




Effect of Isoniazid Prophylaxis Therapy on the Prevention of Tuberculosis Incidence and Associated Factors Among HIV Infected Individuals in Northwest Ethiopia: Retrospective Cohort Study

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Background: Treating latent tuberculosis (TB) infection with Isoniazid (INH) among human immune virus (HIV) infected patients reduces active TB occurrence and death by 62% and 26%, respectively. Even though other studies show aforementioned evidence, TB incidence and its associated factors among HIV-infected individuals who were on INH and never on INH is not well studied in northwest Ethiopia. Therefore, this study tried to assess the effect of INH prophylaxis in TB prevention and associated factors among HIV-infected individuals.

Methods: Data were extracted from charts of HIV-infected clients who completed INH (193) and were never on INH (198) after a simple random sampling selection was done among newly diagnosed patients on follow-up from 2008 to 2015. After data were collected, it was entered into Epi info version 7 and exported to SPSS version 22 for analysis. Cox regression model was fitted and the hazard ratio was reported.

Results: In this study, the overall TB incidence rate among HIV patients was 3.5/100 person-years (PY) [95% CI: 2.55, 4.82]. But it was 7.1/100 PY among patients who were never on INH and 0.35/100 PY among patients who completed INH. INH completed [adjusted hazard ratio (AHR) = 0.08, 95% CI: 0.02–0.37], on anti-retroviral therapy (ART) [AHR = 0.02, 95% CI: 0.01–0.04], baseline World Health Organization (WHO) stage I

Results: & II [AHR = 0.22, 95% CI: 0.08–0.62] and baseline CD4 ≤ 350 [AHR = 3.76, 95% CI: 1.39–10.18] were significantly associated with TB incidence.

Conclusion: Putting patients on INH for 6 months and ART were protective factors against TB. Therefore, health institutions are recommended to provide INH after ruling out active TB and contraindications for HIV-infected individuals.

Keywords: HIV, TB, incidence, isoniazid, prophylaxis, factors

Introduction

People living with HIV develop a variety of different opportunistic infections (OIs) in their life span. Tuberculosis is the most common life-threatening and leading cause of death accounting for approximately 25% of all causes of death among HIV patients.^{1,2}

In 2018, among all TB cases, about 8.6% were HIV infected; TB/HIV co-infection was highest in African countries; more than 50% in the southern part of Africa.³ In Ethiopia, in 2014/15, among 96% of newly enrolled Pre-ART patients

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screened for TB; 9.2% were identified with a TB co-infection.⁴ Moreover, the prevalence of Pulmonary TB among Pre-ART patients was 7.5% according to a study done in Gondar University Hospital.⁵

The risk of developing TB is 20–37 times more among HIV infected than non-HIV infected individuals.⁶ The annual and the lifetime risk of developing active TB in HIV infected and uninfected individuals is 10% respectively.⁷

The WHO as well as the Federal Ministry of Health of Ethiopia formulated and recommend using the three TB/HIV integrated activities to decrease TB/HIV co-infection. These activities are integrating and use of TB/HIV services delivery, use of “three I’s (Intensive TB case finding, Isoniazid preventive therapy (IPT) and Infection prevention at health Institutions)” and minimize the incidence of HIV among TB patients.⁸

INH can prevent both reactivation of latent TB infection to active TB and reinfection with mycobacterium TB bacilli.^{9,10} A retrospective study in public health facilities in Addis Ababa identified that providing INH prophylaxis to HIV infected individuals prevented the occurrence of TB by 96% compared to those who did not take it.¹¹

Although the WHO and the Joint United Nation Program recommend the use of INH prophylaxis at least for 6 months to prevent new TB incidence, the coverage remained very low according to global TB reports in Ethiopia. The IPT coverage was 5.4% among the overall enrolled patients in 2011, and it was more than two-fifths (45%) in 2018 among newly enrolled patients. This low coverage is attributable to weak active TB screening, low INH access, fear of INH resistance, and toxicities.^{12–15}

Before providing INH prophylaxis, every HIV-infected individual should be screened for active TB and contraindications should be excluded. This is because it is not advisable to give INH to HIV-infected individuals if patients have one of the symptoms (“Cough, fever, night sweats, weight loss symptoms”) unless TB is ruled out.^{8,16} Researches had shown that using INH prophylaxis for HIV infected individuals did not increase the risk of developing drug resistance TB.¹⁷

One of the drawbacks of providing INH is its side effects. In a study done in Gondar town, side effects that led to poor INH adherence were “abdominal pain, vomiting, skin rash, jaundice and numbness”.¹⁸

According to TB/HIV co-infection related studies, the most common risk factors that contributed to active TB incidence were INH prophylaxis, WHO clinical staging,

CD4 count, Body Mass Index (BMI), functional status, hemoglobin level, cotrimoxazole prophylaxis.^{11,19–25}

This study was aimed to assess the effect of INH prophylaxis in the prevention of TB incidence and its associated factors among HIV infected individuals. Since it was not well studied and INH implementation varies across different health institutions in northwest Ethiopia, this study will fill knowledge, practice/implementation, and research gaps in the area. Information obtained from this research will help policy makers, health managers, health care workers and other stakeholders. In addition, it will provide additional sources of information for interested researchers to further investigate TB/HIV co-infection and the effectiveness of INH prophylaxis against TB prevention among HIV infected patients.

