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Economic burden of Tuberculosis among Bangladeshi population and Economic Evaluation of the Current Approaches of Tuberculosis Control in Bangladesh

by

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DEDICATION

I dedicate this dissertation to the Tuberculosis patients who participated in the interviews for this study- not for money, just for the betterment of future TB care in the country.



ACKNOWLEDGEMENTS

I am truly grateful for the continuous support and inspiration from my supervisor Dr. M. Mahmud Khan. Without his guidance, supervision, and mentorship, it would not be a reality. I am thankful to Dr. Zaina P. Qureshi for her tremendous support throughout my graduate studies here at the University of South Carolina (USC). I would also like to thank the two other members of my dissertation committee, Dr. James W. Hardin and Dr. Abdul Hamid Salim.

My dissertation work was funded by the post-graduate training grant from TDR, the Special Programme for Research and Training in Tropical Diseases at the World Health Organization (WHO), I am thankful to them. Primary data was collected from all over Bangladesh for this dissertation. I cannot express enough gratitude to the TB patients and health professionals who were interviewed for this dissertation. I also offer my sincere gratitude to the data collectors for their tremendous effort. I would like to thank Dr. Mojibur Rahman (National Tuberculosis Control Program), Dr. Aung Kya Jai Maug (Damien Foundation), Dr. Shayla Islam (BRAC) for helping me with data collection process. Dr. Shahriar Ahmed (icddr,b) indebted me with his tremendous support throughout my research activities. I would like to thank Development Research Initiative (dRi) for helping me out with data collection process. I would also like to thank my co-researchers, Dr. Shakil Ahmed, Dr. Ibrahim Demir, Farahnaz Islam, Dr. Mohammad Masudur Rahman, and Khairul Alam Siddiqi for their continuous support.



Last but not least I would like to remember my family, especially my father whose relentless inspiration helped me to be what I am today, my mother for her monumental support, my siblings and my friends who were always there for me.

ABSTRACT

Introduction: Tuberculosis (TB) is major scourge for human history and causes profound economic burden. Bangladesh is a high burden TB country with 12% of its annual death is caused and 362 thousand people are infected by TB. DS-TB is the most prominent type of TB found in Bangladesh and a 6 month drug regimen (2 month intensive and 4 month continuation phase) is followed. But the directly observed treatment short-course (DOTS) differ in delivery through community health workers (CHW) and community members (CM). Bangladesh has also experienced surge in the number of MDR-TB cases with a 29% of MDR-TB cases were found among the retreatment of pulmonary TB cases in 2015. In Bangladesh, two MDR-TB treatment regimens (9 month and 20-24 month) are practiced. This dissertation aims to estimates the economic burden of TB on the afflicted Bangladeshi population and conducts economic evaluation among different programs for DS-TB and MDR-TB in Bangladesh.

Methods: This study collects direct and indirect cost for TB care data from 1,000 drug sensitive TB (DS-TB) and 145 multi-drug resistant (MDR-TB) patients from all over Bangladesh. Provider cost for TB care was also collected from the health facilities.

Costs for DS-TB and MDR-TB patients were estimated using Generalized Linear Model and summed up with per patient provider level costs to get the total costs per TB patients.

The incremental cost-effectiveness ratio (ICER) of treating DS-TB and MDR-TB patients, CM versus CHW model for DS-TB and 9-month vs. 20-24 month regimen for



MDR-TB were compared using a Markov model with life-time horizon. The measure of effectiveness, Quality adjusted life year (QALY) and cost of treatment was collected from 1,000 DS-TB and 145 MDR-TB patients (598 for CM model and 402 from CHW model; 58 undergone 9 month treatment and 87 from 20-24 month regimen) in Bangladesh. Transition probabilities between Markov states were estimated from quarterly outcomes report collected from health facilities and cost and QALY both were discounted at a rate of 3%. Both deterministic and probabilistic sensitivity analyses were conducted in a Monte Carlo Simulation using R.

Results: Mean age of DS-TB patients under the study was 45.2 years while mean age of MDR-TB patients were 35.5 years. In aggregate, DS-TB patients incurred total average costs of BDT 21,235 (USD 265) for TB illness; while MDR-TB patients' average costs were BDT 34,975 (USD 437). Including provider costs for each patient (USD 9 for DSTB and USD 2,006 for MDR-TB patients) total average costs for each DS-TB patient was BDT 22,003 (USD 275) and for each MDR-TB patient was BDT 1,95,449 (USD 2443).

Assuming 57% case notification rate, the actual costs for treating TB patients in 2015 was USD 55.6 million. If all DS-TB patients were treated the cost would have been 1 billion USD. For MDR-TB treatment, total cost was USD 12.5 million; treating all MDR-TB patients would have costed USD 23 million.

Results show that each DS-TB patient under CM treatment model gains 3.61 QALYs with a cost of BDT 131,555. For the DS-TB patients under the CHW model the cost is 81,650 and the QALY gain is 3.12. The Incremental Cost-Effectiveness Ratio



(ICER) is 103,454, i.e., the CM model is cost-effective if per QALY gain if willingness-to-pay is set to the per capita GDP of Bangladesh (BDT 107,360 in 2015).

Based on the study data, each patient under 9 month regimen gained 6.21 QALY with a total cost of BDT 987,418. Whereas, each patient under CHW model gained 5.74 QALY by incurring costs of BDT 1,501,221. Therefore, 9 month regimen is clearly dominating over the 20-24 month regimen because it costs less while it gains more QALY.

Conclusions: Results show that DS-TB patients incurred about 50% of their household annual income for treatment while that goes up to 66% for the MDR-TB patients. Pre-diagnosis cost constitutes about 63% of total costs for DS-TB patients and 42% of MDR-TB patient costs. This figures show the significant economic burden posed by TB and early diagnosis of the disease can reduce the burden in great extent.

Our study results demonstrate that community based model of DS-TB treatment is cost-effective even with changed costs and utility values in probabilistic sensitivity analysis. Community members as DOTS provider are more capable of reducing stigma related to TB, enhancing patient adherence and thereby reduce costs and increase utility from the treatment. Community members should also be involved in contact tracing and prevention activities to increase the effect of the involvement in TB control.

Our study results also suggest that shorter regimen remains cost-effective in Bangladesh setting with changing costs and utility parameters changed in the probabilistic sensitivity analysis. MDR-TB treatment is itself cost-effective in developed



countries and with cost-effective shorter regimen both treatment adherence and efficacy of the treatment will be improved.



PREFACE

American Psychological Association, 6th edition was used in the dissertation.

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LIST OF ABBREVIATIONS

AIDS
APHA
CEAC
CHW
DALY Disability Adjusted Life Year
DF
DOTs
DS-TB
ERC Ethical Review Committee
EVPI Expected Value of Perfect Information
GFATMGlobal Fund for Tuberculosis and Malaria
HCM Human Capital Method
HIV
IHEAInternational Health Economics Association
ICER
IRBInstitutional Review Board
HRQoLHealth Related Quality of Life
LTBI Latent Tuberculosis Infection
LMICLower Middle-Income Countries
MCMC
MDR-TB

MOHFW Ministry of Health and Family Welfare
NGO
NIDCHNational Institute of Diseases of the Chest and Hospital
NTP
OOPOut-Of-Pocket Payment
PMDTProgrammatic Management of Drug-resistant TB
PPP
QALYQuality Adjusted Life Year
TBTuberculosis
TDR The Special Programme for Research and Training in Tropical Diseases
UHC
UNDP
UNICEF
VIF
WHO
WTP
XDR-TB Extensively Drug Resistant Tuberculosis

CHAPTER 1

INTRODUCTION

1.1 STATEMENT OF THE PROBLEM AND RATIONALE

In 2015, Tuberculosis (TB) ranked 18th among the highest burden diseases globally and it constituted 47% of the global burden attributable to communicable, maternal, neonatal, and nutritional disorders (Kassebaum et al., 2016). In 2015, 10 million new cases of TB were reported and almost two million people died from TB worldwide (World Health Organization (WHO), 2016a). In 2015, TB became the top infectious disease killer by claiming With the me1.1 million lives by matching the death tolls by HIV/AIDS (Kassebaum et al., 2016).

Almost 85% of all new cases of TB and multi-drug resistant TB (MDR-TB) occur in 30 high burden TB countries and Bangladesh is one them (World Health Organization (WHO), 2015b). In 2015, 362,000 Bangladeshis developed TB and 73,000 died from it. TB accounted for 12% of all deaths (609,800) that occurred in 2015 in Bangladesh (Institute for Health Metrics and Evaluation (IHME), 2016). Although case notification rate is only 57%, success of the treatment is high (93%) among DS-TB patients. However, the success rate is 75% among MDR-TB patients which signifies how difficult to treat drug resistant strains (World Health Organization (WHO), 2016a).



Despite having effective treatment, patient adherence to TB treatment remains poor because of long duration of the regimen (six months for newly diagnosed cases) and the need for daily dosing. Failure to adhere to the regimen results in MDR-TB (Gandy & Zumla, 2002). The emergence of drug resistant TB strains has slowed down the progress in global TB epidemic control over the last two decades. Bangladesh has also experienced surge in the number of MDR-TB cases with a 1.6% of new cases are drug resistant and 29% of MDR-TB cases were found among the re-treatment of pulmonary TB cases in 2015 (World Health Organization (WHO), 2016a).

World has experienced a slow progress in TB control. TB incidence has fallen by an average of 1.5% per year since 2000. However, this needs to accelerate to 4-5% annual reduction to reach 2035 milestones of "End TB Strategy" (World Health Organization (WHO), 2016b). End TB Strategy has set ambitious targets of 95% reduction in TB deaths and 90% reductions in TB incidence by 2035 (Uplekar et al., 2015). Bangladesh is also experiencing a slow reduction in TB incidence (360,000 in 2015 from 362,000 in 2014) and incidence rate (225 per 100,000 population in 2015 from 227 per 100,000 population in 2014) (World Health Organization (WHO), 2015a, 2016a).

Economic burden of TB in Bangladesh is a great concern, since it affects a sizable number of people each year and causes 12% of the total death. Both disability and death have grave economic implications in the form of lost income to the persons and their families and lost Gross Domestic Product (GDP) for the country. Expensive treatment of the disease also put burden on the patients, families, and the health system of the country. The most affected group is the working age group persons, that also increases the costs associated with the disease (World Health Organization (WHO), 2016b). Moreover,



almost half (43%) of the patients in Bangladesh is not reported under the national registries and goes untreated (World Health Organization (WHO), 2016a); this makes the control and elimination of the disease extremely hard and expensive. Emergence of drug resistant strain also contributes in escalating costs because of high death rates, costly treatments, and poor outcomes (Fitzpatrick & Floyd, 2012).

Directly Observed Treatment, Short Course (DOTS) strategy for treating drug sensitive TB (DS-TB) has been implemented in Bangladesh since 1993 and all the Upazila Health Complexes (UHCs) have been brought under the purview of the service from where TB detection and treatment services are given free of cost. The essence of the strategy is the diagnosed TB patient has to go to the facility every day for taking the drugs, thus treatment discontinuity and subsequently MDR-TB cases can be averted (World Health Organization (WHO), 2013b).

Menacing drug resistant strains is a growing concern as discontinuity in treatment often results into MDR-TB or XDR-TB. Treatment for both the conditions are costly and also more time consuming as the shortest effective MDR-TB treatment regimen spans over nine months (Deun et al., 2010). Therefore, this prolonged treatment schedule may result into more incidence of treatment discontinuation. WHO in partnership with STOP TB Partnership came up with the response plan in 2007-2008 and Bangladesh is one of the seven countries using the shorter treatment regimens for MDR-TB in June 2013 (World Health Organization (WHO), 2013b).

Given the extent of the incidence and death rate of TB patients worldwide and in Bangladesh, its overwhelming economic impact is of great importance. Few studies have



been conducted to elucidate the social and economic costs of TB (Murrat, Styblo, & Roullion, 1993). Recently few studies have been carried out in this regard (Muniyandi, Ramachandran, Balasubramanian, & Narayanan, 2006; Rajeswari et al., 1999; Russell, 2004) but very few in the context of Bangladesh (Croft & Croft, 1998; Gospodarevskaya et al., 2014; Islam, Wakai, Ishikawa, Chowdhury, & Vaughan, 2002). There is also paucity of evidence on cost-effectiveness of the TB programs. One study compared cost-effectiveness between an NGO and government intervention and found that NGO-driven program is more cost-effective (Islam et al., 2002), however, no study has been conducted after the advent of PPP model or using the cost-utility method.

The costs can be incurred on the patient directly (direct costs) or indirectly (indirect costs) and most of the studies do not capture the whole picture as they often concentrate on pre-diagnosis, pre-treatment or treatment costs (KNCV Tuberculosis Foundation, 2008) only. Besides this the study will also capture the provider's cost of delivery health care services to the TB patients which is often absent in other studies.

This research proposal intends to contribute to the body of TB literature by illustrating the cost associated with different types of TB and to estimate the total economic burden of tuberculosis in Bangladesh. This dissertation also covers the economic evaluation of ongoing treatments approaches for both DS-TB and MDR-TB.



1.2 CURRENT TB CARE APPROACHES IN BANGLADESH

Bangladesh National TB control program (NTP) adopted the Directly Observed Treatment, Short Course (DOTs) strategy in November 1993. By 2007 the DOTS services were available throughout the country including metropolitan areas (National Tuberculosis Control Program (NTP), 2015).

NTP follows a PPP model where NGOs are working in collaboration with the MOHFW. There are about 12 NGOs working in different areas of Bangladesh. Among these BRAC, the largest NGO in the world and Damien Foundation Bangladesh, an affiliate of the Belgian NGO running TB control programs worldwide is the principal NGOs who get fund from Global Fund for Tuberculosis and Malaria (GFTAM) directly. BRAC gives away the funds to number of NGOs as the sub-recipient to work in different areas in Bangladesh. MOHFW also gets funding from the same source and equip the National Institute of Diseases of the Chest and Hospital (NIDCH) and number of Medical College Hospitals with diagnostic and treatment facilities for TB patients infected with both drug sensitive and drug resistant strains(National Tuberculosis Control Program (NTP), 2015).

