes mellitus diabetes were recruited for the study. Persons who showed no interest in the study and are not diabetic were excluded from the study.

Questionnaire and sample size: The sample size used in this research work was obtained from One-Hundred and eighty (180) volunteers, while those recruited were informed about the study and their

consents obtained. Well-structured questionnaires were used to collect demographic data and other pertinent information.

Sample collection and processing: Three-Five (3-5) ml of blood were collected from both diabetic (Cases) and control subjects aseptically and according to standard procedure. Blood samples were allowed to clot at room temperature undisturbed, thereafter Sera obtained were dispensed into a clean, dry cryovial and stored at -20°C prior use. The sera were screened for antibodies to HCV using ELISA kits (Fortress Diagnostic Limited). Standard procedures were strictly adhered during the assay process.

Data analysis: Filled questionnaires were crosschecked manually for correct data entry. The data were analyzed using the SPSS software package, Chi-square test was used to compare several Variables while the critical level for statistical significance was set at P = 5% (0.05).

Sample processing

Sera samples obtained were screened for HCV antibodies using ELISA technique according to the manufactures manual. Serum ALT was also assayed for, following the procedure detailed out in the kit used. Hepatitis C ELISA and ALT kits were stored in the refrigerator at 4°C prior to use. Serum samples were analyzed at Landmark University Medical Laboratory.

3. Results

The total sera samples screened comprises of 71 (39.4%) Males and 109 (60.56%) Females (Table 2). Positive samples obtained showed that, 8(4.4%) were obtained from the Male diabetic patients while the female subjects recorded 16(8.9%) positivity for HCV, (Table 1). The age distribution of the subjects analyzed for the test ranged from 0-20, 21-30, 31-40, 41-50, 51-60 and 60-100 years. Of the 180 serum samples analyzed for HCV, 24 (13.3%) samples tested positive while 156 (86.7%) samples tested negative.

TABLE 1a: Distribution of sera samples assayed among Diabetic subjects screened.

Total number of samples	Number of positive samples	Number of negative samples
180	24 (13.3%)	156 (86.7%)

TABLE 1b Distribution of HCV among Non-Diabetics (Control Subjects) based on gender.

SEX	Number tested	HCV	
		Positive %	Negative%
Male	48	4(4%)	44 (44%)
Female	52	5 (5%)	47 (47%)
Total	100	9 (9%)	91 (91%)

P=0.739; P>0.05

TABLE 2: Distribution of Hepatitis C Virus among diabetic subjects based on Gender

Gender	Total number of samples examined (%)	Number of Positive samples (%)	Number of Negative samples (%)
Male	71(39.4)	8(4.4)	63(35.0)
Female	109(60.6)	16(8.9)	93(51.7)
Total	180(100.0)	24(13.3)	156(86.7)

Chi square (χ^2) = 0.433; df = 1; P value = 0.511

Figure 1: Distribution of Hepatitis C Virus according to Gender among diabetic subjects screened.

TABLE 3: Distribution of Hepatitis C Virus based on Age of subjects screened.

Age	Total number of samples examined (%)	Number of Positive samples (%)	Number of Negative samples (%)
0-20	20 (11.1)	0 (0.0)	20 (11.1)
21-30	20 (11.1)	2 (1.1)	18 (10.0)
31-40	37 (20.6)	6 (3.3)	31 (17.2)
41-50	40 (22.2)	9 (5.0)	31 (17.2)
51-60	24 (13.3)	3 (1.7)	21 (11.7)
60-100	39 (21.7)	4 (2.2)	35 (19.4)
Total	180 (100.0)	24 (13.3)	156 (86.6)

Chi square (χ^2) = 6.778; df = 5; P value = 0.238

Table 3 showed the age group of individuals tested, between 0-20 years, 20 (11.1%) individuals were tested yielding 0 (0.0%) which indicates 20 (100.0%) negative to HCV. For Subjects aged 21-30 years, 20 (11.1%) were screened out of which 2 (10.0%) showed positivity for HCV infection with 18 (90.0%) negative to HCV. Subjects aged 31-40 years, recorded 6(16.2%) positivity. Subjects aged 41- 50 recorded 9 (22.5%) positivity correspondingly, subjects aged 51-60 years, recorded 3 (12.5%) positivity to HCV infection. Interestingly subjects aged 61-100 years, recorded 4 (10.3%) positivity.

Figure 2: Distribution of Hepatitis C Virus according to Age.

TABLE 4: Distribution of Hepatitis C Virus According To Marital Status

Marital Status	Total number of samples examined (%)	Number of Positive samples (%)	Number of Negative samples (%)
Single	41 (22.8)	4 (2.2)	37 (20.6)
Married	130 (72.2)	20 (11.1)	110 (61.1)
Divorced	9 (5.0)	0 (0.0)	9 (5.0)
Total	180 (100.0)	24 (13.3)	156 (86.7)

Chi square (χ^2) = 2.312; df = 2; P value = 0.315

Table 4 showed distribution of subjects based on marital status.130 (72.2%) married subjects were screened. 20 (15.4%) recorded positivity compared to 4 (9.8%) recorded among the single subjects.

Figure 3: Distribution of Hepatitis C Virus according to marital status

TABLE 5: Distribution of Hepatitis C Virus According To Educational Background

Education	Total number of samples examined (%)	Number of Positive samples (%)	Number of Negative samples (%)
Primary	6 (3.3)	0 (0.0)	6 (3.3)
Secondary	25 (13.9)	2 (1.1)	23 (12.8)
Tertiary	148 (82.2)	22 (12.2)	126 (70.0)
No Education	1 (0.6)	0 (0.0)	1 (0.6)
Total	180 (100.0)	24 (13.3)	156 (86.7)

Chi square (χ^2) = 1.993; df = 3; P value = 0.574

Table 5, showed the educational background of the subjects screened. Subjects with secondary education status recorded 2(8.0%) positivity compared to subjects with tertiary level of education recording 22 (14.9%) positivity

Figure 4: Distribution of Hepatitis C Virus according to Educational background.

TABLE 6: Distribution of Hepatitis C Virus based on Demographic Factor.

