

29 cases of HCV also continue to occur as a result of injecting drug use (IDU) and through
30 other means of percutaneous or mucous membrane exposure [2]

31 HIV infection in humans is now pandemic as of January 2006, the Joint United
32 Nations Programme on HIV/AIDS (UNAIDS) and the world health organization (WHO)
33 estimates that AIDS has killed more than 25 million people since it was first recognized
34 in 1981, making it one of the most destructive pandemics in recorded history (. It is
35 estimated that about 0.6% of the world's population is infected with HIV [3]. HIV
36 prevalence varies widely by geographic region and vulnerable population, Nigeria has an
37 overall national prevalence of 3.1% but statewide; HIV prevalence among pregnancy
38 women has ranged from as low as 1.6% in Ekiti in west to 10 % in Benue in south east
39 [4].

40 However, the current estimate state that HIV is set to infect 90 millions people in
41 Africa, resulting in a minimum estimate of 18 million orphans. Antiretroviral drug reduce
42 mortality and morbidity of HIV infection, but routine access to antiretroviral medications
43 is not available in all countries [5]. HCV co-infection with HIV is common and rates
44 among HIV positive population are higher [6]. About 10,000-20,000 death yearly in US
45 is from HCV; expectations are that this mortality rate will increase, as those who were
46 infected by transfusion before HCV testing become apparent. It is responsible for 90-95%
47 of all transfusion-related hepatitis [7].

48 Tuberculosis (TB) has been major public health problems for centuries. The
49 implementation of effective public health interventions for the prevention and control of
50 TB has significantly contributed to a substantial reduction of the global disease but,
51 however, the emergence of the HIV epidemic has posed major challenges to TB control
52 effort globally. In a country with almost 40 % population already infected with TB, an

53 increasing prevalence of HIV will be jeopardize TB central effort with such
54 consequences [8]. HIV has been thought to account for much of the recent increase in the
55 global TB burden, especially in Africa. [9].

56 HIV is the most important risk factor for the development of TB among person
57 infected with *M. tuberculosis* and both **Centres for Disease Control and Prevention**
58 **(CDC)** and WHO guidelines recommend offering HIV testing to those person diagnosed
59 with TB disease [2].

60 The prevalence of HCV infection among persons with TB has been poorly
61 **defined**, and few data are available from most areas around the world. One recent study
62 from US has suggested that veterans with HCV infection are at risk for other selected
63 infectious disease including TB. Part of the lack of data on HCV seroprevalence stems
64 from the fact that there is no recommendation for universal screening of person with TB
65 for HCV infection as there are for HIV testing [2].

66 Richard *et al*, [2] reported that HIV and HCV are both global public health
67 problems infections with HIV and or HCV may have an impact among those with TB.
68 The high presence of HCV co-infection among patients with TB in Georgia has the
69 potentials to have a major impact on TB management, treatment and control.

70 Hepatitis C virus is one of the deadly blood-borne virus that has almost the same
71 route of transmission as of HIV, it is noted to have its major activity in the liver where it
72 causes inflammation of the liver, on the other hand, HIV and TB are closely connected
73 that they are often referred to as co-epidemics among confirmed TB patient, however,
74 drug or regimes given to TB or TB/HIV co-infection patient has hepatotoxicity effect and
75 can eventually lead to hepatocellular carcinoma

76 In this study, the seroprevalence of HCV and HIV in confirmed TB patient were
77 determined in order to provide an updated reference data for effective empiric
78 management of Tuberculosis patients with co-infection of HCV and HIV. Also, the
79 possible predisposition factor(s) to HCV and HIV coinfection in TB patients were
80 identified.

81 **2.0. METHODOLOGY.**

82 **2.1 Study Area.**

83 The study area for this work was Federal Medical Centre (FMC), Ido Ekiti located
84 in Ekiti North senatorial district of Ekiti State, Nigeria.

85 **2.2. Study population**

86 The study population is Tuberculosis confirmed patients attending FMC, Ido
87 Ekiti. A total number of 500 samples were collected from TB confirmed patient after due
88 consultation with the patients.