For the drug sensitive TB patients, standard 6 months' regimen is followed by all participating NGOs. However, the mode of delivery is different for different NGOs. BRAC has employed Community Health Workers (CHWs) besides the DOTs centers to ensure patient compliance, while Damien Foundation (DF) trained and employed influential community members to help the patients to be adhered to the treatment protocol. Another NGO, Salvation Army Bangladesh, is using drug sellers at the



pharmacies as the counselor and drug distributors for the TB patients. Since involving different people, e.g., family members, neighbors, pharmacists falls under common strategy of involving community members. This study will conduct economic evaluation between these two different modes of DS-TB treatment delivery.

National Tuberculosis Control Program (NTP) in Bangladesh follows the 20-24 months treatment regimen for MDR-TB patients. It follows the Programmatic Management of Drug-resistant TB (PMDT) guideline (Falzon et al., 2011). The patients are admitted in the hospitals for first 6-8 months, after the intensive phase they are released to go to their respective home. From then on their treatment is supervised and administered by assigned CHWs (MOHFW, 2012). Intensive phase treatment for MDR-TB patients are provided in NIDCH, which is situated in Dhaka, and Chest Disease Hospitals (CDH) in Chittagong, Sylhet and Khulna.

Damien Foundation (DF) runs shorter protocol of treatment for MDR-TB patients, which span over 9 months. DF generally admits the MDR-TB patients in one of their three hospitals situated at Jalchatra of Madhupur, Tangail, Shomvuganj, Mymensingh and at Netrakona for the intensive phase of treatment which spans over four months followed by five months of continuation phase for which drugs are administered at patients' home (Damien Foundation, 2008). Rajshahi Chest Disease Hospital also provides 9 months treatment. DF also follows the strategy of involving the community members (Sharma, 2002) while BRAC programs deploy community health workers (CHWs) to reach the patients (Liu, Sullivan, Khan, Sachs, & Singh, 2011). This study will assess the cost-effectiveness of the two MDR-TB programs run by NTP and DF in Bangladesh.



1.3 RESEARCH OBJECTIVES

The study will focus on the following general objective:

To assess economic burden of drug sensitive and drug resistant tuberculosis on the afflicted population of Bangladesh and to investigate economic evaluation of the current approaches of Tuberculosis control in Bangladesh.

The specific objectives of the study are as follows:

- To analyze the direct and indirect cost of diagnosis and treatment of drugsensitive TB and MDR-TB in Bangladesh.
- To estimate the health system cost of diagnosis and treatment of drug-sensitive
 TB and MDR-TB in Bangladesh.
- 3. To measure effects as quality-adjusted life-years (QALYs) gained and disability-adjusted life-years (DALYs) averted.
- 4. To conduct a cost effectiveness analysis with QALYs and DALYs as the health outcomes.

1.4 STRUCTURE OF THE DISSERTATION

This dissertation proposal is divided into introduction, literature review, methods which are followed by three sections depicting the three studies conducted under the purview of the dissertation. Three studies are namely economic burden of TB in Bangladesh, economic evaluation of DS-TB treatment approaches and economic evaluation of MDR-TB treatment approaches in Bangladesh.



CHAPTER 2

LITERATURE REVIEW

2.1 Introduction and Scope of Review

This chapter reviews the theoretical concepts of illness with tuberculosis and its impact on individuals, families/households, and society as a whole. Following that the chapter includes literature review of existing studies on economic burden of tuberculosis studies as well as the studies on cost-effectiveness of both drug sensitive and drug resistant TB treatments all around the globe.

2.2 ECONOMIC BURDEN OF TUBERCULOSIS

2.2.1 THEORETICAL CONCEPTS OF ECONOMIC BURDEN OF TUBERCULOSIS

In cost analysis three types of costs are taken into account, direct costs, indirect costs and intangible costs due to illness. Direct costs include healthcare costs (hospital, medication, emergency transportation, outpatient visit charges) and family costs (out-of-pocket payment (OOP), medication, transportation of families etc.). Indirect costs include the opportunity costs of work-loss days, lost productivity/income on the part of both the patients and his/her relatives. Intangible costs can arise from the functional limitations, pains acquired in the process and cannot be quantified and highly subjective in nature (Centers for Disease Control and Prevention (CDC), 2013).



Besides these costs, there is another important cost incurred by the households through coping strategies, which includes sale of assets, taking up debt, saving on food or other items, taking a child out of school to care for the patient or taking up another job (Russell, 2004).

The costs of communicable diseases fall on the households in three distinct phases:

- 1. Pre-diagnosis
- 2. During Diagnosis/Pre-treatment
- 3. During Treatment

The causal linkages of these factors are depicted by Russell (2004):

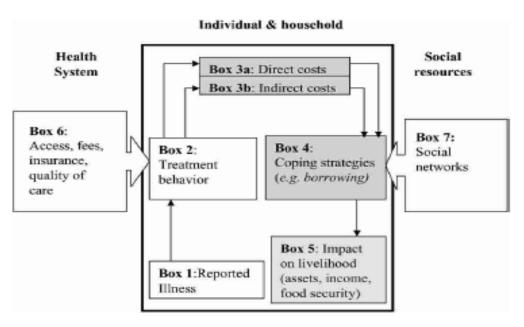


FIGURE 1. Conceptual framework for analyzing the economic burden of illness for households.

Figure 2.1 Conceptual Framework for analyzing the economic burden of illness for households (Russell, 2004)



At the stage of boxes 1 and 2, decisions are made whether and how treatment is sought as a response to the event of illness. The health system is captured in Box 6.

Direct costs capture expenditures related to seeking treatment while indirect costs are loss of labor time for patients and their caregivers. The severity of illness and characteristics of health services affect direct and indirect costs and influence access to and choice of provider. The cost burden and coping strategies of struggling with this burden (mobilizing resources outside the household such as credit) determine household assets and impoverishing processes, hence the link between illness and poverty (Russell, 2014).

McIntyre et al. (McIntyre et al., 2006) provided a flow-chart on economic consequences of TB illness and payment for healthcare (Figure 2.2). According to McIntyre there are four stages of tuberculosis treatment, e.g., illness experience, treatment seeking behavior, economic consequences, and coping strategies and social resources. In preliminary stages, perception of illness and treatment seeking can be affected by the economic status of the person. If the patient with TB does not seek treatment then only indirect costs is incurred in form of lost productivity, while if the patient goes for treatment then direct costs also incurred on top of indirect costs. Indirect costs further subdivided into productivity loss of the patient and the caregiver, whereas direct cost is categorized into financial costs of healthcare (services and medicines) and other financial costs.



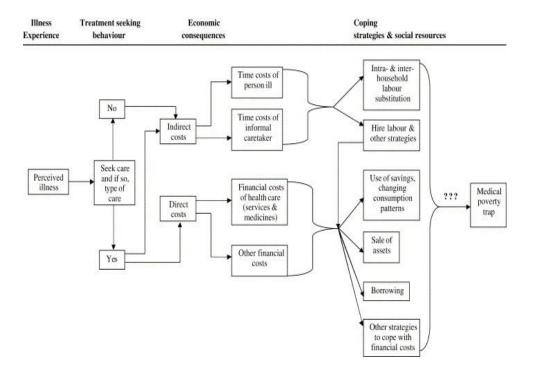


Figure 2.2 Simplified flow-chart of key issues relating to the economic consequences of illness (McIntyre, Thiede, Dahlgren, & Whitehead, 2006)

Based on conceptual model of McIntyre et al. (McIntyre et al., 2006) Laokri et al. (Laokri et al., 2014) (2014) provided an extended conceptual framework incorporating intangible costs, e.g., pain and suffering, and social stigma; elaborating coping costs and social burdens due to illness; and includes societal economic loss along with illness poverty trap ensued due to illness (Figure 2.3). This extended model also includes forgone activities of the informal caregivers as well as those of guardians accompanying the TB patients for treatment. These foregone activities can culminate into labor substitution, withdrawal of children from school, and informal caregiving activities of the family members. Guardians lose time and income. Direct costs have been subdivided into subsidized healthcare costs, non-subsidized healthcare costs, and non-healthcare costs. Non-subsidized healthcare costs and non-healthcare costs along with income loss of the guardian can result in (1) Financial resource mobilization, e.g., borrowing, selling assets,



pledging, extra-earnings etc.; (2) Resource reallocation, e.g., dissaving, budget cuts, deprivation, delayed investment etc. Non-subsidized healthcare costs and non-healthcare costs can also lead to erratic care pathways which include redundant care visits, alternative care seeking, diagnosis and treatment delays, and care interruption.

Intangible costs like pain and sufferings and social stigma can impose social consequences like (1) Low awareness: low awareness of disease, denial of illness status, bad living conditions, fear of losing position, social isolation, lack of family support, patient-related delays; (2) Social exclusion: exclusion from services including public healthcare services, from income and from participation.

Between McIntyre's and Laokri's conceptual frameworks, both of which are built upon the framework proposed by Russell (2004), simpler McIntyre's framework will be adopted for this study. Both frameworks are more or less similar, while Laokri's one is more detailed and includes pain and suffering, and social stigma. It is difficult to assign monetary value to intangible costs like pain and stigma. On the other hand, these intangible phenomena affect the quality of life of the patients. In our adopted quality of life measurement tool pain and stigma have included to offset their absence in the costs estimation. It also prevents double counting; once included in costs and again take into account while measuring quality of life.

There are four approaches to measure the cost of illness, e.g., human capital method, willingness to pay method, production cost and friction cost method (Jo, 2014; Malaney, 2003). Out of these human capital and willingness to pay methods are best suited for calculating the costs of illness from the patient perspective (KNCV



Tuberculosis Foundation, 2008). This study intends to employ both the method to capture different dimensions of costs. As Human Capital method (HCM) captures the valuation of forgone income and productivity and willingness to pay (WTP) captures the subjective reporting of actual costs accrued to the household and the perceived costs of the illness. It is argued that the HCM underestimates the total cost of illness than the WTP method, since it fails to capture the costs which are difficult to measure in numeric terms, e.g., costs associated with pains and sufferings.

The study will estimate the cost to society in the form of lost future productivity discounted to the present. The calculations aim at a sum of future earnings of the premature dead by looking at life expectancy, labor force participation and average salary data. This is sometimes called the 'top-down-approach'. It includes direct and indirect costs. Indirect costs are productivity losses, measured by estimating income foregone due to morbidity and mortality. The cost of morbidity is the value of lost workdays. Future earnings are discounted to assess the present value of lost income.

The study will measure the total cost of illness including the costs of illness in the past and present along with the future projection of the costs based on the collected data. The future projection of the costs is important in the sense that the debilitating disease may accrue a long-term costs burden on the family and the society as a whole and these costs can be quite high, even though often not considered explicitly in cost of illness analysis. As in the case of Measles vaccination, it is found that although measles vaccination prevents deaths within a short time frame (preventing measles death) it also has longer term impact on child survival. In other words, mortality rate among measles vaccinated children were lower than comparable children without vaccination (Koenig et



al., 1990). Productivity effect of malaria persists for many years after the control of malaria outbreaks in an area and imposes huge economic burden in the long run (Breman, Egan, & Keusch, 2001). It is also likely that TB has these indirect long-term effects and this study will try to get a handle on these longer-term outcomes.

From the institutions like DOTS centers, hospitals treating TB patients etc. data will be collected for the variable costs, which is the direct function of number of patients treated and includes costs such as drugs, reagents, and food during hospitalization. Then the fixed costs like personnel salaries, costs of vehicles and their maintenance and other administrative costs will also be enumerated to estimate programmatic costs (Murrat et al., 1993).

2.2.1 EXISTING EVIDENCE ON ECONOMIC BURDEN OF TUBERCULOSIS

Islam et al. (Islam et al., 2002) conducted a cost-effectiveness study comparing between community health workers (CHW) model adopted by BRAC and government run TB program which did not include CHWs. As part of the study the authors estimated the cost of delivering TB treatment from both patient and provider perspectives. From the study areas they collected the costs of all health workers and administrative staff, BRAC capital costs (including building costs) derived from accounting books and financial reports and in the absence of government report on capital costs those were estimated based on the local market price and current replacement costs. Capital costs were annuitized by using 10 years lifetime for furniture, 5 years for vehicles and equipment and were discounted at 5% per annum rate. Training costs were excluded assuming the costs equal across the two types of programs.



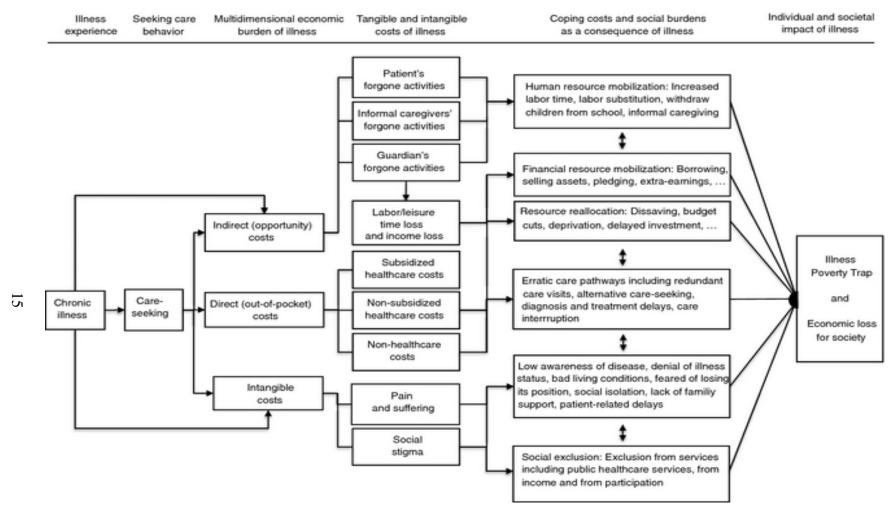


Figure 2.3 Conceptual framework to assess multidimensional economic burden of illness in a user's perspective (adapted from McIntyre et al., 2006) (Laokri, Dramaix-Wilmet, Kassa, Anagonou, & Dujardin, 2014)

Recurrent costs were collected from accounting books and financial reports from both programs and overhead costs for TB programs were calculated to be 10% in BRAC facilities and 5% in government run facilities. Patient's costs were elicited by interviewing 18 BRAC and 20 government patients. Time and travel costs associated with patients visits to health facilities for diagnosis, drug collection, and follow-up tests as well as costs of people accompanying the patients in each visit was included in patient's costs. It was calculated that the total cost was about \$10 (422 BDT in 1996-1997) in BRAC areas while the cost was US\$19 (802 BDT in 1996-1997) in government facilities (Islam et al., 2002). Using the Consumer Price Index (CPI) reported in the World Bank website we get that the costs was in 1,392 in 2015 BDT in BRAC areas, while the costs in government areas were 2,646 in 2015 BDT (The World Bank, 2017a).