Occupation	Total number of samples examined (%)	Number Positive samples (%)	Number of Negative samples (%)
Trading	64 (35.6)	7 (3.9)	57 (31.7)
Civil Servant	64 (35.6)	14 (7.8)	50 (27.8)
Industry	18 (10.0)	2 (1.1)	16 (8.9)
Student	34 (18.9)	1 (0.6)	33 (18.3)
Total	180 (100.0)	24 (13.3)	156 (86.7)

Chi square (χ^2) = 7.613; df = 3; P value = 0.055

Table 6, showed the demographic factor of the individuals was among which the traders recorded 7 (10.9%) positivity while Civil servants screened recorded 14 (21.9%) positivity

Figure 5: Distribution of Hepatitis C Virus according to Age.

TABLE 7: Distribution of Hepatitis C Virus based on Clinical Risk Factors.

Risk Factor		Number of Positive Samples	Number of Negative Samples	Total Number of Samples Examined	Chi-Square
Blood Transfusion	Positive	1 (0.6%)	14 (7.8%)	15 (8.3%)	0.629^a
	Negative	23 (12.8%)	142 (78.9%)	165 (91.7%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	P value = 0.4276
Blood Donation	Positive	3 (1.6%)	10 (5.6%)	13 (7.2%)	1.151^a
	Negative	21 (11.7%)	146 (81.1%)	167 (92.7%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	P value = 0.2833
Care for a Hepatitis C patient	Positive	6 (3.3%)	32 (17.8%)	38 (21.1%)	0.251^a
	Negative	18 (10.0%)	124 (68.9%)	142 (78.9%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	P value = 0.6160

Table 7, Distribution of subjects screened based on clinical risk factors of subjects with history of blood transfusion recorded 1 (6.7%) subjects with history of previously blood donation recorded 3 (23.1%) positivity for HCV infection. Subjects with history of care for a hepatitis patient resulted to 6 (15.8%) positivity

Figure 6: Distribution of Hepatitis C Virus based on Clinical Risk Factors

TABLE 8: Distribution based on lifestyle-risk factors of Subjects Screened.

Risk Factor	Response	Number of Positive Samples	Number of Negative Samples	Total Number of Samples Examined	Chi-Square
Alcohol Consumption	Positive	7 (3.9%)	30 (16.7%)	37 (20.6%)	1.257 ^a
	Negative	17 (9.4%)	126 (70.0%)	143 (79.4%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	Pvalue = 0.2621
Tribal Marks and Tattoos	Positive	4 (2.2%)	15 (8.3%)	19 (10.6%)	1.095 ^a
	Negative	20 (11.1%)	141 (78.4%)	161 (89.4%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	Pvalue = 0.2953
Sharing of Unsterilized equipment	Positive	5 (2.8%)	51 (28.3%)	56 (31.1%)	1.365 ^a
	Negative	19 (10.6%)	105 (58.3%)	124 (68.9%)	df = 1
	Total	24 (13.4%)	156 (86.6%)	180 (100.0%)	Pvalue = 0.2427
Multiple sexual partners	Positive	3 (1.7%)	4 (2.2%)	7 (3.9%)	5.494 ^a
	Negative	21 (11.7%)	152 (84.4%)	173 (96.1%)	df = 1
	Total	24 (13.4%)	156 (86.6%)	180 (100.0%)	Pvalue = 0.0191

Table 8 showed distribution of subjects based on social lifestyles, Subjects with history of alcohol consumption recorded 7 (18.9%) positivity while subjects with tribal marks or tattoos recorded 4 (21.1%) positivity subjects with history of sharing unsterilized equipment yielded 5 (8.9%) positivity to HCV infection. Interestingly, subjects with multiple sexual partners recorded 3 (42.9%) positivity to HCV infection.

TABLE 9: Distribution of Hepatitis C Virus According To Family History

Risk Factor		Number of Positive Samples	Number of Negative Samples	Total Number of Samples Examined	Chi-Square
Previous Record of Hepatitis Virus	Positive	1 (0.6%)	4 (2.2%)	5 (2.8%)	0.198^a
	Negative	23 (12.8%)	152 (84.4%)	175 (97.2%)	df = 1
	Total	24 (13.4%)	156 (86.6%)	180 (100.0%)	Pvalue = 0.6565
Diabetes by Family Members	Positive	12 (6.7%)	88 (48.9%)	100 (55.6%)	0.346^a
	Negative	12 (6.7%)	68 (37.7%)	80 (44.4%)	df = 1
	Total	24 (13.4%)	156 (86.6%)	180 (100.0%)	Pvalue = 0.5563
Hepatitis by Family Members	Positive	3 (1.7%)	29 (16.1%)	32 (17.8%)	0.528^a
	Negative	21 (11.6%)	127 (70.6%)	148 (82.2%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	Pvalue = 0.4676

Table 9, showed the distribution of subjects based clinical history; subjects with previous history of hepatitis virus yielded 1(0.6%) positivity while those with family history of diabetes tested 12 (12.0%) positivity. Individuals with family members infected with hepatitis yielded 3 (9.4%) positivity.

Table 10: Determination of Serum Alanine Aminotransferase on HCV patients.

Age	No. Seropositive For HCV (%)	Normal ALT range (%)	Abnormal ALT range (%)
0-20	0(0%)	Not applicable	Not applicable
21-30	2(1.1%)	2(1.1%)	0(0%)
31-40	6(3.3%)	3(1.7%)	3(1.7%)
41-50	9(5.0%)	6(3.3%)	3(1.7%)
51-60	3(1.7%)	1(0.6%)	2(1.1%)
61-100	4 (2.2%)	2(1.1%)	2(1.1%)
Total	24(13.3%)	14(7.8%)	10(5.5%)

4. Discussion

The present study found a slightly higher prevalence rate of 13.3% among Type 2 Diabetes Mellitus (T2DM) subjects when compared to the global prevalence rate of around (3%) in the general population. The results obtained in this study compares with the work of Nwokediuko *et al.*, (16) in Enugu where HCV occurrence rate among diabetic patients was found to be 14.0%. Although, Ejele *et al.*, (17) and Balogun *et al.*, (18), obtained a prevalence of 3.0% in Niger Delta region and 0.0% in Ibadan which is lower than the result obtained in this study. Similar study conducted among diabetics by Ndako *et al.*, (22) showed 11.0% prevalence in Jos which is also lower than the result obtained in this study, while a prevalence of 5.0% was recorded in a study carried out among diabetes patients at UIITH, Ndako *et al.*,(19) while 5.7% prevalence was reported from India Demitrost *et al.*,(20) which is lower compared to the findings obtained in this study. However, Gray *et al.*, (21) was the first to show a higher prevalence of HCV infection in T2DM patients with a prevalence of 8% among Asian patients. Differences in the incidence rate of HCV results attained from various regions globally depict geographical diversity. The variation in these occurrence rates can be ascribed to exposure to various risk factors which are capable of enhancing the spread and transmission of this virus amongst individuals(19,20).