Methods

Study Area and Design

A retrospective cohort study was conducted in Gondar and Azezo health centers in Gondar town. Gondar is located northwest of Ethiopia. It is 180 km from the capital city of the Amhara region, Bahir Dar and also 748 km from Addis Ababa the capital city of Ethiopia.

According to the Ethiopian population projection census of 2015, the total population in Gondar administrative town was 323,900.²⁶

In Gondar town, there are eight health centers, one referral hospital and 14 health posts.²⁷

The town has also one private general hospital, 14 specialized clinics, 18 medium clinics, 9 first level clinics, 1 basic laboratory, 1 international laboratory and 23 pharmacies.²⁷

Each health center is expected to serve 40,958 of the population. TB screening and treatment is given in all 8 health centers, in the referral hospital and in 12 private clinics. TB/HIV services are given in 20 health institutions. TB and HIV/AIDS services including screening and treatment of opportunistic infection, prevention of mother to child transmission (PMTCT), provision of INH, Cotrimoxazole, ART, anti-TB drugs, care and support for HIV-infected individuals are the major activities for both the health centers. Gondar and Azezo health centers provide health care services for more than 2000 and 754 HIV infected individuals respectively.²⁷

Study Participants

HIV infected adults (≥ 18 years) who completed INH (took INH for 6 months) and who never took the prophylaxis

from September 11, 2008 to March 8, 2015 in both Gondar and Azezo Health Centers were included in the study. These individuals were either on pre-ART or ART. All HIV infected adults who took INH prophylaxis partially, charts with incomplete data and individuals on anti-TB treatment at the beginning of the study were excluded.

Sample Size Estimation and Sampling Procedures

The minimum sample size, 391, was estimated for TB incidence using single population proportion formula (considering HIV infected patients as a single population group as compared to the non-HIV infected population) based on the study in Arba Minch using the following assumption: level of significance (α) = 5%, confidence level = 95%, a margin of error = 4%, incidence proportion of TB = 18%, and 10% attrition rate. Sample size calculation for factors was estimated using Epi info version 7 by taking three factors, INH status, sex, and educational status, with the effect size of 0.313, 0.299, and 0.425 respectively with power = 80%. The sample sizes using the three factors taken from the Arbaminch study²² were (INH = 147, sex = 151, and education = 289). The largest sample size obtained using the single population proportion formula (391) was taken as the sample size of this study. Among the 8 health centers in the town, the two health centers were selected since they were the only health centers that started HIV/AIDS care and treatment services between September 11, 2008, and March 8, 2015. In the beginning, they were checked for the presence of an adequate sample. Then, patients were stratified into two categories based on their INH status; INH completed and never on INH then a sample of patients were selected using simple random sampling (lottery method) from newly and previously enrolled patients from each stratum resulting in 193 INH completed and 198 never on INH patients.

After identifying patients who completed INH starting from September 11, 2008, to March 8, 2015, each individual was followed till either active TB diagnosis (bacteriological or clinical decision) or is censored.

Never on INH individuals were either newly (diagnosed between September 11, 2008 and March 8, 2015) or previously diagnosed (diagnosed before September 11, 2008) HIV infected patients for whom the date of entry was the date of entry into HIV care then followed till either TB diagnosis or censored.

Operational Definitions

Tuberculosis: Is diagnosed using bacteriological confirmation (smear microscopy) or clinical decision.

Clinical TB diagnosis: Is diagnosed when it did not fulfill bacteriological diagnosis (positive smear microscopy) ie diagnosed by clinicians based on either chest x-ray abnormality, histology suggestion (FNAC, biopsy) and failed to improve from repeated broad spectrum antibiotics with clinical suggestive of TB (not using anti-TB).

TB incidence: Occurrence of new TB cases during follow up time period

Separated: Couples who are officially married and living at different places.

INH prophylaxis completed: either Pre-ART or ART individuals taking INH prophylaxis for 6 months to sterilize latent TB infection.

Adherence: Good: $\geq 95\%$ or ≤ 3 doses missed per month.

Fair: (85–94)% or 4–8 doses missed per month.

Poor: $< 85\%$ or ≥ 9 doses missed per month.⁸

Functional status: Working: Able to perform usual work in and out of the house.

Ambulatory: Able to perform activities for daily living.

Bedridden: Unable to perform activities for daily living.⁸

Data Collection Instruments and Procedures

A well structured English version checklist was prepared containing Sociodemographic, clinical, immunological, therapeutic and behavioral factors and it was used to collect data in the charts from June 18 to June 30, 2018 in both ART clinics. The checklist was developed by reviewing related literatures. Charts were extracted by data clerks and the data were collected by the principal investigator.

Data Quality Measures

Data qualities were ensured using a well structured checklist. After selecting chart extractors (data clerks) short training was given on how to select charts randomly for the two groups.

A preliminary review (5%) was done in University of Gondar Referral Hospital to check whether the checklist included adequate information or not.

Data Management and Analysis

After the data were collected, the investigators checked their completeness manually. Then they were entered into

Epi info version 7 and exported to SPSS version 22 for analysis. Before the actual data analysis, data were checked for errors, outlying observations, inconsistencies, and missing observations.

Variables with continuous data were explored for their normality via histogram and Shapiro–Wilk statistics and none of the variables were normally distributed. Descriptive statistics were used to report the frequency, prevalence, median and interquartile range of the variables.