Another study estimated the patient costs during TB treatment in Bangladesh and Tanzania (Gospodarevskaya et al., 2014). Total 96 patients were interviewed to find out the patient costs for six months DSTB treatment. The study sample includes 67 patients from BRAC, 22 from Damien Foundation, and 7 from Population Services and Training Centers. Total costs incurred during the six-month treatment regimen was estimated by combining the cost incurred during two months of intensive phase plus twice the costs incurred during two months of continuation phase. Total cost includes travel costs, guardian/accompanying person costs, caregiver cost, as well as treatment costs, e.g., laboratory tests, administration fees, hospital admission charges, medicines and supplements costs. Income lost due to TB for the patients and the guardians who would otherwise be paid through employment was calculated to determine the indirect costs. Indirect costs for students, prisoners, and those who were unemployed before the TB



illness were not estimated assuming that these groups of people had no foregone costs. But for those who do housework, their foregone wage at the rate paid for maid service was calculated and added to the patient costs. Total patient costs in Bangladesh was \$224 in 2012, which is equivalent to 16,690 in 2012 BDT and 20,720 in 2015 BDT.

Interestingly both of these studies excluded the costs incurred by patients before reaching the treatment facilities. This constitutes a major source of costs for the patients. Since the TB treatment is almost free except for traveling, this cost constitutes a major portion of the total costs. A study conducted by Croft et al. (Croft & Croft, 1998) among TB patients in Nilphamari, Bangladesh showed that a mean financial cost to the patients due to foregone income and payments for doctors' consultation and medicines were \$245 in 1996 which is equal to \$808 in 2015 and BDT 64,663 in 2015 (1 Dollar=80 BDT).

There are several systematic reviews on TB patient and health system costs have been done. Laurence et al. (Laurence, Griffiths, & Vassall, 2015) searched for cost and economic evaluation studies on both DS-TB and MDR-TB between January 1990 and February 2015. The authors found mean DS-TB treatment costs were \$273 in lower middle-income countries (LMICs) and \$258 in low income countries (LICs), whereas the MDR-TB treatment costs were \$6,313 and \$1,218 respectively. Tanimura et al. (Tanimura, Jaramillo, Weil, Raviglione, & Lönnroth, 2014) focused only on LMICs and searched the database from inception to March 31, 2013. Mean total costs ranged from \$55 to \$8,198, with an unweighted average of \$847. Half of the total costs were reported before treatment started, while the composition of costs was 20% to direct non-medical costs, 20% to direct medical care costs, 60% to income loss due to TB illness. In a study conducted with published literature on African countries showed that mean pre-



diagnostic costs were between \$36 and \$196, while post-diagnostic costs were between \$17 and \$448.

2.3 HEALTH RELATED QUALITY OF LIFE (HRQOL)

In this section studies on health related quality of life (HRQoL) of TB patients is reviewed. Main focus is on different measurements of HRQoL, e.g., Quality Adjusted Life Year (QALY) and Disability Adjusted Life Year (DALY). Then different measurement scales used for eliciting HRQoL measures for TB patients is explored.

Quality of life measurement uses utility theory to identify the degree of health concerns related to any disease or health conditions. Quality adjusted life year (QALY) and Disability adjusted life year (DALY) are the two summary estimates widely used in health economic evaluations. While QALY views health outcomes from the perspective of "healthiness", DALY views health outcomes in terms of loss of life years due to disabilities.

QALYs gained =
$$Q^{i} \frac{1-e^{-rL^{i}}}{r} - Q \frac{1-e^{-rL}}{r}$$
 (1)

where L^i and Q^i are, respectively, the period over which treatment affects the individual's quality of life, and the quality of life weight with treatment; while L and Q are the corresponding parameters without treatment.

The formula for calculating the number of QALYs gained through an intervention i is as follows:



$$QALYs \ gained = \sum_{p=1}^{p} Q_{p}^{i} \frac{e^{-r(t_{p}^{i}-a)} - e^{-r-r(t_{p-1}^{i}-a)}}{r} - \sum_{m=1}^{N} Q_{m}^{i} \frac{e^{-r(t_{m}^{i}-a)} - e^{-r-r(t_{m-1}^{i}-a)}}{r}}{r}$$
(2)

here the life expectancy with the intervention (L_i) at age a is divided into P time periods n_p , and Q_p^i is a vector of health-related quality of life weights predicted (or observed) for each time period n_p following the intervention. While Q_m^i is weight associated with the health state before intervention and individual's residual life expectancy is divided into N time periods n_m . Here t_p and t_m are the of individual years within the life expectancy.

On the other hand, Disability Adjusted Life Year (DALY) is the sum of the Years of Life Lost (YLL) due to premature mortality and the Years Lost due to Disability (YLD) for people living with the health condition or its consequence (World Health Organization (WHO), 2013a).

$$DALY = YLL + YLD$$
 (3)

We can derive the formula for YLL and YLD as follows (Diel et al., 2014):

$$YLL = \frac{K \cdot C \cdot e^{r \cdot a}}{\left(r + \beta\right)^2} \cdot \left[\left[1 + \left(r + \beta\right) \cdot a \right] \cdot e^{-\left(r + \beta\right) \cdot a} - \left[1 + \left(r + \beta\right) \left(L + a\right) \right] \cdot e^{-\left(r + \beta\right) \left(L + a\right)} \right] + \frac{1 - K}{r} \left(1 - e^{-r \cdot L} \right)$$
(4)

$$YLD = DW \cdot \left[\frac{K \cdot C \cdot e^{r\alpha}}{(r+\beta)^2} \cdot \left[\left[1 + (r+\beta) \cdot \alpha \right] \cdot e^{-(r+\beta)\alpha} - \left[1 + (r+\beta)(T+\alpha) \right] \cdot e^{-(r+\beta)(T+\alpha)} \right] + \frac{1-K}{r} (1 - e^{-r.T}) \right]$$
(5)

Where, K = Age-weighting modulation constant (1.00), C= Age-weighting scaling constant, L= country-specific standard life expectancy at age of death (years),



DW = Disability Weight (0.333; 95% CI= 0.224-0.454) as per the Global Burden of Disease 2013 study weights (Salomon et al., 2015), T= treatment duration, and α = age of onset of disability.

2.3.1. HEALTH RELATED QUALITY OF LIFE OF TUBERCULOSIS PATIENTS

Although several studies have been conducted to assess the health-related quality-of-life (HRQoL) for tuberculosis patients (Brown et al., 2015), there is no well-accepted tuberculosis-specific HRQoL measurement instrument available. Most studies use EQ-5D, SF-36, SF-6D and other generic HRQoL instruments (Guo, Marra, & Marra, 2009). Only one study conducted in India used a TB-specific tool named DR-12, which has 12 items each ranked on a scale of 1–3 (Dhingra & Rajpal, 2003). Recently a multidimensional TB-specific HRQoL instrument named Functional Assessment of Chronic Illness Therapy-Tuberculosis (FACIT-TB) was developed and psychometrically validated in Iraq (Dujaili et al., 2015). This FACIT-TB instrument includes physical, mental, social and economic, functional, as well as spiritual well-being of the TB patients. This instrument is unique in incorporating questions on adverse drug reaction (ADR), perception about social stigma, and spirituality related with TB.

The scale comprises 45 items: 17 items covering physical well-being (possible score range 0–68), seven items covering social and economic well-being (possible score: range 0–28), 11 items covering emotional well-being/living with TB (possible score range 0–44), seven items covering functional well-being (possible score range 0–28), and three items covering spiritual well-being (possible score range 0–12). A 5-point Likert type scale ranging from 0 (not at all) to 4 (very much) is assigned to each item.



2.4 Cost-Effectiveness of Tuberculosis Programs

Till date, to our best knowledge, the only cost-effectiveness analysis between two TB control programs run by BRAC and government was conducted by Islam et al. (2002). It showed that the government program was 50% more expensive for similar outcomes.

Many cost-effectiveness analyses have been done in order to determine the cost-effective diagnostic techniques, e.g., sputum examination (Walker et al., 2000), serological tests vs. other diagnostic tests (Dowdy, Steingart, & Pai, 2011), dual or single test for detection of latent tuberculosis infection (LTBI) (Pooran et al., 2010), . However, since the focus of this thesis is to compare between two TB control programs we restrict our review among those studies which conducted cost-effectiveness analyses between programs.

Using Denver General Hospital data Burman et al. showed that although DOT is costly at the outset it turns to be cost-effective than Self-administered Therapy (SAT) because of higher cure rates (Burman, Dalton, Cohn, Butler, & Reves, 1997). The outcome variable for this study was cure rate per cost unit.

Using published literature, records, and expert opinions Baltussen et al. showed that DOT as well as incremental programs like DOTS plus, Full combination of DS-TB and MDR-TB strategies all are cost-effective in terms of DALYs averted per cost unit in high burden TB countries in Africa and South-East Asia (Baltussen, Floyd, & Dye, 2005).



Several other studies have been conducted in different countries, e.g., Thailand (Hunchangsith, Barendregt, Vos, & Bertram, 2012), Egypt and Syria (Vassall, Bagdadi, Bashour, Zaher, & Maaren, 2002), Botswana (Moalosi et al., 2003), Haiti (Jacquet et al., 2006), Uganda (Okello, Floyd, Adatu, Odeke, & Gargioni, 2003), Brazil(Mohan, Bishai, Cavalcante, & Chaisson, 2007). These studies invariably documented that the DOTs strategy or involving the communities in the care process is cost-effective over SAT.

In two studies conducted in South Africa (Sinanovic et al., 2003), and in India (Pantoja et al., 2009) the authored showed that PPP models were more cost-effective by virtue of reducing costs to patients by 64-100% in South Africa; while the patient cost fell from US\$154 to US\$132 over four-years period in India.

A recent study shows that shortening of the DS-TB treatment from six-months to four-months remain cost-effective option for Brazil, South Africa, Bangladesh and Tanzania (Gomez et al., 2016). Another study results also support this finding in South Africa (Knight et al., 2015).

Several studies have also been conducted to assess the cost-effectiveness of different MDR-TB treatment regimens. Fitzpatrick et al. (2012) conducted a systemic review of studies which used primary data and outcome which eventually includes only four studies conducted in Estonia, Peru, the Philippines, and Tomsk, Russia. Cost per DALY averted were \$598, \$163, \$143, \$745 respectively. The cost per DALY averted was lower than GDP per capita in all 14 WHO sub-regions considered.

However, there was no study comparing between the shorter and longer regimen for MDR-TB treatment. Two separate studies conducted by DF scientists assessed the



effect of two regimes. One study which was conducted for the standardized regimen of 21-24 months published in 2004 (Van Deun, Salim, Kumar Das, Bastian, & Portaels, 2004) and another study on shorter regimen was published in 2010 (Deun et al., 2010). Both of the studies showed that both treatment strategies are successful in treating MDR-TB patients; however, in absence of any comparative cost-effectiveness analysis between them we cannot tell which one is better.

2.4.1 METHODS OF COST-EFFECTIVENESS OF TUBERCULOSIS PROGRAMS

Cost-effectiveness analyses with DALY or QALY as outcome variable usually employs various models like population model PopMod (Baltussen et al., 2005), Monte Carlo simulation technique (Tupasi et al., 2003), dynamic state-transition model of TB (Resch, Salomon, Murray, & Weinstein, 2006) etc. In this study, to conduct the economic evaluations of different tuberculosis control programs, we shall use the Monte Carlo simulation technique. A probabilistic Markov Chain Monte Carlo (MCMC) simulation model will be fitted. The patient level data on treatment outcomes will help to get the transitional probabilities between states as well as we can use regression techniques to get the probabilities along with the uncertainties. That will help us to conduct the sensitivity analysis of the results. Finally, we will estimate the Incremental Cost-Effectiveness Ratio (ICER) between the different comparators and use the acceptability curves approach to find out the cost-effective TB control program in Bangladesh.

Markov model has some unique characteristics which fit the progression of TB well. For example, in Markov model states are mutually exclusive, states are complete (i.e. no people are lost) and people remain in that state for a fixed period of time. Also,



Markov model is preferred over the decision trees when health event repeats over time, or have longer term health effects, effect of treatment either stops quickly after initial treatment or continue at an earlier level, and the risk of different health events does not depend on patient's prior history (Briggs, Claxton, & Sculpher, 2006).

Here we can represent the Markov model for TB as below:

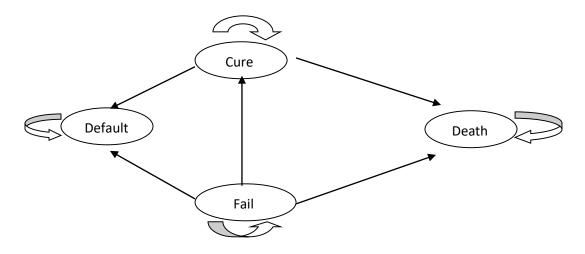


Figure 2.4 Simplified Markov Model for Outcomes of Illness with Tuberculosis

From the Markov model we can find that after starting of the treatment the MDR-TB patient can move to any of the four states, e.g., cure/treatment complete, failure/relapse, default, and death. Here default and death are the absorbing states. If any patient is cured he/she can remain cured, relapse/reinfection may occur, or can be dead. On the other hand, the failed/relapsed patients undergo another cycle of treatment and can culminate into cure, remain failed, can default, or can be dead as well.