89 **2.3 Ethical consideration**

90 The ethical clearance for this research was granted by Federal Medical Centre
91 (FMC) (Now Federal Teaching Hospital) Ido-Ekiti ethical committee after due processes
92 had been followed. Before the collection of the sample, information regarding the study
93 was explained to the subjects.

94 **2.4 Questionnaire and informed consent**

95 Questionnaire to obtain the demographic characteristics, possible risk factors and
96 other relevant information to the study as well as an informed consent were administered
97 to the participant.

98 **2.5 Sample collection**

99 About 5mls of blood was collected by venous puncture from the antecubital foci
100 of the arm after disinfecting the area with 70% alcohol. The blood was allowed to clot
101 and was spun at 1000rpm for 5 minutes and the serum was aseptically collected into
102 sterile cryovials bottles, appropriately labeled and stored at -20⁰C until the test was
103 performed.

104 **2.5 Sample processing**

105 **2.5.1 HCV Detection**

106 **Clinotech diagnosis anti-HCV cassette detection** test was used which is a rapid
107 direct binding procedure, which visually determines antibodies to hepatitis C infection.

108 **2.5.2. Detection of HIV**

109 The **Abbot Determine** HIV-1/2 was used in conjunction with STAT-PAK which
110 are invitro, visually ready, qualitative immunoassays for the detection antibodies to HIV-
111 1 and HIV-2 in human serum, plasma or whole blood.

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113 **2.6. Statistical analysis**

114 The data generated from this study were analysed using SPSS version 16 (SPSS Inc.
115 Chigago IL).

116

117 **3.0 RESULT**

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119 **3.1. Response rate**

120 A total of 500 questionnaires and consent forms were distributed to the patients
121 screened and out of 500 questionnaires distributed, 500 were returned indicating a 100%
122 response rate.

123 The overall seroprevalence of HCV, HIV and HCV/HIV antibodies in tuberculosis
124 confirmed patients are shown in table 1. It shows that out of 500 samples tested for HCV,
125 HIV antibodies, 10(2.0%) are positive for HCV, 21(4.2%) are positive for HIV and
126 3(0.6%) are positive for both HCV and HIV antibodies.

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128 **TABLE 1: OVERALL SEROPREVALENCE OF HCV, HIV AND HCV/HIV**
129 **ANTIBODIES IN TB CONFIRMED PATIENT.**

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| INFECTION | NO OF SAMPLES | NO OF POSITIVE (%) |
|-----------|---------------|--------------------|
| HIV | 500 | 21(4.2) |
| HCV | 500 | 10(2.0) |
| HIV/HCV | 500 | 3 (0.6) |

144 The demographic relationships in respect to sex are shown in table 2, table 3 and
145 table 4 for HCV, HIV and HCV/HIV co-infection respectively. They revealed that out of
146 302 male subjects that participated, 7(2.32%) positive for HCV (table 2), 14(4.64%)
147 positive for HIV (table 3) and 2 (0.66%) were positive for HCV/HIV antibodies (table
148 4), while out of 198 female, 3(1.52%) positive for HCV(table 2), 7(3.54%) positive for
149 HIV (table 3), 1(0.51%) positive for HCV/HIV antibodies (table 4).

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152 **TABLE 2: SEROPREVALENCE OF HCV IN RELATION TO SEX.**

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| SEX | NO POSITIVE (%) | P VALUE |
|--------|-----------------|---------|
| Male | 7.0 (2.32) | 0.531 |
| Female | 3.0 (1.52) | |

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158 Male 7.0 (2.32)

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161 0.531

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163 Female 3.0 (1.52)

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170 **TABLE 3: SEROPREVALENCE OF HIV IN RELATION TO SEX**

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| SEX | NO POSITIVE (%) | P VALUE |
|--------|-----------------|---------|
| Male | 14.0 (4.64) | 0.549 |
| Female | 7.0 (3.54) | |

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174 SEX NO POSITIVE (%) P VALUE

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178 Male 14.0 (4.64)

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181 0.549

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183 Female 7.0 (3.54)

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192 **TABLE 4: SEROPREVALENCE OF HCV/HIV CO-INFECTION IN RELATION TO SEX**

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| SEX | NO POSITIVE (%) | P VALUE |
|------|-----------------|---------|
| Male | 2.0 (0.66) | |