The life table was used to estimate the risk of TB occurrence among INH completed and never on INH individuals at different time intervals, and the Kaplan Meier estimate was used to compare the survival experience of the two groups. The Cox regression model was used to identify the independent variables that were associated with the time to TB incidence. Bivariable Cox regression analysis was done first for each variable then variables with P-value ≤ 0.2 in the bivariable were taken into the multivariable analysis to control the effect of confounders. Variables in the multivariable analysis were considered as significant at $P < 0.05$ and adjusted hazard ratios with 95% confidence level were reported to show the strength of the association.

The Schonfeld and Cox-Snell residual tests were used to check proportional hazard assumptions and goodness of fit of the model; the assumptions were satisfied.

Results

Baseline Sociodemographic Characteristics

About 450 charts were extracted from both health centers (Figure 1). Of the 391 HIV infected individuals included in the study, 193 completed INH and 198 never took INH.

The median age of the study participants was 32 [Inter quartile range (IQR): 27–40] years. The majority, 276 (70.6%) of study participants were females; and 337 (86.1%) were urban dwellers. Ninety-seven (24.8%) study participants were housewives, 89 (22.8%) were daily laborers and 39.9% were illiterates (Table 1).

Baseline Clinical Characteristics of Study Participants

The vast majority, 336 (85.9%) of the study participants were on ART and 158 (40.4%) were stage III patients. The median follow-up time of study participants was 3.001 years (IQR: 2.998–3.001) with minimum and maximum

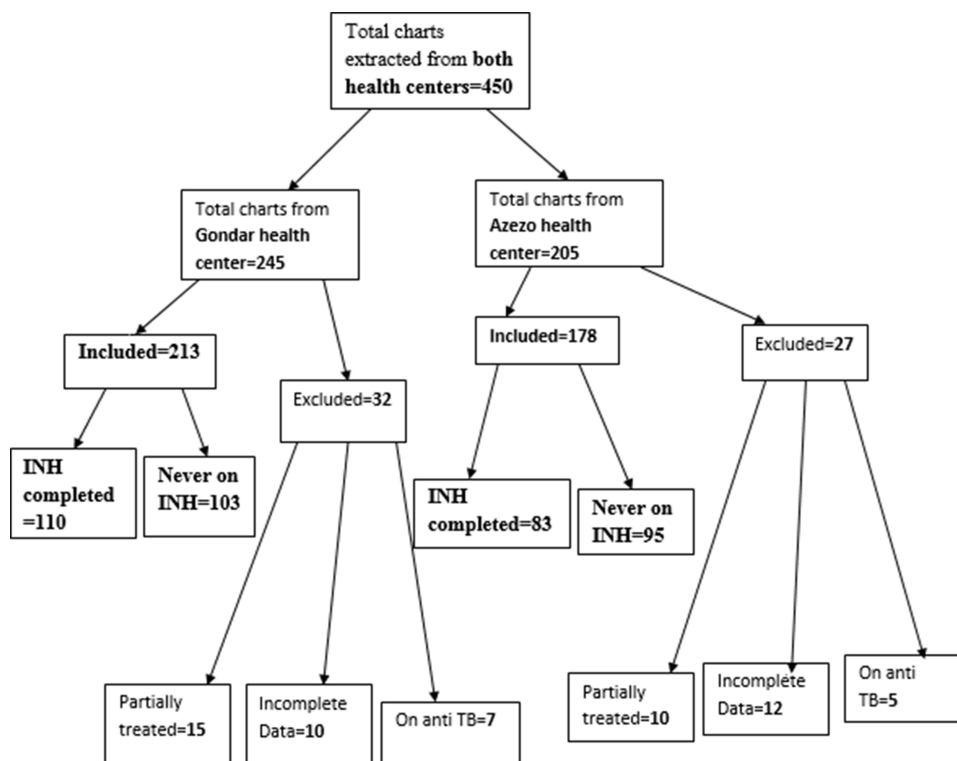


Figure 1 Sample size and sampling procedure (included and excluded participants of both health centers) in northwest Ethiopia from September 11, 2008 to March, 2015.

Table I Baseline Socio Demographic Characteristics with INH Status of Study Participants in Northwest Ethiopia from September 11, 2008 to March, 2015, N = 391

Variables	Sub Category	INH (%)	Never on INH (%)
Age	18–29	72(37.3)	73(36.9)
	30–39	78(40.4)	70(35.4)
	40–49	32(16.6)	31(15.7)
	≥50	11(5.7)	24(12.1)
Sex	Male	58(30.1)	57(28.8)
	Female	135(69.9)	141(71.2)
Residence	Rural	19(9.8)	35(17.7)
	Urban	174(90.2)	163(82.3)
Marital status	Married & Sepa	97(50.3)	98(49.5)
	Single	33(17.1)	27(13.6)
	Widowed	23(11.9)	23(11.9)
	Divorced	40(20.7)	50(25.3)
Educational st.	Illiterate	74(38.3)	82(41.4)
	Primary (1–8) grade	49(25.4)	58(29.3)
	Secondary (9–10)	59(30.6)	43(21.7)
	Tertiary & above (11 and above)	11(5.7)	15(7.6)
Occupational.	Gov & NGOs	14(7.3)	17(8.6)
	Self em. and Mer.	34(17.6)	51(25.8)
	Farmer	14(7.3)	25(12.6)
	House wife	48(24.9)	49(24.7)
	Jobless & Daily	83(43.0)	56(28.3)
Religion	Orthodox	177(91.7)	187(94.4)
	Protestant	1(0.5)	7(3.5)
	Muslim	15(7.8)	4(2.0)