Infectious disease often requires dynamic models which reflect the rate of transmission of disease among the population. The rate of infection is a function of the number of infected individuals in the community (Briggs et al., 2006). Epidemiology of an infectious disease is important to take into account because due to change in the



natural history of disease can affect the outcome of the disease and thereby, the costeffectiveness analysis will be flawed (Jit & Brisson, 2011).

TB is an infectious disease and have latent period and many latent cases which does not turn into a full blown disease. But these latent TB infection (LTBI) can be activated upon proper stimuli or absence of immunity, as in the case of HIV/AIDS. Therefore, many researchers included these aspects in the infectious disease modeling (Jacquet et al., 2006; Menzies, Cohen, Lin, Murray, & Salomon, 2012; Oxlade, Piatek, Vincent, & Menzies, 2015; White & Abubakar, 2016). Dowdy et al. (2013) synthesized that a single model is unlikely meet all criteria for all studies and prepared a wish list for the TB modelers would love to have (Dowdy, Dye, & Cohen, 2013)



CHAPTER 3

METHODS

3.1 Introduction

Bangladesh National Tuberculosis control program (NTP) follows a PPP model where NGOs are working in collaboration with the MOHFW. There are about 12 NGOs working in different areas of Bangladesh. Among these the largest NGO in the world-BRAC and Damien Foundation Bangladesh, an affiliate of the Belgian NGO running TB control programs worldwide are the principal NGOs. These NGOs are recipients of the funds provided by Global Fund for Tuberculosis and Malaria (GFTAM) along with NTP being the principal recipient. BRAC gives away the funds to number of NGOs as the sub-recipient to work in different areas in Bangladesh. NTP uses the fund for equipping the National Institute of Diseases of the Chest and Hospital (NIDCH) and number of Medical College Hospitals with diagnostic and treatment facilities for tuberculosis (TB) patients infected with both drug sensitive and drug resistant strains (MOHFW, 2014).

For the drug sensitive TB patients standard 6 months regimen is followed by all participating NGOs. However, the mode of delivery is different for different NGOs. BRAC has employed Community Health Workers (CHWs) besides the DOTs centers to ensure patient compliance, while Damien Foundation (DF) trained and employed influential community members to help the patients to be adhered to the treatment protocol.



Another NGO, Salvation Army Bangladesh, is using drug sellers at the pharmacies as the counselor and drug distributors for the TB patients. This study will conduct economic evaluation between these two different programs.

NTP in Bangladesh follows the 20-24 months treatment regimen for Multi-Drug Resistant TB (MDR-TB) patients. It follows the Programmatic Management of Drugresistant TB (PMDT) guideline (Falzon et al., 2011). Initially MDR-TB patients are admitted to designated hospitals for intensive phase of treatment which generally last for 6-8 months. Then the patients are released to community and their treatment is supervised and administered by CHWs for another 14-16 months (MOHFW, 2012). Damien Foundation (DF) runs their own protocol of treatment for MDR-TB patients, which span over 9 months and differs in mode of treatment. DF generally admits the MDR-TB patients in one of their three hospitals situated at Jalchatra of Madhupur, Tangail, Shomvuganj, Mymensingh and at Netrakona for the continuation phase of the treatment which spans over four months followed by five months of continuation phase for which drugs are administered at patients' home (Damien Foundation, 2008). DF also follows the strategy of involving the community members (Sharma, 2002) while BRAC programs deploy community health workers (CHWs) to reach the patients (Liu et al., 2011). This study will assess the cost-effectiveness of the two MDR-TB programs run by NTP and DF in Bangladesh.

3.2 STUDY DESIGN

The study follows a stratified random sampling method. From the 64 districts of Bangladesh nine districts from the eight divisions (at least one from each division) were selected based on the high and low burden of TB cases. Then from each district two



upazilas (sub-districts) will be selected randomly. From the registry of the DOTS center of the UHCs of these eighteen upazilas lists of TB patients currently undergoing treatment or recently finished will be collected.

3.3 STUDY SITE

3.3.1 Drug Sensitive TB (DS-TB)

For total representation at least one district from all eight divisions of Bangladesh was selected for the study. Since Dhaka division is bigger in size, three districts was selected from Dhaka division including an urban area of Dhaka city. Selection was made based upon the high and low TB burden among all districts. Therefore, five high burden and four low burden districts have been selected under the purview of the study. Two upazilas (sub-districts) from each of the selected districts was selected randomly. Following is the list of all districts and upazilas covered under the study. From each upazila 50 DS-TB patients were selected randomly for interview.

3.3.2 Multi-Drug Sensitive TB (MDR-TB)

For assessing economic burden of MDR-TB patients and the economic evaluation of comparator MDR-TB control programs about 175 MDR-TB patients will be selected purposively. According to the recent estimates in 2014 number of laboratory-confirmed MDR-TB patients was 994 in Bangladesh (World Health Organization (WHO), 2015a) and the prevalence of MDR-TB is 5,100 in 2015 (World Health Organization (WHO), 2016a). In our study areas the number will be clearly significantly lower. Therefore, we collected the information of the MDR-TB patients from the TB



control programs and reach those who (Brazier, Roberts, & Deverill, 2002) were accessible.

3.4 STUDY PARTICIPANTS

The inclusion criteria for the study participants will be as follows:

- a) Older than 18 years of age,
- b) Suffering or recently suffered from pulmonary TB (DS-TB/MDR-TB),
- c) Undergoing treatment or finished treatment within last 6 months.

3.5 SAMPLE SIZE

Glick (H. A. Glick, 2011) proposed a sample size formula for cost-effectiveness evaluation of clinical trials. Although our study is not a typical clinical trial, given the nature of the intervention and the study design we can apply the formula for calculating the required sample size for our study. The formula calculates the sample size for each of the two groups with similar standard deviation of costs and effect and same sample size:

$$n = \frac{2(Z_{\alpha} + Z_{\beta})^{2}[sd_{c}^{2} + (W * sd_{q}^{2})^{2} - (2W\rho * sd_{c} * sd_{q})}{(WQ - C)^{2}}$$

Where:

 Z_{α} is the Z-statistic for the level of Type I error (set at 95%)

 Z_{β} is the Z-statistic for the level of Type II error (set at 80%)

sdq, sdc are the std deviations for each group for treatment effect and cost respectively



W is the Maximum Willingness to Pay

Q is the expected mean difference in treatment effectiveness

C is the expected mean difference in treatment cost

 ρ is the expected correlation of the difference in cost (C) and effect (Q)

This is a measure of the covariance of changes in effectiveness and changes in cost. Negative covariance, where cost decreases with increasing effectiveness result in a larger sample size. Positive covariance where cost increases with increasing effectiveness result in smaller sample sizes.

DSTB:

With 95% confidence interval and 80% power of the test, we assumed that the standard deviation of costs (sd_c) is 400 USD, standard deviation of effect (sd_q) is 0.2 QALY, ρ , correlation of difference in cost (C) and effect (Q) is 0.4. The expected mean difference in treatment effectiveness (Q) is 0.4 QALY and expected mean difference in treatment cost (C) is 500 USD. We set the willingness-to-pay threshold (W) at the three times of GDP of Bangladesh which is 3942 USD (Macroeconomics, 2001). We found the sample size for both treatment groups is 405.

MDR-TB:

With 95% confidence interval and 80% power of the test, we assumed that the standard deviation of costs (sd_c) is 100 USD, standard deviation of effect (sd_q) is 0.25 QALY, ρ , correlation of difference in cost (C) and effect (Q) is 0.5. The expected mean difference in treatment effectiveness (Q) is 0.15 QALY and expected mean difference in



treatment cost (C) is 1000 USD. We set the willingness-to-pay threshold (W) at the three times of GDP of Bangladesh which is 3942 USD (Macroeconomics, 2001). We found the sample size for one group is 70 and another is 104 with a 2:1 sample size ratio.

Table 3.1 List of Districts and Sub-districts where the survey among DS-TB patients was conducted

Division	District	Sub-District	Number of Patients Surveyed	Treatment Delivery
Barisal	Pirojpur	Pirojpur Sadar	50	Community Health Worker
		Najirpur	50	
Chittagong	Laxmipur	Laxmipur Sadar	50	Community Health Worker
		Ramganj	50	
Dhaka	Faridpur	Faridpur Sadar	50	Community
		Nagarkanda	50	Member
	Manikganj	Manikganj Sadar	50	Community
		Shingair	50	Health Worker
	Dhaka	Mirpur (Urban Area)	100	DOTs center, Pharmacists
Khulna	Kushtia	Kushtia Sadar	50	Community
		Doulatpur	50	Health Worker
Mayoranainah	Netrokona	Netrokona Sadar	50	Community
Mymensingh		Kendua	50	Member
Rajshahi	ChapaiNawabganj	ChapaiNawabganj Sadar	50	Community
		Shibganj	50	Member
D	Panchangarh	Panchagarh Sadar	50	Community Member
Rangpur		Debiganj	49	
Sylhet	Habiganj	Habiganj Sadar	50	Community
		Bahubal	51	Member
Total			1,000	



Study Sites (DS-TB)

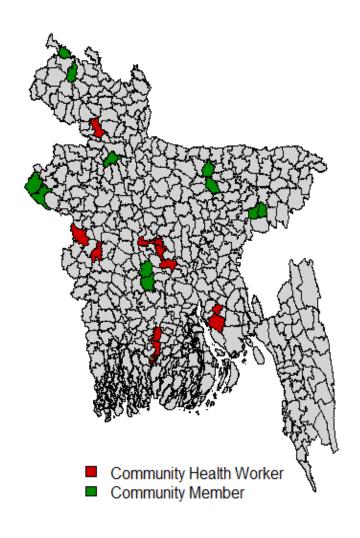


Figure 3.1 Study Sites (Sub-districts) where DS-TB patients were interviewed

Table 3.2 List of Districts where the survey among MDR-TB patients was conducted

Division	District	Number of Patients Surveyed	Treatment Regimen
Chittagong	Chittagong	78	20-24 Months Regimen
Dhaka	Dhaka	23	20-24 Months Regimen
Mymensingh	Mymensingh	17	9 Months Regimen
	Netrokona	32	9 Months Regimen
Rajshahi	Rajshahi	18	9 Months Regimen
Total		168	

3.6 ETHICAL CONSIDERATION

The study has already got ethics approval from University of South Carolina in the USA where the PI is a PhD student. Institutional Board Review (IRB) approval will also be taken from Jahangirnagar University in Bangladesh. A third and final approval was obtained from WHO Research Ethics Review Committee (WHO ERC).

The participants were approached at their households by the trained enumerators.

At first the enumerators politely introduced themselves and asked pleasantries. After establishing rapport, the enumerators conveyed their purpose of the visit.



Study Sites (MDR-TB)

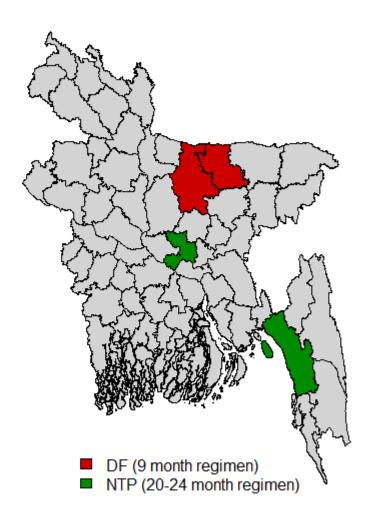


Figure 3.2 Study Sites (Districts) where MDR-TB patients were interviewed

An informed consent was obtained from each participants of the study. At the

beginning the enumerator read the introduction in comprehensible manner to the patient.

In the introduction the name and whereabouts of the investigators, name of the funding agency, and the purpose of the study are furnished. It also attests the right of the



respondent to withdraw any time during the interview. The benefit of participating in the study is also described in that part.

The data is kept in one laptop and under lock and key in the office of the principal investigator. No one except the investigator have access to the stored data. For data analysis the patient's information is de-identified. Thus the data analysis and presentation in the report is completely anonymous and in any circumstances it will be kept confidential.

3.7 Data Collection Instruments

3.7.1 Patient Questionnaire

Stop-TB questionnaire on patient's cost has been adopted for the study. The patient questionnaire includes questions on the previous TB treatment costs including number of visits, tests, drugs, travel, food, accommodation, and out-of-pocket and insurance costs etc. The questionnaire also includes questions on the current or recent treatment for TB, which comprises of treatment costs, follow-up costs, costs borne by family or friends, hospitalization costs, food costs, other comorbidities cost, insurance, coping costs. Therefore, the cost instrument is a comprehensive tool to capture all costs incurred by the patients and their families for TB. The questionnaire was field tested among the TB patients and changes were made accordingly to make the question understandable and answerable.

The patient questionnaire also includes the health-related quality of life questions. In this study Functional Assessment of Chronic Illness Therapy-TB (FACIT-TB) questionnaire (Dujaili et al., 2015), which includes 45 items under five sub-groups, will



be incorporated along with smaller generic instruments like EuroQol-5 Dimensions-5 Levels (EQ-5D-5L) with visual analogue scale (VAS), and SF-6D, which is an abridged well-validated version from SF-36 (Brazier et al., 2002). These generic instruments will be used to assign utility scores for various components of the FACIT-TB instruments. Mapping function will be used to predict the utility values. This approach involves estimating the relation between a non-preference-based measure (like FACIT-TB) and generic preference-based measure using statistical association and this approach requires overlap between the two measures applied on the same population (Young, Mukuria, Rowen, Brazier, & Longworth, 2015). Multinomial logistic regression models will be estimated for each dimension, and the estimates from these regressions will be used to categorize respondents into five levels of each of the EQ-5D dimensions and thus predict the EQ- 5D health state for each respondent. A total of 1000 Monte Carlo simulations will be run to estimate EQ-5D health states. The standard set of UK general population values will be then applied to each predicted health state to obtain EQ-5D values. Mapping is usually performed using regression analysis and often preferred regressions are OLS or tobit.