The Prevalence rate of HCV infection among males, recorded 8(4.4%) while the female subjects had 14(8.9%) seropositivity. This finding agrees with the result obtained from a similar work by Ndako *et al.*, (22) and Gacche *et al.*, (23) where the incidence of anti-HCV in diabetic among the female subjects were higher compared to the male subjects. Increased rate of occurrence in females could be attributed to various risk factor HCV infection, which was quite evident from the life style and history of the individuals recruited for this study(22,23).

A higher prevalence rate of 9(5.0%) was observed amongst subjects aged 41-50, this result is in accordance with the findings of Tessema *et al.*,(24), where the sero-prevalence of HCV increases as age of participants increased and it was significantly higher in the age group of 41 – 50 years, which is also closer to the results obtained by Klevens *et al.*, (25) which showed a higher incidence rate among subjects

aged 35-44 and this concurs with the result obtained in this study. The high seropositivity observed in older age group could be attributed to possible differences in social practices, parenteral exposures, decline in physical mobility and a reduced rate of medical examination compared to younger individuals thus increasing chances of transmission of infection.(26).

The result obtained in this study showed that having multiple sexual partners is a major means of spreading HCV; however, it was observed that the prevalence of HCV was higher among married subjects (15.4%), similar study carried out by Qureshi *et al.*, (27) recorded a higher prevalence among married subjects. This findings could be as a result of increased exposure to several risk factors to this infectious agent Simo *et al.*, (28).

Regardless of the fact that HCV is a blood-borne virus, the sero-prevalence among individuals that had undergone blood transfusion or donation is of no statistical significant difference and this agrees with the work of Simo *et al.*, (28).Among subjects that had history of blood donation in this study, 3 (1.7%) were found to be positive to HCV. However, it is observed that blood and blood products are potential sources of transmission for HCV infection Ndako *et al.*, (22). In the present study, sharing personal items that may be contaminated with infected blood and tattooing were defined as HCV predictors. The risk of HCV continues to be a great occupational threat. Consequently, blood transfusion was also identified as a predictor of HCV Infection(30).

High rate of positivity was observed in subjects screened for alcohol consumption, sharing of unsterilized objects and subjects that had tribal marks or tattoo ranging from 3.9%, 2.2% and 2.8% respectively with no statistical significant difference. However, participants with family history of diabetes recorded a sero-prevalence of 6.7% to HCV infection, This report is similar to the work of Muller *et al.*, (32) where the increased incidence of HCV was closely related with family history of diabetes mellitus,A significant difference was observed in participants with a family history of diabetes mellitus and those without, (32,33); this might be attributed to multiple sexual partners, transfusion of unscreened blood in hospitals, family history of related infections, risky behaviours such as alcohol intake and other potential unidentified routes of transmission which can only be discovered through advance studies(34).

This study found that elevated liver enzymes; especially ALT has a direct relationship with seropositivity to HCV in the diabetic population studied, showing the relevance of this as a screening test in diabetics. In a study by Mason, (29) more than 20% of diabetes patients with consistently elevated serum aminotransferases had evidence of HCV infection. It was also discovered that most of the anti-HCV positive diabetic patients presented with an abnormal liver function tests, a combination of hepatocellular and cholestasis pattern being the predominant biochemical alteration, Osi and Sanna., (31). Elevation of ALT in hepatitis C positive diabetes patients in this study is usually mild, with most having ALT level between one to two times upper limit of normal.

Conclusion

The outcome of this study showed an increased risk of HCV infection in patients with T2DM, which warrants routine screening of diabetic subjects for HCV. Similarly, the study adds to the limited data on the subject available in this region and will help at increasing awareness regarding association of HCV and diabetes, which will further help to reduce morbidity and possible mortality associated with this comorbidity in the long run. The need for prompt enlightenment of the general public on the threats of the co-infectious nature of HCV and diabetes is strongly advocated. In addition, it is important that health care workers, approach the settings of early diagnosis and management of this condition among infected persons to avert further complications among type 2 diabetics.

Abbreviations

HCV-Hepatitis C Virus; T2DM-Type 2 Diabetes Mellitus; ALT-Alanine Transaminases; ELISA-Enzyme-Linked Immunosorbent Assay.

Declarations

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Authors Contributions:

JAN- Made substantial contributions to the drafting, conception; field analysis and sample collection of this Research.

AOO- Made substantial contributions on design module of this research.

JAO- Made substantial contributions on the Monitoring of the subjects recruited for this study.

JAA- Made substantial contributions on the laboratory analysis of the samples obtained.

OO- Made substantial contributions on the interpretation of the statistical analysis used.

BAA-Made substantial contributions on the field work and the analysis of the samples obtained.

All authors have read and approved the manuscript.

Authors Information:

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Ethics approval and consent to participate: Ethical clearance and approval was obtained from the appropriate ethical committee of the Federal Teaching Hospital Idi-Nigeria.

Consent for publication: Only consented volunteers, obtained through volunteer/consent forms filled by respective subjects were enrolled for the study. I wish to declare that consent for participation obtained from study participants was firstly verbal and later documented through a consent form. Study participants were also counseled on the relevance of the screening before sample collection was embarked upon. This was explained to the ethics committee and was so approved.

Availability of supporting data and material: The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Competing interests: The authors declare that they have no competing interests.