Abbreviations: Gov, government; NGOs, no governmental organization; sepa, separated; mer, merchant; INH, isoniazid.

value of 0.01 and 3.001 years respectively while the median follow up time for INH completed individuals and those who did not take is similar (3.001). The median time of study participants since HIV diagnosis was 3.6 (IQR: 3.001–6.489) years. Of all study participants, 312 (79.8%) had working functional status. The median baseline CD4 count was 212 (IQR: 117–359) cells/ μ L. Among INH completed individuals 89.1% were on ART and 86.5% were on cotrimoxazole preventive therapy (CPT) (Table 2).

Incidence of TB Among HIV Infected Individuals

The overall incidence rate of TB among 391 study participants was 38 (3.5/100 PY) [95% CI: 2.55, 4.82] whereas, it was 0.35/100 PY (INH completed) and 7.1/100 PY (never on INH) individuals. Most TB incidences occurred

within the first year of follow-up time, in particular within the first 6 months.

According to results from Kaplan Meier and log rank tests, there was a significant difference in acquiring TB during the follow-up period between INH completed and never on INH groups (P-value <0.001). The estimated average TB free time among INH completed individuals was 2.99 years and 2.56 years among never on INH individuals (Figure 2).

TB Incidence and Factors Among HIV Infected Individuals

Sociodemographic and Behavioral Factors

TB incidence rate increased as the age of the study participants increased. TB was 29 (7.4%) among urban dwellers while it was 9 (2.3%) among rural dwellers. However, the incidence in rural areas among individuals who never

Table 2 Baseline Clinical and Immunological Characteristics with INH Status of Study Participants in Northwest Ethiopia from September 11, 2008 to March, 2015, N = 391

Clinical Variables	Sub Category	INH (%)	Never on INH (%)
WHO stage	Stage I	63(32.6)	58(29.3)
	Stage II	45(23.3)	49(24.7)
	Stage III	80(41.5)	78(39.4)
	Stage IV	5(2.6)	13(6.6)
Previous TB rx.	Yes	23(11.9)	22(11.1)
	No	170(88.1)	176(88.9)
B. functional stat.	Working	155(80.3)	157(79.3)
	Ambula.&B	38(19.7)	41(20.7)
Baseline BMI	<18.5 kg/m ²	64(33.2)	87(43.9)
	≥18.5 kg/m ²	129(66.8)	111(56.1)
Baseline Hgb.	<10 mg/dl	5(2.6)	9(4.5)
	10–12 mg/dl	26(13.5)	43(21.7)
	>12 mg/dl	162(83.9)	146(73.7)
Baseline CD4	≤100 cells/μL	35(18.1)	46(23.2)
	101–200 cells/μL	49(25.4)	57(28.8)
	201–350 cells/μL	50(25.9)	47(23.7)
	>350 cells/μL	59(30.6)	48(24.2)
CD4 at ART initiat.	≤100 cells/μL	35(20.3)	37(22.6)
	101–200 cells/μL	68(39.5)	61(37.2)
	201–350 cells/μL	51(29.7)	39(23.8)
	>350 cells/μL	18(10.5)	27(16.5)
ART status	On ART	172(89.1)	164(82.8)
	On Pre-ART	21(10.9)	34(17.2)
Past OIs than TB	Yes	80(41.5)	96(48.5)
	No	113(58.5)	102(51.5)
CPT initiation	Yes	167(86.5)	169(85.4)
	No	26(13.5)	29(14.6)

Abbreviations: ART initiat, ART initiation; previous TB rx, previous TB treatment history; functional stat, functional status.

took INH was 9 (25.7%) as opposed to 27 (16.6%) urban dwellers (Table 3). TB incidence rate among farmers was 8.47/100 PY [95% CI: 4.24, 16.94] and smokers had a nearly similar TB incidence rate (3.56/100 PY) as compared to nonsmokers (3.50/100 PY). Alcohol drinkers had a TB incident rate of 2.49/100 PY while non-drinkers had 3.97/100 PY.

Clinical and Immunological (Laboratory) Factors for Active TB Incidence

Out of the total study participants, 15 (19%) individuals with ambulatory and bedridden functional status had TB. However, it was 23 (7.4%) in those with working

functional status. Twenty eight (82.4%) of Pre-ART participants who did not take INH developed TB (Table 4).