3.7.2 Provider Questionnaire

Institutional level data was also collected for assessing the direct health system costs associated with TB treatment. From this secondary source the data on the number of patients diagnosed and treated and the outcome of the diseases in terms of complete cure, remission, relapse or death was collected. Another questionnaire for the program managers was used in the study to collect data on the facility and personnel level costs for the TB control programs (included in the Annexure). Market values of the TB drugs was

collected for estimating the drug costs for patients under various programs. Those who are involved in the TB control programs on honorary basis, their opportunity costs of the time were calculated using the average wage rates for service holders using secondary sources.

3.8 DATA COLLECTION

Given the extent of household level primary data collection in 10 randomly selected districts a total of two survey teams consisting of sixteen data collectors were formed with one supervisor, one back checker and six enumerators in each team. The questionnaires were prepared in consultation with my supervisor and mentors at USC. The questionnaires were pre-tested at the field level by the selected survey teams before the actual data collection began.

3.8.1 QUALITY ASSURANCE

The study employed multilevel quality assurance process for data collection.

Researcher will execute a four-step scrutiny process to ensure the reliability and validity of the information. For each of the steps, certain team members will be given specific responsibility to manage the quality assurance process.

3.8.2 Cross Check

Enumerators collected quantitative data directly from households. Later, enumerators went to the households with filled questionnaire where other enumerators had collected the answers. This cross check by other enumerators helped to recover any primary mistakes in collecting data.



3.8.3 ACCOMPANY CHECK AND SPOT CHECK

Field team supervisor carried out next level data check through accompany check and spot check. Accompany check includes accompanying enumerators during data collection, validating the information provided by the respondents, throwing of questions, examining proper coding and collection of information. Field team supervisor also scrutinized the information through spot check by going to the households after the enumerators leave the households. Such meticulous checking system ensured the quality of data effectively.

3.8.4 Back Check

The third level quality assurance was carried out by the team supervisors. Team supervisors visited households randomly with filled up questionnaires to examine the accuracy and reliability of information. Three-layer cross checking in the field ensured high quality data collection.

3.8.5 OTHER QUALITY CHECK AND FEEDBACK

After data compilation, the data set was sent to the PI for his feedbacks and other quality measures. The Principal Investigator travelled intensely during the period of data collection and ensured the quality of the data collection.

3.9 Data Analysis

Data analysis is primarily performed in STATA 14.2. Patient and provider level costs data are being managed and analyzed. Regression modelling for finding important predictors for patient costs.



For cost-effectiveness analysis R will be used for the ease of estimation. Markov simulation modeling will be performed in Winbugs and Just Another Gibbs Sampler (JAGS). Both of these softwares are open source and can easily be used from within R. R is also an open source software for which many packages are found suitable for cost-effectiveness analysis (Sutton, Welton, Cooper, Ades, & Abrams, 2012). BCEA is one of those packages. It helps to analyze cost-effectiveness within a Bayesian framework (Baio, 2012)

3.10 EXPECTED OUTCOME OF THE STUDY

Tuberculosis is a deadly tropical disease affecting the people of developing countries and incurring huge cost on the economy. Thus economic evaluation of this disease in a developing country setting will provide the researchers, policy makers an empirical evidence of the extent of the cost burden.

3.11 DISSEMINATION OF RESULTS AND PUBLICATION POLICY

The results of the research will be disseminated to the global audience through presentations in conferences organized by World Health Organization (WHO),

International Health Economics Association (iHEA), and American Public Health

Association (APHA) etc.

The results will also be communicated with the policy makers and program managers of TB programs in Bangladesh and abroad. Scientific research articles will be prepared after analyzing the data and will be published in reputed peer-reviewed journal. The Principal Investigator of the study, Mohammad Rifat Haider, will take lead in analyzing the data and writing the article and will be the first author. The supervisor of



PhD Study, M Mahmud Khan, PhD, and the mentor, Zaina P. Qureshi, PhD, committee members James W. Hardin, PhD and Md. Abdul Hamid Salim, MBBS will also be coauthors for these studies.

Contribution of other contributors in preparing the papers will also be acknowledged. The sponsorship of TDR, WHO will be acknowledged by quoting: "This investigation received financial support from TDR, the Special Programme for Research and Training in Tropical Diseases, co-sponsored by UNICEF, UNDP, the World Bank and WHO".



CHAPTER 4

STUDY I

4.1 ECONOMIC BURDEN OF TUBERCULOSIS IN BANGLADESH¹





Abstract

Background: Tuberculosis (TB) is major scourge for human-kind and causes profound economic burden. Bangladesh is a high burden TB country by which 12% of its annual deaths are caused and 362,000 people are infected by TB. This study estimates the economic burden of TB on the afflicted Bangladeshi population.

Methods: Based on McIntyre's framework on economic consequences of illness, this study collects direct and indirect cost for TB care data from 1,000 drug sensitive TB (DS-TB) and 145 multi-drug resistant (MDR-TB) patients from all over Bangladesh. Provider cost for TB care was also collected from the health facilities. Costs for DS-TB and MDR-TB patients were estimated using a Generalized Linear Model and summed up with per patient provider level costs to get the total costs per TB patients.

Results: Mean age of DS-TB patients under the study was 45.2 years while mean age of MDR-TB patients were 35.5 years. In aggregate, DS-TB patients incurred total average costs of BDT 21,235 (USD 265) for TB illness; while MDR-TB patients' average costs were BDT 34,975 (USD 437). Including provider costs for each patient (USD 9 for DSTB and USD 2,006 for MDR-TB patients) total average costs for each DS-TB patient was BDT 22,003 (USD 275) and for each MDR-TB patient was BDT 1,95,449 (USD 2443).

Assuming 57% case notification rate, the actual costs for treating TB patients in 2015 was USD 55.6 million. If all DS-TB patients were treated the cost would have been 1



billion USD. For MDR-TB treatment, total cost was USD 12.5 million; treating all MDR-TB patients would have costed USD 23 million.

Conclusions: Results show that DS-TB patients incurred about 50% of their household annual income for treatment while that goes up to 66% for the MDR-TB patients. Prediagnosis cost constitutes about 63% of total costs for DS-TB patients and 42% of MDR-TB patient costs. This figures show the significant economic burden posed by TB and early diagnosis of the disease can reduce the burden in great extent.

Keywords: Economic Burden, Tuberculosis, Bangladesh, Patient Costs, Provider Costs



Background

Tuberculosis is an ancient disease and has claimed more life than any other microbial pathogens in human history (Daniel, 2006). Despite having effective treatment for TB for more than half a century and an effective vaccine for a century, TB still kills more people now than it ever has in the history of the world (McMillen, 2015). It is the human behavior, non-compliance to the relatively long regimen of drugs that provides the bacteria with opportunity of growing resistance against the anti-TB drugs. Tuberculosis is also a disease of poverty, that means the poor and congested living conditions facilitate the bacteria to strive (Davies, 2003, Gandy et al., 2002). Slow progress in control of a preventable and curable disease over last two decades calls for shift our focus from biomedical research of inventing new drugs with shorter regimen to community and patient-driven approach where a paradigm shift is urged for (Stop TB, 2015).

In 2015, Tuberculosis (TB) ranked 18th among the highest burden diseases globally and it constituted 47% of the global burden attributable to communicable, maternal, neonatal, and nutritional disorders (Kassebaum et al., 2016). In 2015, 10 million new cases of TB were reported and almost two million people died from TB worldwide (World Health Organization (WHO), 2016a). In 2015, TB became the top infectious disease killer by claiming 1.1 million lives by matching the death tolls by HIV/AIDS (Kassebaum et al., 2016).

Almost 85% of all new cases of Drug Sensitive TB (DS-TB) and multi-drug resistant TB (MDR-TB) occur in 30 high burden TB countries and Bangladesh is one them (World Health Organization (WHO), 2015b). In 2015, 362,000 Bangladeshis developed TB and 73,000 died from it. TB accounted for 12% of all deaths (609,800) that



occurred in 2015 in Bangladesh (Institute for Health Metrics and Evaluation (IHME), 2016).

The economic burden of TB in Bangladesh is a great concern, since it affects a sizable number of people each year and causes 12% of the total death. Both disability and death have grave economic implications in the form of lost income to the persons and their families and lost Gross Domestic Product (GDP) for the country. The working age group is more affected by the disease, that also increases the costs associated with the disease (World Health Organization (WHO), 2016b). Besides causing death TB causes significant disability among the afflicted population. Not only are older people more vulnerable to the disease, but their disease is more frequently complicated with adverse drug events which leads to reduced health related quality of life (Negin, Abimbola, & Marais, 2015).

Expensive treatment of the disease also put burden on the patients, families, and the health system of the country. (World Health Organization (WHO), 2016b). Moreover, almost half (43%) of the affected in Bangladesh are not reported under the national registries and go untreated (World Health Organization (WHO), 2016a); this makes the control and elimination of the disease extremely hard and expensive. Emergence of drug resistant strain also contributes in escalating costs because of high death rates, costly treatments, and poor outcomes (Fitzpatrick & Floyd, 2012).

This study intends to assess the patient-level costs for both DSTB and MDR-TB, and the provider-level costs for providing the TB diagnosis and treatment.



Methods

Conceptual Framework

In cost analysis three types of costs are taken into account, direct costs, indirect costs and intangible costs due to illness. Direct costs include healthcare costs (hospital, medication, emergency transportation, outpatient visit charges) and family costs (out-of-pocket payment (OOP), medication, transportation of families etc.). Indirect costs include the opportunity costs of work-loss days, lost productivity/income on the part of both the patients and his/her relatives. Intangible costs can arise from the functional limitations, pains acquired in the process and cannot be quantified and highly subjective in nature (Centers for Disease Control and Prevention (CDC), 2013). Besides these costs, there is another important cost incurred by the households through coping strategies, which includes sale of assets, taking up debt, saving on food or other items, taking a child out of school to care for the patient or taking up another job (Russell, 2004).

The costs of communicable diseases fall on the households in three distinct phases: Pre-diagnosis, During Diagnosis/Pre-treatment, and During Treatment. McIntyre et al. (McIntyre et al., 2006) provided a flow-chart on economic consequences of TB illness and payment for healthcare (Figure 2.2). According to McIntyre there are four stages of tuberculosis treatment, e.g., illness experience, treatment seeking behavior, economic consequences, and coping strategies and social resources. In preliminary stages, perception of illness and treatment seeking can be affected by the economic status of the person. If the patient with TB does not seek treatment then only indirect costs are incurred in form of lost productivity, while if the patient goes for treatment then direct



costs also incurred on top of indirect costs. Indirect costs further subdivided into productivity loss of the patient and the caregiver, whereas direct cost is categorized into financial costs of healthcare (services and medicines) and other financial costs.

Based on the conceptual model of McIntyre et al., (McIntyre et al., 2006) Laokri et al. (Laokri et al., 2014) provided an extended conceptual framework incorporating intangible costs, e.g., pain and suffering, and social stigma; elaborating coping costs and social burdens due to illness; and includes societal economic loss along with illness poverty trap ensued due to illness (Figure 2.3). This extended model also includes forgone activities of the informal caregivers as well as those of guardians accompanying the TB patients for treatment. These foregone activities culminate into labor substitution, withdrawal of children from school, and informal caregiving activities of the family members. Guardians lose time and income. Direct costs have been subdivided into subsidized healthcare costs, non-subsidized healthcare costs, and non-healthcare costs. Non-subsidized healthcare costs and non-healthcare costs along with income loss of the guardian can result in (1) Financial resource mobilization, e.g., borrowing, selling assets, pledging, extra-earnings etc.; (2) Resource reallocation, e.g., dissaving, budget cuts, deprivation, delayed investment etc. Non-subsidized healthcare costs and non-healthcare costs can also lead to erratic care pathways which include redundant care visits, alternative care seeking, diagnosis and treatment delays, and care interruption.

Intangible costs like pain and suffering and social stigma can impose social consequences like (1) Low awareness: low awareness of disease, denial of illness status, bad living conditions, fear of losing position, social isolation, lack of family support,



patient-related delays; (2) Social exclusion: exclusion from services including public healthcare services, from income and from participation.

Between McIntyre's and Laokri's conceptual frameworks, both of which are built upon the framework proposed by Russell (2004), simpler McIntyre's framework will be adopted for this study. Both frameworks are more or less similar, while Laokri's one is more detailed and includes pain and suffering, and social stigma. It is difficult to assign monetary value to intangible costs like pain and stigma. On the other hand, these intangible phenomena affect the quality of life of the patients. In our adopted quality of life measurement tool pain and stigma have included to offset their absence in the costs estimation. It also prevents double counting; once included in costs and again take into account while measuring quality of life.

Study Design

The study follows a stratified random sampling method. From the 64 districts of Bangladesh nine districts from the eight divisions (at least one from each division) were selected based on the high and low burden of TB cases. Then from each district two upazilas (sub-districts) were selected randomly. From the registry of the Directly Observed Treatment-Short-course (DOTS) center of the UHCs of these eighteen sub-districts lists of TB patients currently undergoing treatment or recently finished were collected. Following is the list of all districts and sub-districts covered under the study. From each sub-district 50 DS-TB patients were selected randomly for interview.

For assessing economic burden of MDR-TB patients and the economic evaluation of comparator MDR-TB control programs about 168 MDR-TB patients was selected purposively.



Study Participants

The inclusion criteria for the study participants were older than 18 years of age, suffering or recently suffered from pulmonary TB (DS-TB/MDR-TB), and undergoing treatment or finished treatment within the previous 6 months.