References

1. Rosen HR. Chronic hepatitis C infection. *N Engl J Med*. 2011;364(25):2429–38.
2. Gane EJ. The natural history of recurrent hepatitis C and what influences this. *Liver Transplant*. 2008;14(2):36–44.
3. Foster GR, Goldin R. Management of chronic viral hepatitis. Taylor Fr Gr. 2005;2nd Edition.
4. Shinn JH, Liang KC. Chronic hepatitis C and Diabetes Mellitus. *Chinese J Med Assoc*. 2006;69(4):143–5.
5. Alter MJ. Epidemiology of Hepatitis C virus. *J Hepatol*. 1997;26:625–55.
6. Vescovo T, Refolo G, Vitagliano G, Fimia GM, Piacentini M. Molecular mechanisms of hepatitis C virus–induced hepatocellular carcinoma. *Clin Microbiol Infect*. 2016;22(10):853–61.
7. Elfiky AA, Elshemey WM, Gawad W, Desoky O. Molecular modeling comparison of the performance of NS5b polymerase inhibitor (PSI-7977) on prevalent HCV genotypes. *Protein J*. 2013;32(1):75–80.
8. Mehta, SH., Brancati, FL., Strathdee, SA. Pankow, JS., Netski, D., Coresh, J., Szklo, M., and Thomas, DL. Hepatitis C virus infection and incident type 2 diabetes *Hepatology*. 2003; 38(1):50-56. doi: 10.1053/jhep.50291.
9. Imam K. Clinical features, diagnostic criteria and pathogenesis of diabetes mellitus. *Adv Exp Med Biol*. 2012;771:340–55.
10. Peck T, Price C, English P, Gill G. Oral health in rural South African type 2 diabetic patients. *Trop Doct*. 2006;36(2):111–2.

11. Atkinson MA, Eisenbarth GS, Michels AW. Type 1 diabetes. *Lancet*. 2014;383:69–82.
12. Knobler, H, Schihmanter, R, Zifroni, A, Fenakel, G, Schattner, A. Increased risk of type 2 diabetes in noncirrhotic patients with chronic hepatitis C virus infection. *Mayo Clin Proc* 2000; 75: 355–359.
13. Zein CO, Levy C, Basu A, Zein NN. Chronic hepatitis C and type II diabetes mellitus: a prospective cross-sectional study. *Am. J Gastroenterol*. 2005;100 (1):48–55.
14. Younossi ZM, Stepanova M, Nader F, Younossi Z, Elsheikh E. Associations of chronic hepatitis C with metabolic and cardiac outcomes. *Aliment Pharmacol Ther*. 2013;37(6):647–52.
15. Naing C, Mak JW, Ahmed SI, M. Maung M. Relationship between hepatitis C virus infection and type 2 diabetes mellitus: meta-analysis. *World J Gastroenterol*. 2012;18(14):1642–51.
16. Nwokediuko SC, Oli JM. Hepatitis C Virus Infection in Nigerians with diabetes mellitus. *Niger J Clin Pract*. 2008;11(2):94–9.
17. Ejele O, Erhabor O, Nwauche CA. The risk of transfusion-transmissible viral infections in the Niger-Delta area of Nigeria. *Sahel Med J*. 2005;8(1).
18. Balogun WO, Adeleye JO, Akinlade KS, Kuti M, Otegbayo JA. Low prevalence of hepatitis-C viral seropositivity among patients with type-2 diabetes mellitus in a tertiary hospital. *J Natl Med Assoc*. 2006;98(11):1805.
19. Ndako,J.A., Nwankiti,O.O., Onovoh,E.M., Adekeye,A.E.,Choji,T.P.,Alesa,M.U and Alarape,A.J.. Screening response to Hepatitis C Virus antibodies among Diabetic patients attending UITH Nigeria,Current Research Journal of Biological Sciences.2011;3(6): 542-546
20. Demitrost L, Ranabir S. Thyroid dysfunction in type 2 diabetes mellitus: A retrospective study. *Indian J Endocrinol Metab*. 2012; 16(Suppl 2):S334–335.
21. Gray H, Wreghitt T, Stratton IM, Alexander GJ, Turner RC, O’Rahilly S. High prevalence of hepatitis C infection in Afro-Caribbean patients with type 2 diabetes and abnormal liver function tests. *Diabet Med*. 1995;12:244–249
22. Ndako,J.A., Echeonwu,G.O.N., Shidali,N.N., Bichi,I.A., Paul,G.A., Onovoh,E. and Okeke,L.A. Occurrence of Hepatitis C Virus infection in type 2 diabetic patients attending Plateau state specialist hospital Jos Nigeria *Virology Journal*;2009; 6:98 doi: 10.1186/1743-422X-6-98.
23. Gacche RN, Al-Mohani SK. Seroprevalence and Risk Factors for Hepatitis C Virus Infection among General Population in Central Region of Yemen. *Hepatitis Research and Treatment*. 2012: 1-4.
24. Tessema B, Yismaw G, Kassu A, Amsalu A, Mulu A, Emmrich F, Sack U. Seroprevalence of HIV, HBV, HCV and syphilis infections among blood donors at Gondar University Teaching Hospital, Northwest Ethiopia: declining trends over a period of five years. *BMC Infect. Dis*. 2010;10:111
25. Klevens RM, Miller J, Vonderwahl C, Speers S, Alelis K, Sweet K, et al. Population-based surveillance for hepatitis C virus, United States. *Emerg Infect Dis*. 2009;15(9):1499.
26. Lee SR, Kardos KW, Schiff E. Evaluation of a new, rapid test for detecting HCV infection, suitable for use with blood or oral fluid. *J Virol Methods*. 2011;172(1–2):27–31.

27. Qureshi H, Bile KM, Jooma R, Alam SE, Afrid. HUR. Prevalence of hepatitis B and C viral infections in Pakistan: findings of a national survey appealing for effective prevention and control measures. East Mediterr Heal J. 2010;16(8):15–23.
28. Simo R, Hernadez C, Genesa J, Jardi R, Mesa J. High prevalence of Hepatitis virus infection in diabetic patients. Am Diabetes Assoc. 1996;19(9):998–1000.
29. **Mason, A. L., Lau, J.Y., Hoang, N., Qian, K., Alexander, G,J., Xu, L, Guo,L., Jacob,S., Regenstein, F.G., Zimmerman, R., Everhart, J.E., Wasserfall, C.,Maclaren,N.K., and R. P. Perrillo., R.P.** Association of diabetes mellitus and chronic hepatitis C virus infection. Hepatology 1999; 29:328-333
30. Cardinal GF, Di Martino V, Lambrey G, Nalet B, Anciaux M. Prevalence of hepatitis C infection and risk factors in hospitalized diabetic patients: results of a cross-sectional study. Eur J Gastroenterol Hepatol. 2008;20(9):829–36.
31. Francesca W, Lutje. V, Declan D, Valerie S. Sexual transmission of Hepatitis C Virus infection in a heterosexual population: A systematic review [version 1; referees: 2 approved]. HRB Open Res. 2018
32. Mueller, H. M., Pfaff, E., Goeser, T., Kallinowski, B., Solbach, C. &Theilmann, L. Peripheral blood leukocytes serve as a possibleextrahepatic site for hepatitis C virus replication. Journal of GeneralVirology, 1993; 74, 669-676.
33. Osi O, Sanaa MK. Hepatitis C in Developing countries. Acad Press. 2018;71–81.
34. Ghany MG, Strader DB, Thomas DL, Seeff LB. Diagnosis, management and treatment of hepatitis C. J Hepatol. 2009;

Tables

TABLE 1a: Distribution of sera samples assayed.