TB incidence rate among individuals who never took INH was higher 7.1/100 PY [95% CI: 5.12, 9.84] as compared to the INH completed, 0.35/100 PY. Individuals who started INH before ART had higher TB incidence rate 1.03/100 PY [95% CI: 0.26, 4.14] than those who started after ART with IR of 0/100 PY. TB incidence rate among individuals who were treated for TB previously was 4.94/100 PY and those individuals who were not treated for TB was 3.32/100 PY. As duration of HIV diagnosis increased, TB incidence decreased. TB incidence rate was higher 0.86/100 PY among individuals

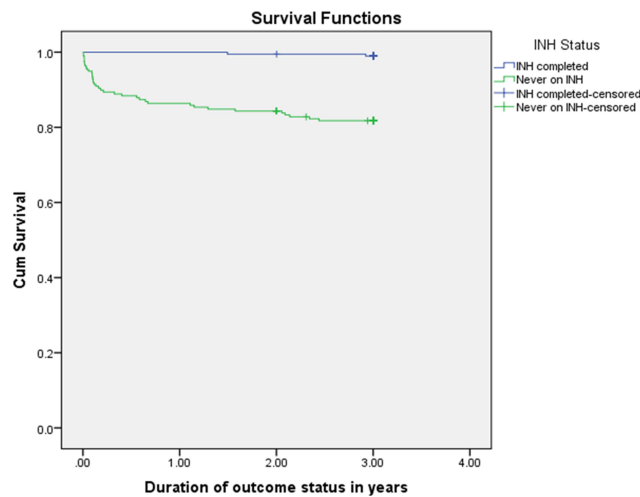


Figure 2 Survival function curve both for INH completed and never on INH groups in northwest Ethiopia from September 11, 2008 to March, 2015.

with good ART adherence and TB case was zero with fair and poor ART adherence. Interestingly, the incidence rate among individuals with no INH or ART was 86.96/100 PY (Table 5 and Figure 3).

Cox Regression Survival Analysis

In bivariable analysis, sex, INH status, ART status, baseline WHO stage, baseline BMI, baseline functional status and duration of HIV diagnosis were significantly associated with TB incidence.

In multivariable analysis, INH, ART status, WHO stage and baseline CD4 were significantly associated with TB incidence. Individuals who completed INH were protected from TB as compared to individuals who were never on INH [AHR = 0.08, 95% CI: 0.02, 0.37]. On ART individuals were at lower risk of developing TB as compared to pre-ART individuals [AHR = 0.02, 95% CI: 0.01, 0.04]. Stage I and II individuals were at lower risk of developing TB than stage III and IV individuals [AHR = 0.22, 95% CI: 0.08, 0.62]. The hazard of developing TB among patients with baseline CD4 count ≤ 350 was significantly higher than those with CD4 > 350 [AHR = 3.76, 95% CI: 1.39, 10.18] (Table 5).

Discussion

In this study, we determined the incidence of tuberculosis and associated factors among HIV patients who completed INH therapy and those who never took INH. The overall incidence was 3.5/100 PY; factors independently associated with it were INH, ART status, WHO stage, and baseline CD4.

Table 3 Socio Demographic and Behavioral Factors with Corresponding INH Status and Cumulative TB Incidence in HIV Infected Individuals in Northwest Ethiopia from September 11, 2008 to March 8, 2015, N = 391

Socio Demo	INH Completed		Never on INH	
	TB (%)	Total	TB (%)	Total
Age				
18–29	0(0.0)	72.0	11(15.1)	73.0
30–39	2(2.6)	78.0	12(17.1)	70.0
40–49	0(0.0)	32.0	7(22.6)	31.0
≥50	0(0.0)	11.0	6(25.0)	24.0
Sex				
Male	1(1.7)	58.0	17(29.8)	57.0
Female	1(0.7)	135	19(13.5)	141.0
Residence				
Rural	0(0.0)	19.0	9(25.7)	35.0
Urban	2(1.1)	174	27(16.6)	163
Marital stat.				
Married & Se.	2(2.1)	97.0	14(14.3)	98.0
Single	0(0.0)	33.0	9(33.3)	27.0
Widowed	0(0.0)	23.0	6(26.1)	23.0
Divorced	0(0.0)	40.0	7(14.0)	50.0
Education				
Illiterate	1(1.4)	75.0	12(14.6)	94.0
Primary	1(2.0)	50.0	8(13.8)	66.0
Secondary	0(0.0)	59.0	14(32.6)	57.0
Tertiary & ab	0(0.0)	11.0	2(13.3)	17.0
Occupation				
Gov & NGOs	0(0.0)	14.0	6(35.3)	17.0
Self e. & m	2(5.9)	34.0	11(21.6)	51.0
14.0	8	8	Farmer (32.0)	0(0.0)
House wife	0(0.0)	48.0	5(10.2)	49.0
Jobless & Da.	0(0.0)	48.0	6(10.7)	56.0
Religion				
Orthodox	2(1.1)	177	33(17.6)	187
Protestant	0(0.0)	1.00	1(14.3)	7.00
Muslim	0(0.0)	15.0	2(50.0)	4.00
Cigarette sm.				
Yes	1(6.3)	16.0	2(14.3)	14.00
No	1(0.6)	177	34(18.5)	184
Alcohol drink				
Yes	1(1.4)	69.0	1(0.8)	124
No	7(14.9)	47.0	29(19.2)	151

(Continued)

Table 3 (Continued).

Socio Demo	INH Completed		Never on INH	
	TB (%)	Total	TB (%)	Total
Khat chewing				
Yes	0(0.0)	14.0	3(25.0)	12.0
No	2(1.1)	179	33(17.7)	186

Abbreviations: Socio demo, socio demographic; Se., separated; Gov, governmental; NGOs, nongovernmental; self e & m, self employee & merchant; Da., daily laborer; ab, above; sm, smoking.