Sample Size

The sample size for estimation of the costs incurred by DS and MDR-TB treatment will be calculated using the following formula:

$$n = \left(\frac{Z * SD}{d}\right)^2$$

Where,

Z= 1.96, the right-tail quantile value of a standard normal variable Z at $\,^{lpha}$ =0.05 d= margin of error

SD= Standard Deviation of the mean costs

For DS TB in a recent study we find that the patient cost for the treatment was \$224 (Tanzania). This cost does not include the cost for providing the treatment, i.e., health care delivery costs. If we guess that the total cost would be \$400 including all other costs. We also assume that the standard deviation would be \$400 and with the margin of error of \$50 we get the sample size at 5% significance level is 246. However, 1,000 DS-TB patients were interviewed under the study; out of them 404 undergone treatment under CHW model and 598 got treatment under CM model.

NTP PMDT Expansion plan, Bangladesh (2013 - 2017) estimates the MDR-TB treatment cost is \$6000. If we take the similar figure as the standard deviation and \$1200 as the margin of error, we get the sample size for MDR-TB patients at 5% significance



level is 96. Under this study, 145 MDR-TB patients were interviewed; 58 patients were treated with 9 month regimen while 87 patients were under 20-24 month regimen.

Providers

Under the purview of the study 16 DOTs center managers and 3 representatives from MDR-TB care providing hospitals were interviewed for collecting cost of providing treatment to DS-TB patients. In each of the TB treatment facilities the facility manger was interviewed using a pre-set questionnaire.

Variables

Patient Costs	Provider Costs	Patients' Socio-economic	
 Direct costs Medical costs Non-medical costs Indirect (opportunity) costs Coping costs Costs made in the facility that were not obligatory to get the diagnosis and treatment (i.e., costs of food). Other costs: (in)direct costs made by or for accompanying persons (attendants) 	 Prevention and Promotion Costs Contact Tracing Costs Diagnosis Costs Drugs Costs Treatment Costs Other TB activities Costs Human Resources Costs Capital Costs 	 Characteristics Age Sex Education Occupation Religion Current health status Location (Rural, urban, urban slum) Wealth (asset) index Type of TB patient (New, relapse, failure, transfer in) Household income/number of earning members 	



Data Analysis

Data analysis was performed by Stata 14.2 (StataCorp, 2015). Descriptive statistics like means, standard deviation, frequency, and percentages will be reported. Multivariable regression analysis will be performed for finding the important variables for treatment and access costs.

Measurement of Average Total Direct Costs

Average total direct costs will be measured by combining all the out-of-pocket medical and non-medical costs for TB treatment. Costs for DS-TB and MDR-TB patients will be separately calculated. These costs include drugs, diagnostic tests, fees, consultation fees, food costs, travel costs, accompanying person costs. That means direct costs include all costs incurred directly out-of-pocket for the treatment of TB patient.

Costs incurred by each patient, e.g., both DS-TB and MDR-TB, have been calculated. Patients were interviewed on each and every visits they made to any type of provider for the TB illness were tried to track down. For each visit the patients reported their (including their accompanying persons) incurred direct costs.

Measurement of Average Total Indirect Costs

Average total indirect costs will be measured by foregone income due to inability to do normal daily activities. These activities can be formal or informal. Valuation of productivity losses has been done using per capita gross domestic product (GDP) of Bangladesh (USD 1342/year that means 294 BDT/day). Valuation using GDP is preferred because this approach gives same weight to rich and poor people's income.

Regression Analysis

Multivariable regression analysis will be performed using socio-demographic variables as the predictors. The cost data for both DS-TB and MDR-TB patients were



skewed to the right with cost amounts concentrating near zero values. However, the costs are not exactly zeros because all TB patients incurred some costs. Given the distribution of the costs data, normality assumption for OLS regression has been checked using histogram and normal probability plot, and heteroscedasticity was tested using the Breusch-Pagan test. We found that the normality assumption was violated and there was evidence of heteroscedasticity (unequal variance).

Although log transformation of the cost variable is a common way to deal with this skewness, it still suffers from the problem of heteroscedasticity and the transformation and retransformation would lead to biased estimate of cost. Therefore, estimation of a Generalized Linear Model (GLM) has been preferred because it is particularly helpful in avoiding the log retransformation problem and it does not require the normality assumption to hold true. For GLM one has to specify the correct link and variance (family) function. The modified Park test was used for selecting family, while the Pregibon link test (checking linearity of response on scale of estimation) was used to assess the choice of link function, The large sample Pearson correlation test and Modified Hosmer-Lemeshow test (checks for systematic bias in fit on raw scale) were used for specifying link. Based on these tests, a GLM model with identity link and inverse Gaussian family, in which variance is proportional to the cube of mean, was found best suited for both DS-TB and MDR-TB cost models.

We can specify the model as below:

$$c_{i,j} = \beta_0 + \beta_1 x_{i,j1} + \beta_2 x_{i,j2} + \sum_{k=0}^{n} \alpha_k z_{i,j}$$



Here, $c_{i,j}$ is the patient cost patient (i=1 to N) and suffering from DS-TB or MDR-TB (j=0 or 1). $x_{i,j1}$, $x_{i,j2}$ are dummy variables for treatment through CHW model and CM model for DS-TB treatment costs respectively; while these two represents 20-24 month regimen and 9 month regimen for MDR-TB treatment model, $z_{i,j}$ represents different covariates to control for across individuals.

Provider Costs

These costs can be called as health system costs as it contains the costs from the health system perspective. It includes capital costs, personnel costs, drugs cost, laboratory costs, and programmatic costs. Total cost for each facility has been summed up and then was divided by the number of total TB patients served by each facilities to come up with the per patient costs incurred by the provider. In other words, the health system cost will be described as the cost from the provider side to treat each patient under each type of treatment modality.

Estimating Economic Burden of TB in Bangladesh

Finally, the total economic burden of TB in Bangladesh has been calculated using the TB prevalence data. This data was obtained from Global Tuberculosis Report 2015 published by World Health Organization (WHO). Recently a TB prevalence survey has been conducted by Institute of Epidemiology, Disease Control, and Research (IEDCR) in Bangladesh. The preliminary findings of that survey found that the prevalence rate was lower (295 per 100,000 population) than WHO estimate of 362 per 100,000 in 2015. This rate has also been used to estimate a comparative economic burden of TB in Bangladesh.



Results

Patient Characteristics

Mean age of DS-TB patients under the study was 45.2 years while mean age of MDR-TB patients were 35.5 years. Most MDR-TB patients were under the age of 45 years, whereas DS-TB patients were more dispersed among the age groups. In both DS-TB and MDR-TB samples majority of the patients were male, had no education, did informal work or did not work before occurrence of TB, of Islamic faith, and resided in rural areas. Most of the DS-TB patients were newly diagnosed, while most MDR-TB patients suffered from relapse or treatment failure. In the same vein, most of the DS-TB patients had no previous history of TB treatment, but almost two-third of the MDR-TB patients had previous history of TB treatment and almost one-fifth of the total sample did not complete the treatment. Since, wealth index was calculated separately among two samples, one-fifth (20%) of each population belonged to each quintile except the poorest quintile contained more (22.4%) patient than the poorest one (17.6%) (Table 4.1).

Table 4.1 DSTB and MDR-TB Patient characteristics under the study

Characteristics	DS-TB Patients	MDR-TB Patients 145	
N	1,000		
	% (n)	%(n)	
Age			
18-25 Years	14.9 (149)	29.0 (42)	
26-35 Years	18.4 (184)	33.8 (49)	
36-45 Years	17.5 (175)	13.8 (20)	
46-55 Years	20.6 (206)	9.6 (14)	
56-65 Years	17.5 (175)	9.0 (13)	
66+ Years	11.1 (111)	4.8 (7)	
Sex			
Female	37.5 (375)	42.1 (61)	
Male	62.5 (625)	57.9 (84)	
Education			
No Education	42.1 (421)	33.1 (48)	
Primary	31.7 (317)	31.0 (45)	



Secondary	20.6 (206)	29.0 (42)
Higher Secondary and	5.6 (56)	6.9 (10)
Higher		
Occupation before TB		
Formal	11.6 (116)	21.4 (31)
Agriculture or Household	13.4 (134)	8.3 (12)
Jobs		
Informal	37.6 (376)	33.1 (48)
Did not work	37.4 (374)	37.2 (54)
Religion		
Other	10.6 (106)	7.6 (11)
Islam	89.4 (894)	92.4 (134)
Current Health Status		
(VAS)		
050	23.2 (232)	34.5 (50)
51-65	16.9 (169)	11.0 (16)
66-80	37.9 (379)	30.3 (44)
81-100	22.0 (220)	24.1 (35)
Location		
Urban	18.2 (182)	37.9 (55)
Rural	77.2 (772)	47.6 (69)
Urban Slum	4.6 (46)	14.5 (21)
Type of TB Patient*		
New	94.2 (909)	4.6 (6)
Relapse/Failure	4.9 (47)	87.7 (114)
Loss to Follow Up	0.9 (9)	7.7 (10)
Household Income Before TB	15,282 [24,916]	17,355 [11,307]
Previous TB Treatment		
No Previous Treatment	94.0 (940)	35.2 (51)
Not Completed	1.0 (10)	17.9 (26)
Completed	5.0 (50)	46.9 (68)
DS-TB Program Model		
Community Health Worker	40.3 (404)	-
Community Member	59.7 (598)	-
MDR-TB Program		
Regimen		
9 Month Regimen	=	40.0 (58)
24 Month Regimen	-	60.0 (87)
Wealth Index		
Poorest	22.4 (224)	20.0 (29)
Poorer	17.6 (176)	20.0 (29)
Middle	20.0 (200)	20.0 (29)
Richer	20.0 (200)	20.0 (29)
Richest	20.0 (200)	20.0 (29)



Total	100 (1,000)	100.0 (145)
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^{* 15} missing for MDR-TB

Patient Level Average Costs

Direct Costs

In Table 4.2, average direct costs borne by DS-TB and MDR-TB patients have been illustrated. Direct costs include all the out-of-pocket medical and non-medical costs for TB treatment. Total average direct costs for DS-TB patients were BDT 20,154 (USD 252) and for MDR-TB the amount was BDT 30,858 (USD 386). The highest costs were incurred by the patients during the pre-diagnosis phase of the illness. DS-TB patients spent BDT 13,287 (USD 166) and MDR-TB patients incurred BDT 14844 (USD 186) before the diagnosis of TB disease was confirmed. TB diagnosis was costly for DS-TB patients (BDT 1,107; USD 14) than MDR-TB patients (BDT 685; USD 9). Hospital costs were way higher for the MDR-TB patients (BDT 7,669; USD 96) than DS-TB patients (BDT 2,515; USD 31). Additional food costs were more or less same for both DS-TB and MDR-TB patients. Accompanying person's costs was higher for MDR-TB patients (BDT 2,114.70; USD 26) than DS-TB patients (BDT 380; USD 5). Costs for side-effects of TB drugs were higher for the MDR-TB patients (BDT 1,647; USD 21) than DS-TB patients (BDT 435; USD 5). Relocation costs was incurred by only the MDR-TB patients during hospitalization (initial incentive phase of treatment). On an average, the relocation cost was BDT 341.24/USD 4.

Indirect Costs

In table 4.2, average indirect costs borne by DS-TB and MDR-TB patients have also been shown. Indirect costs include the income loss by the patients as well as their accompanying persons. MDR-TB patients incurred more indirect costs (BDT 1,523; USD



19) than DS-TB patients (BDT 407; USD 5). Similarly, the persons accompanied MDR-TB patients lost more income (BDT 2,594; USD 32) than persons accompanied DS-TB patients (BDT 674; USD 8). In total, MDR-TB patients incurred more indirect costs (BDT 4,117; USD 51) than DS-TB patients (BDT 1,081; USD 14).

Total Average Patient Level Costs

In aggregate, DS-TB patients incurred total average costs of BDT 21,235 (USD 265) for TB illness; while MDR-TB patients' average costs were BDT 34,975 (USD 437).

Table 4.2 Patient Level Average Costs (BDT)

Costs	DS-TB Patients	MDR-TB Patients
N	1,000	145
	Mean [SD]	Mean [SD]
Direct Costs		
Before Diagnosis Costs	13,287.16 [55002.53]	14,844.23 [25589.24]
TB Diagnosis Costs	1,106.69 [5725.75]	684.63 [2177.10]
Follow-up Costs	172.16 [640.63]	877.17 [1825.46]
Drug Collection Costs	14.6 [195.91]	1.17 [6.18]
Hospital Costs	2,515.23 [14813.86]	7,669.40 [10521.34]
Additional Food Costs	2,244.01 [1915.92]	2,678.39 [1595.63]
Accompanying Person	380.03 [739.85]	2,114.70 [5309.25]
Costs		
MDR-TB Relocation Costs	-	341.24 [1386.08]
Drug Side-effects Costs	4,34.58 [2998.73]	1,647.28 [5463.01]
Sub-Total (Direct Costs)	20,154.43 [60241.68]	30,858.22 [39964.31]
Indirect Costs		
Patient Opportunity Costs	407.07 [378.39]	1,522.86 [942.80]
(Income Loss)		
Accompanying Person	673.57 [1952.26]	2,593.91 [6765.72]
Opportunity Costs		
Sub-Total (Indirect Costs)	1,080.64 [2004.40]	4,116.76 [6887.64]
Total Costs	21,235.10 [60841.03]	34,974.098 [43635.95]

Provider Level Average Costs

In Table 4.3, facility level data was used to illustrate the average per person health system costs to provide TB treatment. DS-TB treatment facilities did not report any costs



for prevention and promotional activities, e.g., contact tracing, health promotional activities, vaccination activities etc. In treating each DS-TB patients the health system incurred only BDT 768 (USD 9.60), for MDR-TB patients the cost rose to BDT 160,474 (USD 2006). Drugs (BDT 59276; USD 741), human resources (BDT 51826; USD 648), and diagnostic (BDT 35554; USD 444) were the highest cost incurring areas for MDR-TB patients. Similarly, for DS-TB patients the highest cost-incurring areas were drugs (BDT 439; USD 5), diagnostics cost (BDT 184; USD 2), and human resources (BDT 70; USD 1).