Total number of samples	Number of positive samples	Number of negative samples
180	24 (13.3%)	156 (86.7%)

TABLE 1bDistribution of HBV among Non-Diabetics (Control Subjects) based on gender.

SEX	Number tested	HBV	
		Positive %	Negative%
Male	48	4(4%)	44 (44%)
Female	52	5 (5%)	47 (47%)
Total	100	9 (9%)	91 (91%)

P=0.739; P>0.05

TABLE 2: Distribution of Hepatitis C Virus among diabetic subjects based on Gender

Gender	Total number of samples examined (%)	Number of samples (%)	Positive Number of samples (%)	Negative
Male	71(39.4)	8(4.4)	63(35.0)	
Female	109(60.6)	16(8.9)	93(51.7)	
Total	180(100.0)	24(13.3)	156(86.7)	

Chi square (χ^2) = 0.433; df = 1; P value = 0.511

TABLE 3: Distribution of Hepatitis C Virus based on Age of subjects screened.

Age	Total number of samples examined (%)	Number of samples (%)	Positive Number of samples (%)	Negative
0-20	20 (11.1)	0 (0.0)	20 (11.1)	
21-30	20 (11.1)	2 (1.1)	18 (10.0)	
31-40	37 (20.6)	6 (3.3)	31 (17.2)	
41-50	40 (22.2)	9 (5.0)	31 (17.2)	
51-60	24 (13.3)	3 (1.7)	21 (11.7)	
60-100	39 (21.7)	4 (2.2)	35 (19.4)	
Total	180 (100.0)	24 (13.3)	156 (86.6)	

Chi square (χ^2) = 6.778; df = 5; P value = 0.238

TABLE 4: Distribution of Hepatitis C Virus According To Marital Status

Marital Status	Total number of samples examined (%)	Number of samples (%)	Positive	Number of Negative samples (%)
Single	41 (22.8)	4 (2.2)		37 (20.6)
Married	130 (72.2)	20 (11.1)		110 (61.1)
Divorced	9 (5.0)	0 (0.0)		9 (5.0)
Total	180 (100.0)	24 (13.3)		156 (86.7)

Chi square (x^2) = 2.312; df = 2; P value = 0.315

TABLE 5: Distribution of Hepatitis C Virus According To Educational Background

Education	Total number of samples examined (%)	Number of samples (%)	Positive	Number of Negative samples (%)
Primary	6 (3.3)	0 (0.0)		6 (3.3)
Secondary	25 (13.9)	2 (1.1)		23 (12.8)
Tertiary	148 (82.2)	22 (12.2)		126 (70.0)
No Education	1 (0.6)	0 (0.0)		1 (0.6)
Total	180 (100.0)	24 (13.3)		156 (86.7)

Chi square (x^2) = 1.993; df = 3; P value = 0.574

TABLE 6: Distribution of Hepatitis C Virus based on Demographic Factor.

Occupation	Total number of samples examined (%)	Number of samples (%)	Positive Number of samples (%)	Negative
Trading	64 (35.6)	7 (3.9)	57 (31.7)	
Civil Servant	64 (35.6)	14 (7.8)	50 (27.8)	
Industry	18 (10.0)	2 (1.1)	16 (8.9)	
Student	34 (18.9)	1 (0.6)	33 (18.3)	
Total	180 (100.0)	24 (13.3)	156 (86.7)	

Chi square (χ^2) = 7.613; df = 3; P value = 0.055

TABLE 7: Distribution of Hepatitis C Virus based on Clinical Risk Factors.

Risk Factor		Number of Positive Samples	of Number of Negative Samples	of Total Number of Samples Examined	Chi-Square
	Positive	1 (0.6%)	14 (7.8%)	15 (8.3%)	0.629^a
	Negative	23 (12.8%)	142 (78.9%)	165 (91.7%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	P value = 0.4276
Blood Transfusion Blood Donation	Positive	3 (1.6%)	10 (5.6%)	13 (7.2%)	1.151^a
	Negative	21 (11.7%)	146 (81.1%)	167 (92.7%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	P value = 0.2833
CareforanHepatitis C patient	Positive	6 (3.3%)	32 (17.8%)	38 (21.1%)	0.251^a
	Negative	18 (10.0%)	124 (68.9%)	142 (78.9%)	df = 1
	Total	24 (13.3%)	156 (86.7%)	180 (100.0%)	P value = 0.6160

TABLE 8: Distribution based on lifestyle-risk factors of Subjects Screened.

Factor	Response	Number Positive Samples	of	Number Negative Samples	of	Total Number of Samples Examined	Chi-Square
Alcohol consumption	Positive	7 (3.9%)		30 (16.7%)		37 (20.6%)	1.257 ^a
	Negative	17 (9.4%)		126 (70.0%)		143 (79.4%)	df = 1
	Total	24 (13.3%)		156 (86.7%)		180 (100.0%)	Pvalue = 0.2621
Facial Marks Tattoos	Positive	4 (2.2%)		15 (8.3%)		19 (10.6%)	1.095 ^a
	Negative	20 (11.1%)		141 (78.4%)		161 (89.4%)	df = 1
	Total	24 (13.3%)		156 (86.7%)		180 (100.0%)	Pvalue = 0.2953
Wearing of sterilized garment	Positive	5 (2.8%)		51 (28.3%)		56 (31.1%)	1.365 ^a
	Negative	19 (10.6%)		105 (58.3%)		124 (68.9%)	df = 1
	Total	24 (13.4%)		156 (86.6%)		180 (100.0%)	Pvalue = 0.2427
Multiple sexual partners	Positive	3 (1.7%)	4 (2.2%)	7 (3.9%)		5.494 ^a	
	Negative	21 (11.7%)	152 (84.4%)	173 (96.1%)		df = 1	
	Total	24 (13.4%)	156 (86.6%)	180 (100.0%)		Pvalue = 0.0191	