Overall TB incidence was relatively lower as compared to findings in other studies.

The TB incidence rate increased as the age of the study participants increased which was in line with the studies done in SNNPR, Cambodia and Iran in which the incidence rate of TB increased as age increased.^{23,28,29} This would be due to the decline in immune status as age increased, age related comorbidities and poor standard of living conditions.

Individuals who lived in rural areas (6.6 PY) had two times the TB incidence rate of urban dwellers (3.03 PY). This finding was in line with a study done in South Africa and in contrast to a study done in the Amhara region in which TB burden was higher among urban dwellers.^{24,25} This discrepancy might be due to the immune status of study participants, living environment, and standard of living conditions.

In our study, smokers had nearly similar TB incidence rate (3.56/100 PY) as compared to non smokers (3.50/100 PY). This finding was against to studies done in Amhara region, Jimma and Mettu Karl hospital, Addis Ababa and Cambodia in which smokers were at higher risk of TB.^{19,23,24,30} This might be due to differences in the amount of cigarettes smoked/individual/day, duration of smoking, ART status, living standards and immune status.

About 78% of study participants with baseline WHO stage I and II were protected from developing TB as compared stage III and IV. This was comparable with studies done in Addis Ababa and JUSH that showed stage III and IV individuals were at a higher risk of developing active TB.^{19,20} This might be due to the presence of the better immunity among stage I and II individuals that prevent mycobacterium TB infection and reactivation of latent TB infection into active TB.

Individuals with baseline CD4 count \leq 350 were 3.76 times at higher risk of TB as compared to those

Table 4 Clinical and Immunological Factors with INH Status and Cumulative TB Incidence in HIV Infected Individuals in Northwest Ethiopia, September 11, 2008 to March, 2015, N = 391

Clinical Variables	INH Completed		Never on INH	
	TB (%)	Total	TB (%)	Total
WHO stage				
Stage I	1(1.6)	63.0	2(3.4)	58.0
Stage II	0(0.0)	45.0	2(4.1)	47.0
Stage III	1(1.2)	74.0	23(29.5)	78.0
Stage IV	0(0.0)	5.00	9(69.2)	13.0
Previous TB rx				
Yes	0(0.0)	23.0	6(27.3)	22.0
No	2(1.2)	170	30(17.0)	176
B. functional sta.				
Working	2(1.3)	155	21(13.4)	157
Ambulatory & Bed.	0(0.0)	38.0	15(36.6)	41.0
Baseline BMI				
<18.5 kg/m ²	1(1.6)	64.0	22(25.3)	87.0
\geq 18.5 kg/m ²	1(0.8)	129	14(12.6)	111
Baseline Hgb.				
<10 mg/dl	0(0.0)	5.00	1(11.1)	9.00
10–12 mg/dl	0(0.0)	26.0	21(25.6)	53.0
>12 mg/dl	2(1.2)	162	24(16.4)	146
Baseline CD4				
\leq 100 cells/ μ L	0(0.0)	35.0	14(30.4)	56.0
101–200 cells/ μ L	0(0.0)	49.0	6(10.5)	57.0
201–350 cells/ μ L	0(0.0)	50.0	11(23.4)	47.0
>350 cells/ μ L	2(3.4)	59.0	5(10.4)	48.0
CD4 at ART in.				
\leq 100 cells/ μ L	0(0.0)	35.0	3(8.1)	37.0
101–200 cells/ μ L	0(0.0)	68.0	1(1.6)	61.0
201–350 cells/ μ L	0(0.0)	51.0	2(5.1)	39.0
>350 cells/ μ L	0(0.0)	18.0	2(7.4)	27.0
ART status				
On ART	0(0.0)	172	8(4.9)	164
On Pre-ART	2(9.5)	21.0	28(82.4)	34.0
Duration of ART				
< 2 years	0(0.0)	27.0	6(18.8)	32.0
2–3 years	0(0.0)	18.0	1(0.8)	118
>3 years	0(0.0)	127	1(7.1)	14.0
Past OIs other than TB				
Yes	0(0.0)	80.0	19(19.8)	96.0

(Continued)

Table 4 (Continued).

Clinical Variables	INH Completed		Never on INH	
	TB (%)	Total	TB (%)	Total
No	2(1.8)	113	17(16.7)	102
CPT initiation				
Yes	0(0.0)	167	30(17.8)	169
No	2(7.7)	25.0	6(20.7)	29.0

Abbreviations: TB rx hx, tuberculosis treatment history; sta, status; BMI, body mass index; Hgb, hemoglobin; CPT, cotrimoxazole preventive therapy; OIs, opportunistic infections; μ L, micro liters; mg/dl, milligram per deciliter.

with $CD4 > 350$. Comparable findings were obtained in studies done in Harrar which was 62% higher in those with $CD4 \leq 350$; in the Amhara region, it was 1.71 times at higher risk among those with $CD4 < 200$; in South Africa individuals with $CD4$ count < 100 cells/ μ L had 2.22 times higher risk compared to $CD4 > 349$.^{24,25,31} Although TB can occur at any $CD4$ count, individuals with advanced WHO stage and severe immune suppression are at increased risk of conversion of latent TB infection to active TB; this might be the causes for these similarities.