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197 Male 2.0 (0.66)

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|--------|-------------|-------|
| Female | 1.00 (0.51) | 0.824 |
|--------|-------------|-------|

Age group distributions for HCV, HIV and HCV/HIV antibodies are shown in table 5, table 6 and table 7 respectively. In age group 18-25, out of 4 (0.8%) subjects that participated, no subjects were positive for HCV and HIV antibodies as shown in table 5 and table 6 respectively. In age group 26-30, 20 (4.0%) subjects participated, 4 (0.8%) positive for HCV(table 5), 2(0.4%) positive for HIV (table 6), but no subject had HCV and HIV together (table 7). 70(14%) subjects are within 36-45 age group, 5(1%) had HCV (table 5), 10(2%) had HIV(table 6), 2(0.4%) had HCV/HIV antibodies (table 7). In age group 46-55, 136(27.2%) participated, 1(0.2%), 5(1%), and 1(0.2%) were seropositive for HCV, HIV, and HCV/HIV antibodies and these were shown in table 5, table 6 and table 7 respectively. Out of 150(30%) subjects within 56-65 age group, 3(0.6%) were seropositive for HIV (table 6), no subject was positive for HCV (table 5) and HCV/HIV antibodies (table 7). In age 66-75, 70(14%) participated, 1(0.2%) was positive for HIV (table 6), no seropositivity in HCV and HIV/HCV as shown in table 5 and table 7 respectively. 50(10%) subjects participated in age group 75- above, no seropositivity was recorded in both HCV and HIV as shown in table 5 and table 6 respectively.

224 **TABLE 5: SEROPREVALENCE OF HCV IN RELATION TO AGE.**

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227 **AGE GROUP** **NOT EXAMINED (%)** **NO POSITIVE (%)** **P**

228 **VALUE**

229 **IN YEARS**

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| | | | |
|--------------|-------------|----------|-------|
| 233 18-25 | 4 (0.8) | 0 (0.0) | |
| 234 26-35 | 20 (4.0) | 4 (0.8) | |
| 235 36-45 | 70 (14.0) | 5 (1.0) | |
| 236 46-55 | 136 (27.2) | 1 (0.2) | 0.000 |
| 237 56-65 | 150 (30.0) | 0 (0.0) | |
| 238 66-75 | 70 (14.0) | 0 (0.0) | |
| 239 75-above | 50 (10.0) | 0 (0.0) | |
| 240 | | | |
| 241 TOTAL | 500 (100.0) | 10 (2.0) | |

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247 **TABLE 6: SEROPREVALENCE OF HIV IN RELATION TO AGE.**

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249 **AGE GROUP** **NO EXAMINED (%)** **NO POSITIVE (%)** **P VALUE**

250 **IN YEARS**

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| | | | |
|--------------|-------------|----------|-------|
| 255 18-25 | 4 (0.8) | 0 (0.0) | |
| 256 26-35 | 20 (4.0) | 2 (0.4) | |
| 257 36-45 | 70 (14.0) | 10 (2.0) | 0.000 |
| 258 46-55 | 136 (27.2) | 5 (1.0) | |
| 259 56-65 | 150 (30.0) | 3 (0.6) | |
| 260 66-75 | 70 (14.0) | 1 (0.2) | |
| 261 75-Above | 50 (10.0) | 0 (0.0) | |
| 262 | | | |
| 263 TOTAL | 500 (100.0) | 21 (4.2) | |

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269 **TABLE 7: SEROPREVALENCE OF HCV/HIV CO-INFECTION IN RELATION TO AGE**