TABLE 9: Distribution of Hepatitis C Virus According To Family History

Risk Factor		Number Positive Samples	of	Number Negative Samples	of	Total Number of Samples Examined	Chi-Square	
Previous Record Hepatitis Virus	of	Positive		1 (0.6%)		4 (2.2%)	5 (2.8%)	0.198 ^a
		Negative		23 (12.8%)		152 (84.4%)	175 (97.2%)	df = 1
		Total		24 (13.4%)		156 (86.6%)	180 (100.0%)	Pvalue = 0.6565
Diabetes Family Members	by	Positive		12 (6.7%)		88 (48.9%)	100 (55.6%)	0.346 ^a
		Negative		12 (6.7%)		68 (37.7%)	80 (44.4%)	df = 1
		Total		24 (13.4%)		156 (86.6%)	180 (100.0%)	Pvalue = 0.5563
Hepatitis Family Members	by	Positive		3 (1.7%)		29 (16.1%)	32 (17.8%)	0.528 ^a
		Negative		21 (11.6%)		127 (70.6%)	148 (82.2%)	df = 1
		Total		24 (13.3%)		156 (86.7%)	180 (100.0%)	Pvalue = 0.4676

Table 10: Determination of Serum Alanine Aminotransferase on Hepatitis CVirus patients.

Age	No. Seropositive For HCV (%)	Normal ALT range (%)	Abnormal ALT range (%)
0-20	0(0%)	Not applicable	Not applicable
21-30	2(1.1%)	2(1.1%)	0(0%)
31-40	6(3.3%)	3(1.7%)	3(1.7%)
41-50	9(5.0%)	6(3.3%)	3(1.7%)
51-60	3(1.7%)	1(0.6%)	2(1.1%)
61-100	4 (2.2%)	2(1.1%)	2(1.1%)
Total	24(13.3%)	14(7.8%)	10(5.5%)