The overall incidence rate of TB for the cohort was found to be 3.5/100 PY whereas the incidence of TB among INH completed individuals was 0.35/100 PY and 7.1 in those who never took INH. These findings were nearly similar with studies done in SNNPR overall TB (2.6/100 PY), among INH (0.7/100 PY) and for those who did not take INH (6.1/100 PY); in another study done in public health facilities of Addis Ababa, overall TB was 6.7/100 PY, INH group was 0.21/100 PY and among never on INH groups 7.18/100PY.^{11,28}

INH completed individuals were 92% at lower risk of developing TB compared to those never on INH. In line with this study, INH completed individuals were 96%, 85% and 68% at lower risk of developing TB according to studies conducted in public health facilities of Addis Ababa, SNNPR and Arba Minch Hospital respectively.^{11,22,28} This increased INH effectiveness might be due to synergistic effect with ART in the prevention of TB, good INH adherence, being on CPT and better immune status.

On ART patients were 98% at lower risk of developing TB disease compared to the Pre-ART counterparts. In support of this study, a study in Jimma and Mettu Karl hospital, not on ART participants (pre-ART) were 3.1

times more at higher risk compared to being on ART; the risk was also reduced by half in the study conducted in Harrar.^{30,31}

There was no TB incident case among individuals who were on both INH and ART. Similarly, a study in South Africa showed TB incidence was reduced by 89% among these groups of people. Moreover, a study done in SNNPR, TB incidence was reduced only by 57% among these groups of individuals.^{25,28} This might be due to the synergistic effect of INH and ART, being on CPT, good INH adherence and good clinical care.

As the duration of HIV diagnosis increased, the incidence rate of TB was decreased. In contrast to this finding, in Guangxi, China individuals with a long duration of HIV diagnosis were at higher risk of TB.³² This might be due to the inclusion of nearly half of the participants from INH completed, ART status, and having good medication adherence.

Strength and Limitations of This Study

It will be used as a baseline source of information for researchers who are interested to further investigate TB/HIV co-infection and the effectiveness of INH against active TB incidence among different HIV infected individuals and health institutions.

Being a retrospective study, variables such as viral load, diabetes mellitus, and other chronic diseases were missed from the charts during data collection these variables were significantly associated with TB incidence in related studies. We were also unable to determine the amount of alcohol, cigarettes, and khat being consumed. Since we excluded partially treated individuals we might have resulted in an overestimation of INH efficacy despite their numbers being small, 25.

Conclusions and Recommendations

According to this study, INH status, Baseline $CD4$ count, baseline WHO stage and ART status were significantly associated with TB incidence. Provision of INH for 6 months was significantly associated with a reduction of TB incidence. Individuals who were never on INH had 20 times higher TB incidence rate (7.1/100 PY) than INH completed individuals (0.35/100 PY). Individuals who started INH before ART had higher TB incidence rate than those started after ART. Individuals who were both INH completed and on ART were 100% protected from

Table 5 Bi Variable and Multivariable Cox Hazard Regression Analysis for Socio Demographic, Clinical and Immunological Factors with TB Incidence Among HIV Infected Individuals in Northwest Ethiopia from September 11, 2008 to March, 2015, N = 391

Variables	Outcome Status				
	TB (%)	Total	IR/100PY	CHR (95% CI)	AHR (95% CI)
Age					
18–29	11(7.6)	145	2.67	0.41[0.15,1.11]	1.41[0.43,4.64]
30–39	14(9.5)	148	3.39	0.52[0.20,1.35]	1.83[0.66,5.09]
40–49	7(11.1)	63.0	4.08	0.62[0.21,1.80]	1.38[0.43,4.44]
≥50	6(17.1)	35.0	6.81		
Sex					
Male	18(15.7)	115	6.01	2.29[1.21,4.33]*	1.98[0.97,4.07]
Female	20(7.2)	276	2.55		
Residence					
Rural	9(16.7)	54.0	6.60	2.10[0.99,4.44]	1.02[0.38,2.70]
Urban	29(8.6)	337	3.06		
INH status					
Completed	2(1.0)	193	0.35	0.05[0.01,0.21]*	0.08[0.02,0.37]*
Never on INH	36(18.2)	198	7.10		
ART status					
On ART	8(2.4)	336	0.81	0.03[0.01,0.07]*	0.02[0.01,0.04]*
Pre-ART	30(54.5)	55.0	32.04		
B. WHO stage					
Stage I and II	5(2.3)	215	0.78	0.11[0.04,0.29]*	0.22[0.08,0.62]*
Stage III and IV	33(18.8)	176	7.42		
B. Functional					
Ambula. & Be.	15(19.0)	78.0	7.61	2.79[1.46,5.35]*	0.95[0.43,2.10]
Working	23(7.4)	312	2.59		
Baseline BMI					
<18.5 kg/m ²	23(15.2)	151	5.79	2.56[1.34,4.91]*	1.90[0.24,15.21]
≥18.5 kg/m ²	15(6.3)	238	2.18		
BMI at start S.					
<18.5 kg/m ²	22(19)	116	7.52	3.53[1.86,6.73]*	0.73[0.36,1.48]
≥18.5 kg/m ²	16(5.8)	275	2.02		
Baseline CD4					
≤350 cells/μL	31(10.9)	284	4.02	1.76[0.78,7.24]	3.76[1.39,10.18]*
>350 cells/μL	7(6.5)	107	2.24		
Duration of HI.					
<5 years	37(14.9)	249	5.60	22.8[3.12,165.9]*	1.82[0.16,20.75]
≥5 years	1(0.7)	142	0.24		

Notes: *Statistically significant at P-value≤0.05 or one not included in CI, confidence interval; inf, infection; B. hemoglobin, baseline hemoglobin; B. functional status, baseline functional status; CHR, crude hazard ratio; AHR, adjusted hazard ratio; IR, incidence rate; PY, person years; ab, above; Dx, diagnosis; o & b, only and both; duration of HI, duration of HIV diagnosis, ART initi- initiation.