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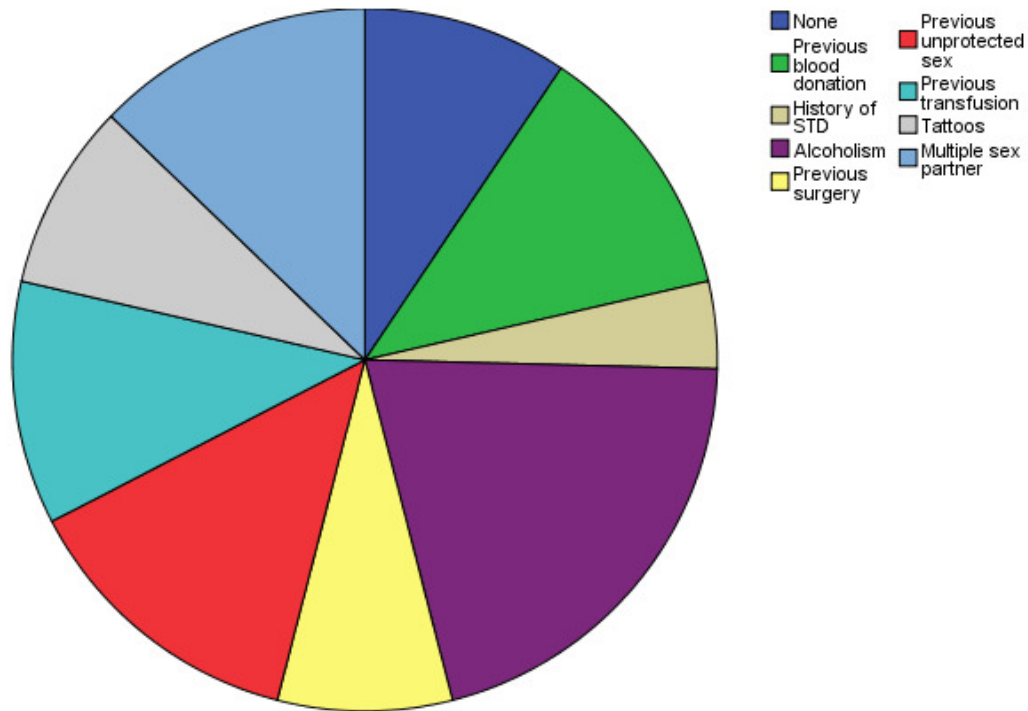
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| AGE GROUP IN YEARS | NO EXAMINED (%) | NO POSITIVE (%) | P VALUE |
|-----------------------|-----------------|-----------------|---------|
| 18-25 | 4 (0.8) | 0 (0.0) | 0.002 |
| 26-35 | 20 (4.0) | 0 (0.0) | |
| 36-45 | 70 (14.0) | 2 (0.4) | |
| 46-55 | 136 (27.2) | 1 (0.2) | |
| 56-65 | 150 (30.0) | 0 (0.0) | |
| 66-75 | 70 (14.0) | 0 (0.0) | |
| 75-above | 50 (10.0) | 0 (0.0) | |
| TOTAL | 500 (100.0) | 3 (0.6) | |

290 The risk factors associated with HCV, HIV and co-infection of HIV/HCV in TB
291 patient was based on patient self-report. Alcoholism, previous unprotected sex, multiple
292 sex partner, previous blood donation, Previous transfusion, Tattoos and History of Sexual
293 Transmitted disease are the risk factors. Out of 31 infected subjects, 14(45.2%) identified
294 with alcoholism, previous unprotected sex 11(35.5%), Multiple sex partner 10(32.3%),
295 Previous blood donation 8(25.8%), previous transfusion 7(22.6%), others are Tattoos
296 7(22.6%) and history of STD 3(9.68%). The risk factors were represented in pie chart
297 shows in Figure I.

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PATIENTS AS PREDISPOSING TO HIV AND HCV INFECTIONS



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307 **FIGURE I: PIE CHART OF RISK FACTORS WITH HCV AND HIV**

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310 **4.0 DISCUSSION, CONCLUSION AND RECOMMENDATION**

311 The HIV and HCV are both global public health problems. Infections with HIV and or
312 HCV may have a major impact on those with TB. **HIV is the most important risk factor
313 for the development of TB among person infected** with *M. tuberculosis* and both CDC
314 and WHO guidelines recommended offering HIV testing to those person diagnosed with
315 TB disease [2, 10]

316 However, the prevalence of HCV infection among persons with TB has been
317 poorly defined and few data are available from the most area around the world. One study
318 in US suggested that veterans with HCV infection are at risk for other selected infectious
319 disease including TB [2]. Of 500 (100%) samples collected and tested against HCV and
320 HIV antibodies, HIV antibodies were positive in 21(4.2%) which might be due to the fact