Figures

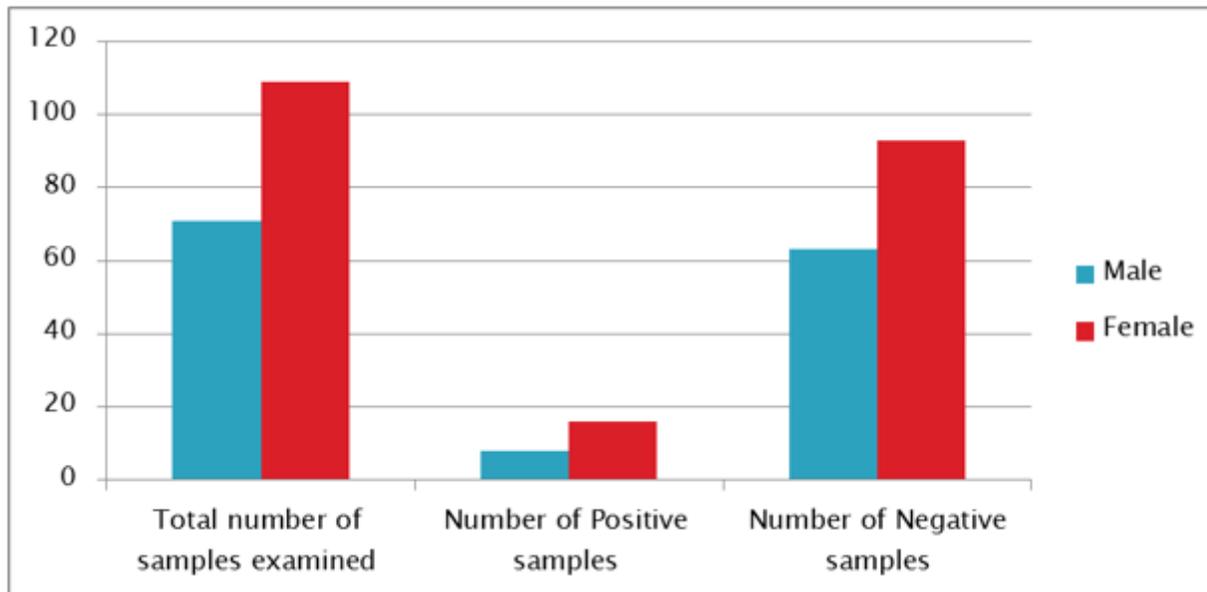


Figure 1

Distribution of Hepatitis C Virus according to Gender

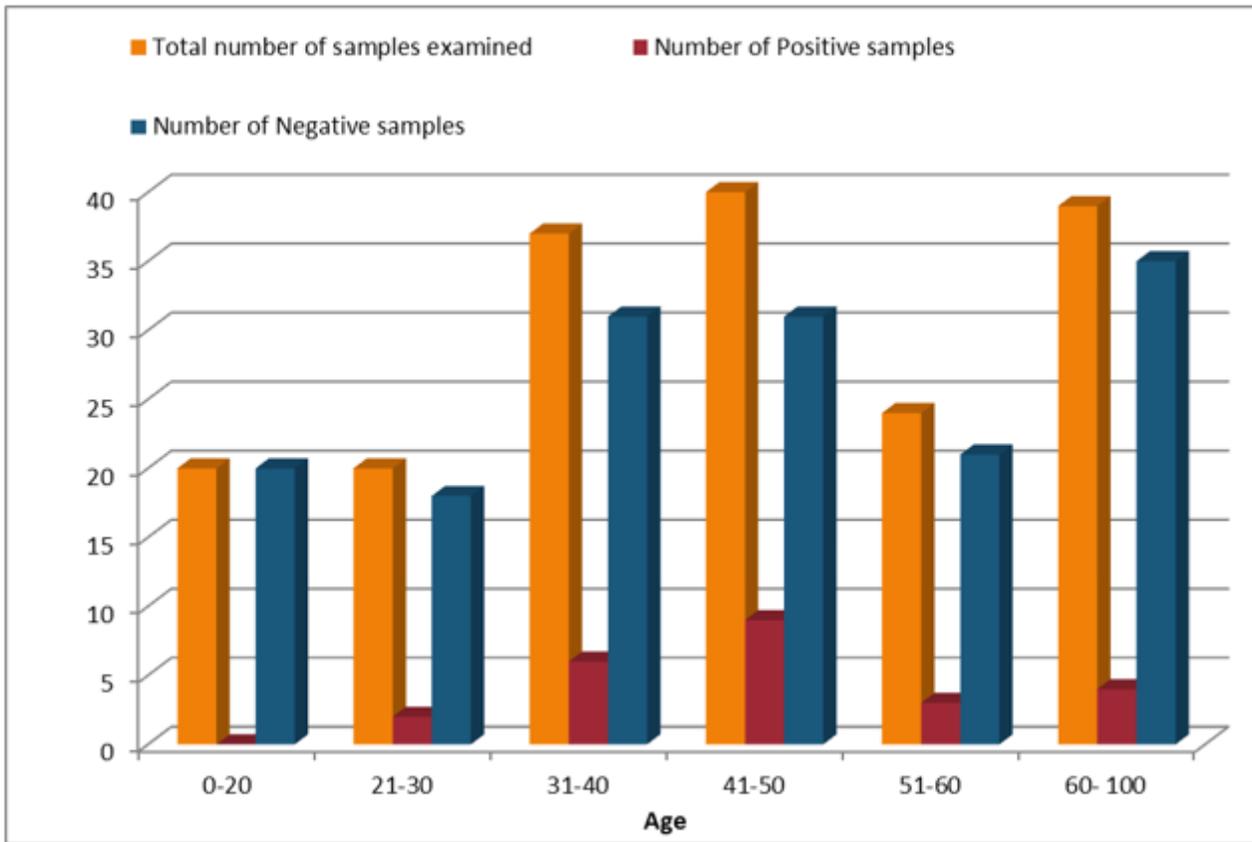


Figure 2

Distribution of Hepatitis C Virus according to Age

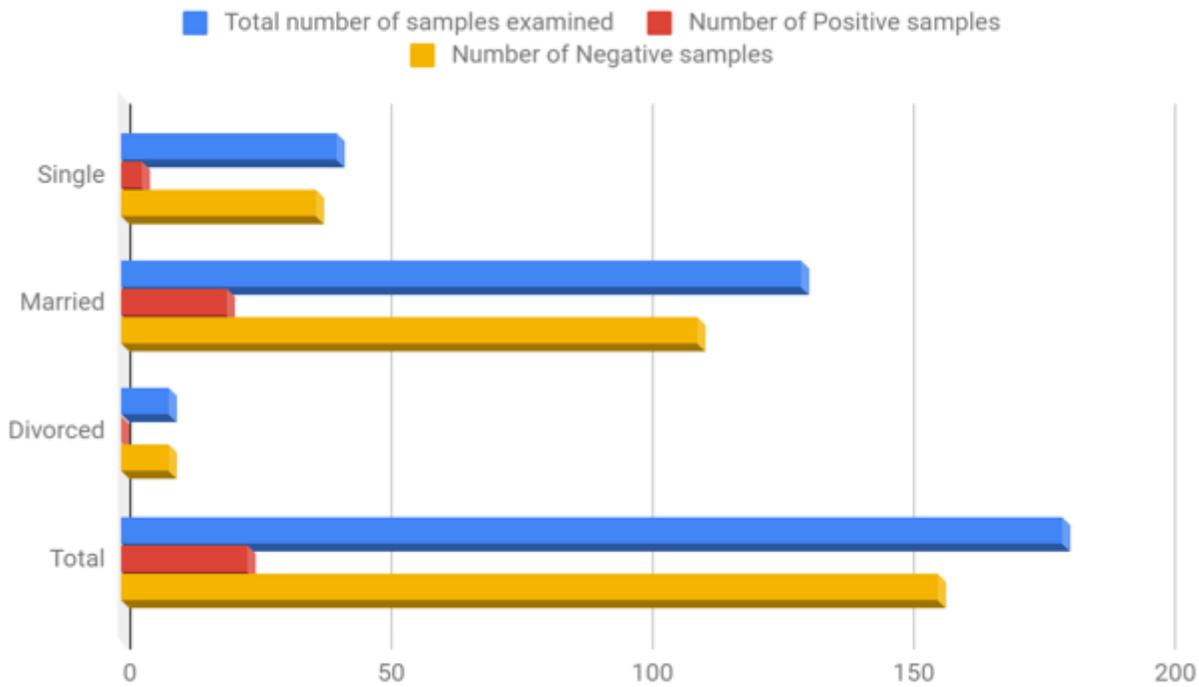


Figure 3

Distribution of Hepatitis C Virus according to marital status

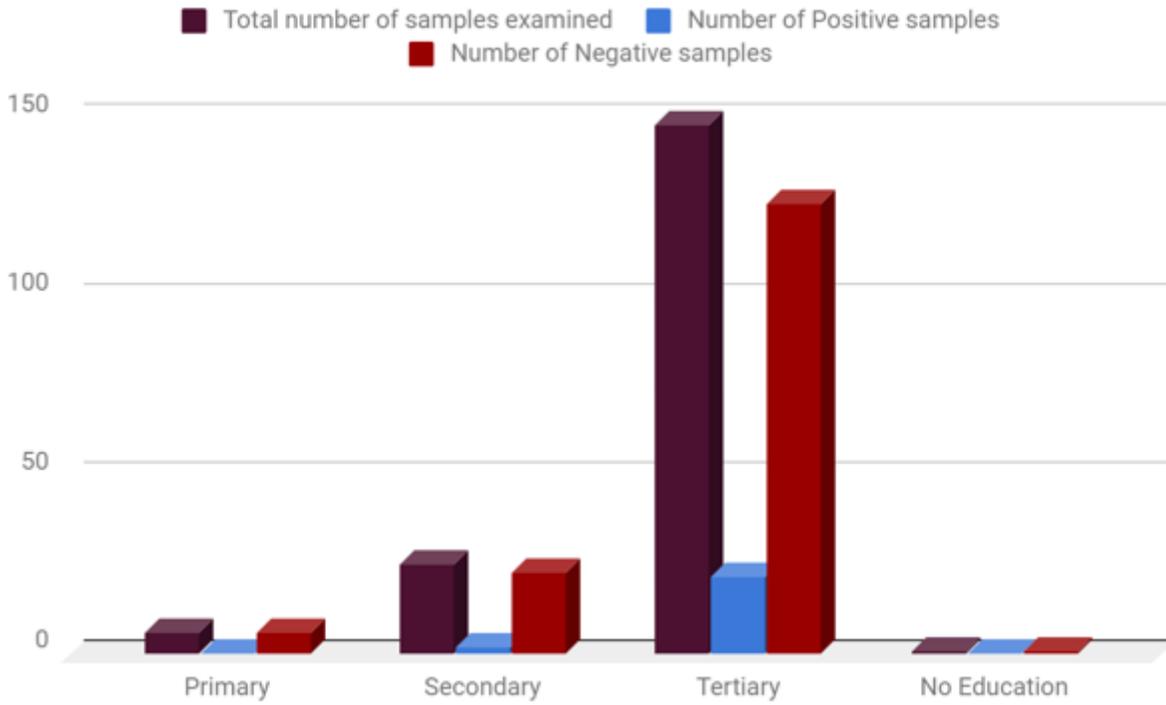


Figure 4

Distribution of Hepatitis C Virus according to Educational background.