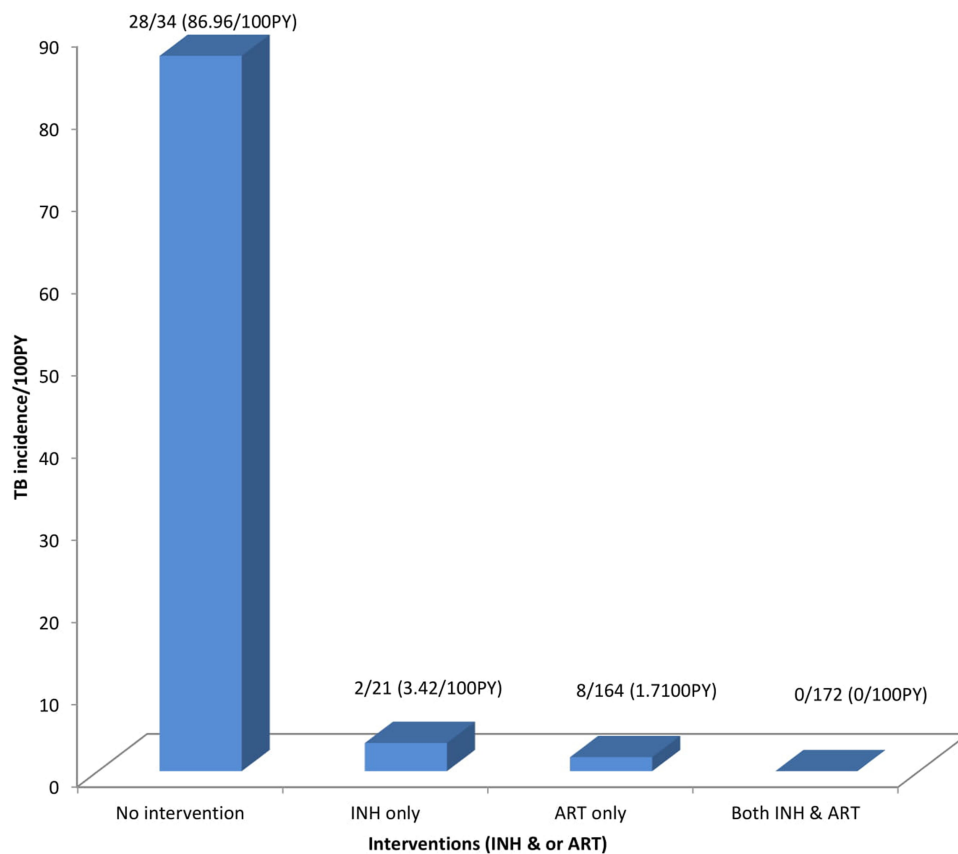


Figure 3 TB Incidence rate with corresponding INH and or ART intervention status in northwest Ethiopia from September 11, 2008 to March, 2015.

new TB incidence. However, ART alone was more protective than INH alone. As individuals lived longer and being on ART for a long time, TB incidence was decreased. Therefore, it is recommended to strengthen the provision of INH prophylaxis to all individuals in all health institutions with HIV/AIDS care and treatment centers after screening active TB and excluding contra indication.

Abbreviations

AIDS, acquired immune deficiency syndrome; ART, anti retro viral therapy; BMI, body mass index; CD4, cluster of differentiation; CPT, cotrimoxazole preventive therapy; HIV, human immune deficiency virus; HURH, Hawassa University Referral Hospital; INH, isoniazid; IPT, isoniazid preventive therapy; IQR, inter quartile range; IR, incidence rate; JUSH, Jimma University Specialized Hospital; OIs, opportunistic infections; PMTCT, prevention of mother to child transmission; PY, person year; TB, tuberculosis; UOGRH, University of Gondar Referral Hospital; WHO, World Health Organization; CI, confidence

interval; SNNPR, southern nations nationalities and peoples region.

Data Sharing Statement

All relevant information is included within the manuscript. The data upon which the result was based can be available upon request.

Ethics Approval and Consent to Participate

This research was conducted after ethical clearance was obtained from the Ethical Review Board of School of Medicine, University of Gondar. The Department of Internal Medicine wrote an official letter to the Ethical Review Board of School of Medicine. After ethical approval, the Ethical Review Board of School of Medicine wrote official letter to the town health bureau then the health bureau to both health centers. As this study was a retrospective cohort, verbal consent was requested from and approved by Heads of Health Centers to review and use medical records of the study participants. In addition, confidentiality of study

participants was maintained by avoiding personal identifiers during data collection and all check lists were kept locked.

The study was conducted in accordance with the Declaration of Helsinki 1964 and its consecutive amendments.

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Author Contributions

All authors made a significant contribution to the work reported; in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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