321 that HIV prevalence in Ekiti State is low and this agreed with previous work of USAIDS
322 [11]. This also agreed with the work of Idigbe *et al*, [12] that prevalence of HIV in TB in
323 Nigeria, Lagos to be specific is 5.3%. However, HCV antibodies were positive in
324 10(2.0%) patients which also agreed with previous work of Mwangi [13] that the
325 prevalence in Ekiti State and in Nigeria is low. This might be due to proper screening of
326 donor which is one of the major predisposing factors to increase in the incidence of HCV
327 [13]. However, the prevalence of HCV among TB has been poorly defined and few data
328 are available around the world. Part of the lack of data on HCV seroprevalence stem from
329 the fact that there are no recommendations for universal screening of TB patient for HCV
330 as there for HIV testing [2]. Although, Halim and Ajayi [14] reported 12.3%
331 seroprevalence of HCV in Nigeria among the donor and the findings from Richard *et al*,
332 [2] reveals that patient with TB may have among the highest prevalence of HCV
333 infection.

334 The prevalence of HCV and HIV antibodies together in TB patient is 3(0.6%) which
335 appears to be low but can pose a major threat to the management of TB patients and this
336 agreed with previous work of Richard *et al*, [2] who reported 0.4% prevalence rate of
337 HCV and HIV antibodies in TB patient in Georgia.

338 Gender wise distribution of seroprevalence of HCV and HIV in TB patients revealed that
339 although the number of males that participated is more than female but there was no
340 significant difference (0.531, 0.549, 0.824) between male and female for HCV, HIV, and
341 HCV/HIV co-infection respectively, which shows that HCV, HIV, and HCV/HIV can
342 infect any sex and this agreed with previous work of Richard *et al*, [2].

343 Age distribution revealed that age group 36-45 had the highest prevalence of
344 HCV, HIV and both HCV/HIV antibodies and this is statically significant (P value 0.000
345 for HCV, 0.000 for HIV and HCV/HIV is 0.002). This might be due to the fact that at this
346 age, subjects are sexually active and are involved in some of the risk factor(s) that
347 predispose them to the infections. This agreed with work of Watanabe *et al*, [15] which
348 revealed 25-45years as the most predispose age group to HIV. The predisposing risk
349 factors to seroprevalence of HCV, HIV and HCV/HIV antibodies in the study population,
350 showed in figure I. Alcoholism is the highest of the factors, follow by previous
351 unprotected sex, multiple sex partner, previous transfusion, tattoos and the least is a

352 history of sexually transmitted disease. Although HCV and HIV are blood borne disease,
353 the route of transmission is similar, nevertheless, a number of investigations have
354 indicated that acquisition of HCV through sexual contact is uncommon and have
355 suggested that HCV is inefficiently transferred through this mechanism [16, 17] despite
356 these findings, a number of studies had found that high-risk sexual behaviour or history
357 of STD are associated with an increased risk of HCV infection [16] and so, high-risk
358 sexual behaviours and /or a history of STI may be a maker for other risks that have been
359 implicated as mechanism of transmission of HCV. However, Richard *et al*, [2] revealed
360 that most common route of HCV transmission worldwide is through hematogeneous
361 transmission, tattoos and nevertheless, hematogeneous transmission may not be too
362 implicated in this study because in Ekiti state, the WHO guidelines of screening of blood
363 donor are followed strictly and this has contributed to the low prevalence of HCV, HIV in
364 this part of Nigeria [11].

365 **CONCLUSION**

366 Since, HCV and HIV co-infection in TB patients increased the risk of antituberculosis
367 drug-induced hepatotoxicity and that there is an even greater risk for drug-induced
368 hepatotoxicity among those undergoing treatment for TB who had both HCV and HIV
369 co-infection, to this end, more active screening for HCV should be done in this
370 population (TB) as was done for HIV. There is also a need to know the underlying health
371 status of TB patient as regards the HCV and HIV before administering drugs. Above all,
372 there is a need for a sample of TB patient to send to the laboratory for liver function test
373 because of the effects of the regimes on the liver.

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375 **RECOMMENDATION**

376 It is recommended that there should be a universal screening of person with TB for HCV
377 infection as there are for HIV testing.

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