Table 4.3 Provider Level Average Costs (Per Patient)

Costs	DS-TB Facilities	
N	5648	576
	BDT	BDT
Prevention and Promotion	0.00	524.78
Costs	0.00	324.78
Diagnostic Costs	183.51	35,554.08
Drug Costs	438.79	59,275.55
Training Costs	10.63	1,014.58
Meeting Costs	1.34	874.64
Incentive Payment	31.28	3,100.00
Human Resources Costs	69.86	51,825.73
Capital Costs	29.37	8,211.78
Other Costs	2.92	92.73
Total Costs (BDT)	767.69	16,0473.86
Total Costs (USD)	9.60	2,005.92

Per Patient Total Average Costs

Average per patient total costs including patient and provider level costs have been shown in Table 4.4. On an average, each DS-TB patient incurs BDT 22,003 (USD 275) and each MDR-TB patient incurs BDT 1,95,449 (USD 2443).

Table 4.4 Per Patient total average cost (Including patient and provider level costs)

Costs	DS-TB	MDR-TB
	BDT	BDT



Patient Level Costs	21,235.10	27,809.67
(BDT)		
Provider Level Costs (BDT)	767.69	16,0473.86
Total Costs (BDT)	22,002.79	19,5448.84
Total Costs (USD)	275.03	2,443.11

Bivariate Analysis of Patient Level Cost Data

Table 4.5 shows the results from bivariate analysis of patient level cost for both DS-TB and MDR-TB patients. For DS-TB costs only mean total costs of different wealth quintiles were significantly different. In case of MDR-TB patients, categories of previous history of TB treatment and MDR-TB regimen had significantly different mean total costs.

Table 4.5 Bivariate Analysis of Patient Level Cost for DS-TB and MDR-TB patients

Characteristics	DS-TB Patients		MDI	R-TB Patier	nts	
N		1,000			145	
	Mean	Median	p-Value	Mean	Median	p-
						Value
Age			0.056^{a}			0.438
18-25 Years	15,178	9,529	0.813	43,3348	26,616	0.269
			(kwallis)			
26-35 Years	17,788	8,806		31,536	18,732	
36-45 Years	23,763	8430		23,988	16,135	
46-55 Years	22,795	8,887		25,703	14,609	
56-65 Years	15,993	8,802		47,734	19,504	
66+ Years	36,463	9,378		35,049	16,476	
Sex			0.206			0.179
Female	18,092	8,802	0.812	40,703	24,217	0.239
Male	23,121	8,849		30,816	17,816	
Education			0.107			0.289
No Education	16,888	7,130	0.000	25,946	14,923	0.075
Primary	21,531	8,646		36,153	27,196	
Secondary	26,083	12,830		40,912	18,467	
Higher Secondary	34,404	21,196		48,076	25,419	
and higher						
Occupation			0.079			0.097
Did not work	2320	10,095	0.001	46,270	26,431	0.133
Formal	31,086	10,478		32,054	26,898	
Agriculture or	22,693	7,552		30,992	16,102	
Household Jobs						



Informal	15,602	7,533		25,150	16,201	
Religion		,	0.168	,	,	0.210
Other	13,532	7,234	0.110	19,083	14,428	0.058
Islam	22,148	9,194		36,280	18,665	
Location			0.205			0.141
Urban	26,814	10,416	0.0015	44,143	29,782	0.003
Rural	20,585	8,494		29,566	14,127	
Urban Slum	10,071	8,319		28,737	17,709	
Previous TB			0.108			0.041
Treatment						
No	20,453	8,515	0.001	45,025	35,200	0.004
Yes	33,484	11,988	0.021	29,523	16,377	
DS-TB Program			0.069			
Model						
Community Health	16,971	7,574	0.000	-	-	
Worker						
Community	24,102	9,519		-	-	
Member						
MDR-TB						0.006
Program Regimen						
9 Month Regimen	-	-		22,975	13,406	0.000
20-24 Month	-	-		42,975	28,261	
Regimen						
Wealth Index			0.015			0.120
Poorest	12,375	7,267	0.000	30,846	16,542	0.071
Poorer	27,450	7,072		30,180	16,476	
Middle	18,483	8,340		27,372	16,147	
Richer	19,175	9,972		32,204	21,801	
Richest	30,776	12,826		54,273	30,770	
Comorbidity			0.238			0.927
No	19,610	8,180	0.006	35,146	18,012	0.693
Yes	24,404	9,752		34,319	20,607	

^a p-values are obtained from univariate analysis

Multivariable Analysis of Patient Level Cost Data

Two separate GLMs were estimated with the patient level cost data for DS-TB and MDR-TB patients (Table 4.6). The modified Park test showed that best the GLM model for DS-TB costs belonged utilized the inverse Gaussian family, but for MDR-TB costs the Gamma family was best suited. The log link was best for both models. Results from the GLM post-estimation for selecting best model are shown in table 4.7.



Results show that education is a significant cost driver for both DS-TB and MDR-TB patients. With higher education the costs tend to become higher in both cases.

Improved health status (measured by VAS) was associated with decreased expense on TB care for DS-TB patients. This effect of better health status did not hold true for MDR-TB patients. Similarly, while location of the patient had no effect in case of MDR-TB patients, DS-TB patients resided in the urban slums incurred less cost than their urban counterparts.

Previous TB treatment had a negative effect on MDR-TB patients cost, but it had no effect on costs of DS-TB patients. Community member model incurred more cost than the community health care model in case of DS-TB treatment. On the other hand, 20-24 month regimen incurred more cost on MDR-TB patients than 9 month regimen.

Wealth has no effect on cost of MDR-TB patients, while DS-TB patients belonged to middle, richer, richest quintiles incurred more costs than the poorest patients.

Table 4.6 Multivariable Analysis of Patient Level Cost with Generalized Linear Model

Characteristics	DS-TB Patients		MDR-TB Patients	
N	1,0	000	14	5
	Log Lin	k Inverse	Log Link Gar	nma Family
	Gaussia	n Family		
	Coefficient	95% CI	Coefficient	95% CI
Age				
18-25 Years	Ref.	-	Ref.	-
26-35 Years	0.25	-0.19-0.68	0.05	-0.39-0.48
36-45 Years	0.30	-0.15-0.76	-0.30	-0.79-0.23
46-55 Years	0.42	-0.03-0.86	0.20	-0.46-0.87
56-65 Years	0.36	-0.14-0.85	0.28	-0.34-0.90
66+ Years	0.41	-0.18-1.00	0.16	-0.68-0.99
Sex				
Female	Ref.	-	Ref.	-
Male	0.18	-0.15-0.52	-0.10	-0.48-0.27
Education				
No Education	Ref.	-	Ref.	-
Primary	0.39*	0.08-0.70	0.57**	0.15-0.99



Secondary	0.45*	0.07-0.84	0.59**	0.15-1.02
Higher Secondary and	0.93*	0.07-1.80	0.30	-0.41-1.01
higher				
Occupation				
Did not work	Ref.	-	Ref.	-
Formal	0.09	-0.42-0.60	-0.33	-0.80-0.14
Agriculture or Household	0.02	-0.50-0.54	0.28	-0.44-1.00
Jobs				
Informal	-0.31	-0.680.07	-0.49	-0.92-0.07
Religion				
Other	Ref.		Ref.	-
Islam	0.28	-0.07-0.62	0.43	-0.20-1.05
Current Health Status	-0.01*	-0.020.002	-0.004	-0.01-0.005
(VAS)				
Location				
Urban	Ref.	-	Ref.	-
Rural	-0.02	-0.46-0.41	0.42	-0.19-1.04
Urban Slum	-1.07**	-1.740.40	-0.004	-0.58-0.58
Previous TB Treatment				
No	Ref.	-	Ref.	-
Yes	0.40	-0.23-1.03	-0.49**	-0.83
				0.15
DS-TB Program Model				
Community Health	Ref.	-	-	-
Worker				
Community Member	0.43**	0.18-0.69	-	-
MDR-TB Program				
Regimen				
9 Month Regimen	-	-	Ref.	-
20-24 Month Regimen	-	-	1.17***	0.60- 1.74
Wealth Index				
Poorest	Ref.	-	Ref.	-
Poorer	0.15	-0.18-0.48	-0.03	-0.52-0.46
Middle	0.38*	0.04-0.72	-0.37	-0.93-0.20
Richer	0.41*	0.04-0.78	-0.32	-0.95-0.31
Richest	0.54*	0.03-1.05	0.10	-0.63- 0.82
Comorbidity	_			
No	Ref.	-	Ref.	-
Yes	0.15	-0.13-0.44	0.02	-0.40-0.44
AIC		9.30	22.	
BIC		58.78	-515	
Log Likelihood	-131	159.45	-1636.64	

Table 4.7 Results from GLM post-estimation for selecting best model



Test	DS-TB	DS-TB Patients		3 Patients
	Test	Decision	Test	Decision
	Statistic		Statistic	
	p-Value		p-Value	
Modified Park Test	0.27	Inverse	0.15	Gamma
		Gaussian		Family
		Family		
Pearson Correlation	0.83	Log Link	0.15	Log Link
Test				
Pregibon Link Test	0.21	Log Link	0.12	Log Link
Modified Hosmer-	0.57	Log Link	0.66	Log Link
Lemeshow Test				

Economic Burden of TB Care in Bangladesh

Based on the World TB Report 2016, total 209,438 DS-TB patients were under treatment in Bangladesh. Assuming 57% case notification rate, we get the actual number of TB patients in Bangladesh in 2015 was 367,435. Therefore, the actual costs incurred by Bangladesh have been calculated as USD 55.6 million. Whereas, if all DS-TB patients were treated the cost would have been 1 billion USD.

For MDR-TB treatment, total USD 12.5 million was incurred in Bangladesh in 2015. If all MDR-TB patients were treated the total cost would have been USD 23 million (Table 4.8).

Table 4.8 Economic Burden of TB care in Bangladesh in 2015

Type of TB Patients	Average cost	Total TB patients under	Total actual costs for TB Care		Total TB patients in 2015	require	l costs d for TB are
		treatment in 2015	BDT	USD		BDT	USD
DS-TB	22,003	209,438	4.45 Billion	55.59 Million	367,435	8.08 Billion	101.06 Million
MDR- TB	195,449	5,100	0.18 Billion	12.46 Million	9,700	1.90 Billion	23.70 Million
Total			4.63 Billion	68.05 Million		9.98 Billion	124.76 Million



Discussions

Study results show that average per DS-TB patient costs were BDT 21,235 (USD 265), which is almost similar to the findings of a recent study (BDT 20,720; USD 224) (Gospodarevskaya et al., 2014). However, the study reported only treatment cost and excluded the pre-diagnosis cost incurred by TB patients. Excluding the whooping \$166 for pre-diagnosis cost, the actual treatment level costs for DS-TB in Bangladesh from our study comes to \$100. The main difference between the study and our result stems from the estimation of productivity losses by patients and guardians. That study used the household level income but we used per capita GDP as the basis for calculation of lost productivity. Given our sample size of 1,000 in comparison to their 96, we can confidently claim that our result is more authentic and does not differ significantly between different types of providers.

Both types of delivery modalities for DS-TB patients in Bangladesh, e.g., CHW and CM models, ensure patients can get their medicines at their doorsteps or very near to their houses supervised by community health workers and community members. MDR-TB patients also get their medicines from assigned community DOTS providers. Yet, TB patients incur considerable expenditure for TB treatment. From the results we can see that a significant portion of the expenditure is incurred before TB diagnosis. It constitutes about 63% of total costs for DS-TB patients and 42% of MDR-TB patient costs. This shows that once the patient is diagnosed and under the treatment stream, patient level costs and health system costs constitute only one-third of the total cost. Delay in TB diagnosis is the major cost driver for the patients. Patients may visit number of providers from informal to formal, even are hospitalized in the course, and incur a great loss in



terms of medical and non-medical costs. This finding is similar to other studies where pre-diagnosis cost of TB treatment constituted more than half of the total costs (Tanimura et al., 2014). High pre-diagnosis cost of TB care is a phenomenon ubiquitous in LMICs, e.g., in Malawi the patients incur a significant pre-diagnosis cost which offset the free TB care and make the TB treatment unaffordable (Kemp, Mann, Simwaka, Salaniponi, & Squire, 2007).

The study results slightly differ with the findings from a systematic review done by Laurence et al. The average provider level cost was \$273 in that study (Laurence et al., 2015) in comparison to \$9 per DS-TB patient in ours, since they included hospitalization cost (\$215). But in Bangladesh, the DS-TB treatment protocol does not require hospitalization, therefore, our results do not include any hospitalization costs and that makes the two estimates very close. For MDR-TB costs, provider level costs were calculated \$6313 in LMICs and \$1218 in LICs, and patient level costs were calculated \$1616 total direct costs in LMICs and \$1662 total costs in LICs (Laurence et al., 2015). In our study we found different results- provider level costs were \$2006 and patient level costs were \$437. It may be due to their estimation of life-time productivity loss, which we confined within the period of illness only.

We found that health system costs for providing treatment costs is nominal (\$9 per patient) in Bangladesh. But the high number of patients make the total burden high. For treating DS-TB patients Bangladesh incurred 55.6 million USD in 2015, and for MDR-TB patients USD 12.5 million. In total, expenditure for TB treatment in Bangladesh was 68 million USD. In Bangladesh, total health expenditure was 325,094 million BDT (4,063 million USD) in 2012. Therefore, in 2015 the amount would have



been 4,545 million USD. So, TB care expenditure constitutes about 1.67% of total healthcare expenditure in Bangladesh. This figure does not look so ominous, but the potential of TB treatment is very high in terms of future benefits. In a recent study commissioned by the Copenhagen Consensus Group, Vassal showed that TB treatment was ranked first among all priorities because its huge potential in future befits. If one taka is spent for TB treatment, the economic return would be in the range of 29 to over 162 BDT (Anna Vassal, 2016).