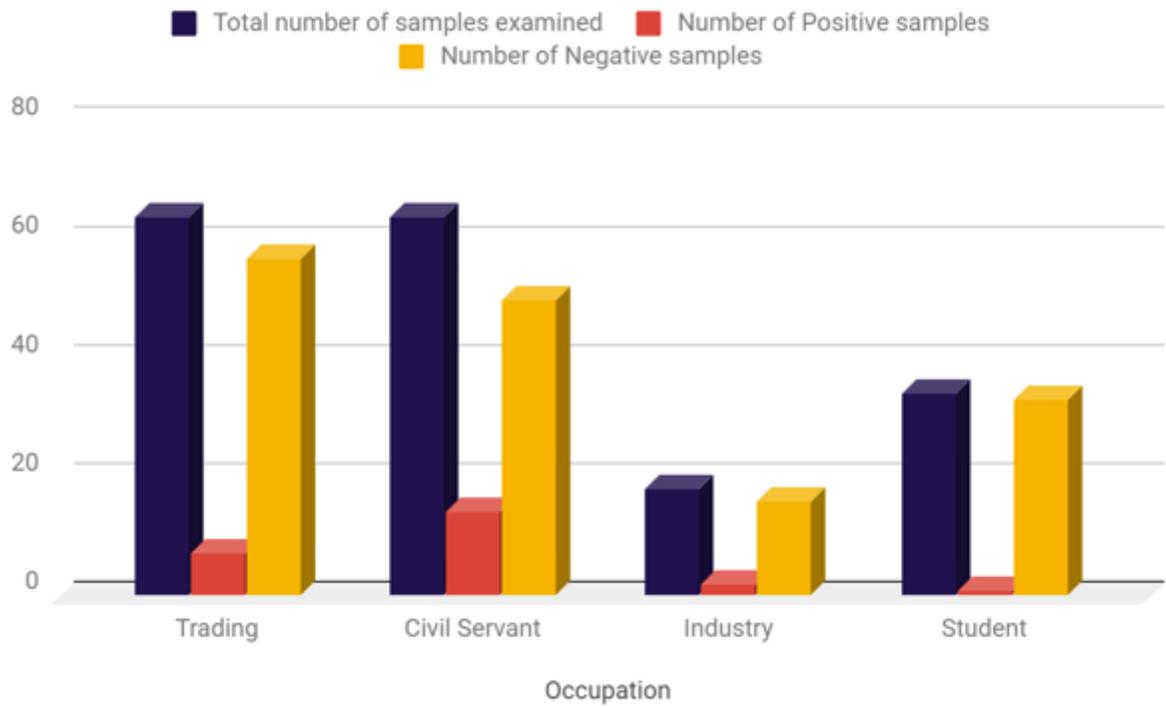


Figure 5

Distribution of Hepatitis C Virus according to Age.

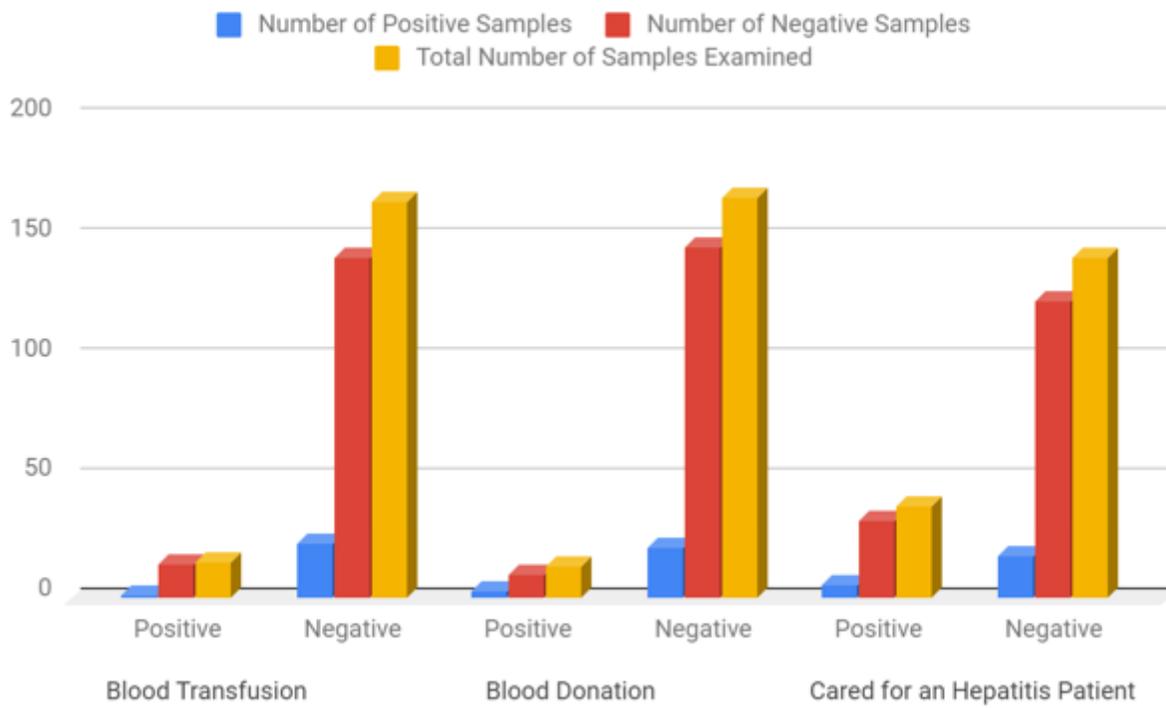


Figure 6

Distribution of Hepatitis C Virus based on Clinical Risk Factors