TB also poses great financial hardship on the afflicted population. Often times poor people are the sufferer and their economic condition does not help them either. In our study results we see that DS-TB patients incurred about 50% of their household annual income for TB treatment while that goes up to 66% for the MDR-TB patients. This catastrophic health expenditure is multiplied in severity due to the absence of any health insurance or other healthcare financing mechanism in Bangladesh (Nazmul, Abul Quasem, Howlader, & Kabir, 2015). People tend to resort to sale of assets or savings and borrowing with or without interest to cope with this catastrophic cost, which, in the long run, make the household poorer (Khan, Ahmed, & Evans, 2017; Rahman, Gilmour, Saito, Sultana, & Shibuya, 2013).

Data collected from the DS-TB health facilities show that no cost was incurred for health promotional and preventive activities in past year. That means there was no such activities in place. But contact tracing, promotional activities like making people aware of the signs and symptoms of TB illness, informing the people on the treatment availability and the successful cure is possible upon completion of treatment, the place where TB treatment is available etc. are deemed to be instrumental in combating TB in LMICs like



Bangladesh. There are some top-down promotional activities done from the central level, but that may not reach the grass-root level. Different means of behavioral change communication should be introduced; otherwise the ambitious target of reducing TB deaths by 95% and curbing new cases by 90% from 2015 to 2035 (World Health Organization (WHO), 2017) would not be achieved.

Urban slum is another hot spot which can serve as the new foci of TB in Bangladesh. With highest urbanization rate in the world (6.5%) the capital Dhaka city experiencing a burgeoning urban slum population. This population often resides in the most inhuman condition and lack basic needs like health, education, and proper housing. This close proximity of people (200,000.people in 1 square kilometer in Bangladeshi slums (Angeles et al., 2009)) and poor living conditions facilitate transmission of the TB bacillus and containment of the disease makes so challenging. We find from our results that slum dwellers can spend much lower than their urban counterparts and still that lower spending leaves them as poverty-stricken for rest of their lives due to the long term effect on their income generating potentials.

Conclusions

Since the lion share of the total costs was incurred before diagnosis, it is imperative to strengthen the early diagnosis and treatment of TB disease in Bangladesh. CHW or CM who are involved with the DOTS treatment and act as a DOTS provider should also act as the counselor for the patients and their family members. DOTS supporter should also keep eyes open to find any potential TB cases and refer them to the nearest facility.



Contract tracing can be another useful way for detecting TB patients. Although it was a regular activity in the past, during the survey we did not find any such activities performed nowadays. It is a reminder that we should not move away from the basic prevention techniques of the public health.

Although, treating DS-TB patients is more beneficial in the long-run, proper treatment of MDR-TB patient is very important in further spread of the deadly disease. MDR-TB needs special attention, because of the length of the treatment and potential for loss to follow up and relapse. Therefore, MDR-TB patients should also be given proper treatment and special arrangements, like tracking patients uptake of drugs by novel means, can be thought of.

TB has been cited as the most effective health intervention in terms of costbenefit ratio (Anna Vassal, 2014). It is high time to get all TB patients under treatment and thereby improve the economy of the country. It will also help us to reach the ambitious goal of the WHO End TB strategy by 2035.

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CHAPTER 5

STUDY II

5.1 ECONOMIC EVALUATION OF DRUG-SENSITIVE TUBERCULOSIS (DS-TB)
TREATMENT APPROACHES IN BANGLADESH²

² Haider, M. R. To be submitted.



Abstract

Introduction: Bangladesh is a high burden Tuberculosis (TB) country that experienced 362,000 new TB patients and 73,000 TB deaths in 2015. Drug sensitive TB (DS-TB) is the most prominent type of TB found in Bangladesh and a 6-month drug regimen (2 month intensive and 4 month continuation phase) is prescribed. However, the directly observed treatment short-course (DOTS) differs in delivery through community health workers (CHWs) and community members (CMs). This study compares these two delivery models and conducts a cost-effectiveness analysis.

Methods: The incremental cost-effectiveness ratio (ICER) of treating DS-TB patients, 45 years old on average, through CM versus CHW delivery was compared using a Markov model with life-time horizon (27 years). The measure of effectiveness, Quality adjusted life year (QALY) and cost of treatment was collected from 1,000 MDR-TB patients (598 for CM model and 402 from CHW model) in Bangladesh. Transition probabilities between Markov states were estimated from quarterly outcomes report collected from health facilities and cost and QALY both were discounted at a rate of 3%. Both deterministic and probabilistic sensitivity analyses were conducted in a Monte Carlo Simulation using the R programming language.

Results: Results show that each DS-TB patient under CM treatment model gains 3.61 QALYs with a cost of BDT 131,555. For the DS-TB patients under the CHW model the cost is 81,650 and the QALY gain is 3.12. The Incremental Cost-Effectiveness Ratio (ICER) is 103,454, i.e., the CM model is cost-effective if per QALY gain if willingness-to-pay is set to the per capita GDP of Bangladesh (BDT 107,360 in 2015).



Conclusions: Our study results suggest that a community-based model of DS-TB treatment is cost-effective even with changed costs and utility values in probabilistic sensitivity analysis. Community members as DOTS provider are more capable of reducing stigma related to TB, enhancing patient adherence and thereby reduce costs and increase utility from the treatment. Community members should also be involved in contact tracing and prevention activities to increase the effect of the involvement in TB control.

Keywords: Economic Evaluation, Drug Sensitive Tuberculosis, Cost-effectiveness, Cost, QALY, Community Based Treatment, Bangladesh



Introduction

In 2015, tuberculosis (TB) ranked 18th among the highest burden diseases globally and it constituted 47% of the global burden attributable to communicable, maternal, neonatal, and nutritional disorders (Kassebaum et al., 2016). During that year, 10 million new cases of TB were reported and almost two million people died from TB worldwide (World Health Organization (WHO), 2016a). In the same year, TB became the joint top infectious disease killer by claiming 1.1 million lives and matching the death tolls by HIV/AIDS (Kassebaum et al., 2016).

Almost 85% of all new cases of TB and multi-drug resistant TB (MDR-TB) occur in 30 high burden TB countries including Bangladesh (World Health Organization (WHO), 2015b). According to one estimate in Bangladesh during the whole year of 2015, approximately 362,000 people developed TB and 73,000 died from it. In Bangladesh, TB accounted for 12% of all deaths (609,800) that occurred in 2015 (Institute for Health Metrics and Evaluation (IHME), 2016). Although case notification rate is only 57%, success of the treatment is high (93%) among DS-TB patients. However, the success rate is 75% among MDR-TB patients which signifies how difficult to treat drug resistant strains (World Health Organization (WHO), 2016a). Despite having effective treatment, patient adherence to TB treatment remains poor because of the long duration of the regimen (six months for newly diagnosed cases) and the need for daily dosing. Failure to adhere to the regimen results in MDR-TB (Gandy & Zumla, 2002).

The predominant method of detecting TB at the community level is examining the sputum sample with Acid Fast Bacilli (AFB) technique. In the case of detection of MDR-TB, culture and sensitivity analyses are done in laboratories. Directly Observed



Treatment Short-Course (DOTS) strategy has been implemented in Bangladesh since 1993 and all the Upazila Health Complexes (UHCs) have been brought under the purview of the service from where TB detection and treatment services are given free of cost. The essence of the strategy is the diagnosed TB patient has to go to the facility every day for taking the drugs, thus treatment discontinuity and subsequently MDR-TB cases can be averted (World Health Organization (WHO), 2013d).

The vibrant presence of Non-governmental organizations (NGOs) in the health sector and TB control endeavor compels the National Tuberculosis Control Program (NTP) under Ministry of Health and Family Welfare (MoHFW) of Bangladesh to incorporate them into the public-private partnership (PPP) model of combating TB since 2003. In the recent guideline the role of government and private sector partnership was reiterated again (Guideline). It was also found in different studies that this PPP model in TB control was effective in achieving relatively high case detection (Ullah et al., 2012; Ullah, Newell, Ahmed, Hyder, & Islam, 2006).

For the drug sensitive TB patients, a standard 6-month regimen is followed by all participating NGOs. However, the mode of delivery is different for different NGOs. BRAC has employed Community Health Workers (CHWs) besides the DOTS centers to ensure patient compliance, while Damien Foundation (DF) trained and employed influential community members to help the patients to be adhered to the treatment protocol. Another NGO, Salvation Army Bangladesh, is using drug sellers at the pharmacies as the counselor and drug distributors for the TB patients. Since involving different people, e.g., family members, neighbors, pharmacists falls under common strategy of involving community members.



Using Denver General Hospital data Burman et al. showed that although DOTS is costly at the outset it turns to be cost-effective than Self-administered Therapy (SAT) because of higher cure rates (Burman et al., 1997). The outcome variable for this study was cure rate per cost unit. Using published literature, records, and expert opinions Baltussen et al. showed that DOTS as well as incremental programs like DOTS plus, Full combination of DS-TB and MDR-TB strategies all are cost-effective in terms of DALYs averted per cost unit in high burden TB countries in Africa and South-East Asia (Baltussen et al., 2005).

A recent study shows that shortening of the DS-TB treatment from six-months to four-months remain cost-effective option for Brazil, South Africa, Bangladesh and Tanzania (Gomez et al., 2016). Another study results also support this finding in South Africa (Knight et al., 2015).

Although several studies conducted economic evaluation between different types of treatment model or regimen, economic evaluation between CHW and CM models has not been performed. The two methods of delivering DS-TB care in Bangladesh based on the service area of particular NGOs provide us with the opportunity to evaluate the cost-effectiveness of two methods of delivering DOTS to DS-TB patients. This study aims to conduct an economic evaluation between CHW and CM models of delivering DS-TB care with a societal perspective and Quality Adjusted Life Years (QALY) as the outcome.



Methods

Study Design and Data Sources

The study follows a stratified random sampling method. From the 64 districts of Bangladesh nine districts from the eight divisions (at least one from each division) were selected based on the high and low burden of TB cases. Then from each district two upazilas (sub-districts) will be selected randomly. From the registry of the DOTS center of the UHCs of these eighteen upazilas, lists of TB patients currently undergoing treatment or recently completed treatment will be collected. From each upazila 50 DS-TB patients were selected randomly for interview. The list of all districts and upazilas covered under the study is shown in Table 3.1.

Institutional level data was also collected for assessing the provider level costs associated with TB treatment using a pre-set and pre-tested provider questionnaire. We interviewed the healthcare facility manager to get the annual human resources costs, costs for training, meeting, incentive payments, capital costs, and other costs. Drug costs and diagnostic costs are assumed to be equal for both treatment model since both model follows the same guidelines published by NTP (National Tuberculosis Control Program (NTP), 2014). Capital costs were annuitized to get the annual costs. Facility quarterly reports from the year 2015 was used for estimating the total number of patients treated in each facility. Finally, average per patient costs were calculated for each type of treatment model.

The DS-TB patients were interviewed using a pre-tested questionnaire adopted from the Stop-TB questionnaire on patient's cost (Stop TB Partnership DOTS Expansion Working Group (TB and Poverty subgroup), 2008). The variables of interest are the



duration of illness, time elapsed before diagnosis, present status of the illness, how many healthcare providers has been consulted, the direct costs incurred in each encounter, duration of DOTS treatment, transportation cost to DOTS center, any friend/relative accompanying with and the opportunity cost of their time, and the lost work days and income of the patients among others.

Target Population and Study Sample

This study covers pulmonary form of DS-TB patients aged 18 years or more from all administrative divisions of Bangladesh. Total 1,000 DS-TB patients' data was collected for patient level cost and outcomes (QALY) estimation under this study. Out of these 1,000 patients, 402 were under CHW treatment model and 598 were under CM treatment model. Study locations are shown in the Appendix Figure 1A.

Glick (H. A. Glick, 2011) proposed a sample size formula for cost-effectiveness evaluation of clinical trials. Although our study is not a typical clinical trial, given the nature of the intervention and the study design we can apply the formula for calculating the required sample size for our study. The formula calculates the sample size for each of the two groups with similar standard deviation of costs and effect and same sample size:

$$n = \frac{2(Z_{\alpha} + Z_{\beta})^{2}[sd_{c}^{2} + (W * sd_{q}^{2})^{2} - (2W\rho * sd_{c} * sd_{q})}{(WQ - C)^{2}}$$

Where:

 Z_{α} is the Z-statistic for the level of Type I error (set at 95%)

 Z_{β} is the Z-statistic for the level of Type II error (set at 80%)



 sd_q , sd_c are the std deviations for each group for treatment effect and cost respectively

W is the Maximum Willingness to Pay

Q is the expected mean difference in treatment effectiveness

C is the expected mean difference in treatment cost

 ρ is the expected correlation of the difference in cost (C) and effect (Q)

This is a measure of the covariance of changes in effectiveness and changes in cost. Negative covariance, where cost decreases with increasing effectiveness result in a larger sample size. Positive covariance where cost increases with increasing effectiveness result in smaller sample sizes.

With 95% confidence interval and 80% power of the test, we assumed that the standard deviation of costs (sd_c) is 400 USD, standard deviation of effect (sd_q) is 0.2 QALY, ρ , correlation of difference in cost (C) and effect (Q) is 0.4. The expected mean difference in treatment effectiveness (Q) is 0.4 QALY and expected mean difference in treatment cost (C) is 500 USD. We set the willingness-to-pay threshold (W) at the three times of GDP of Bangladesh which is 3942 USD (BDT 315,360) (Macroeconomics, 2001). We found the sample size for both treatment groups is 405 which is equal to our study sample size.

Ethical Consideration

The study has already got ethics approval from University of South Carolina in the USA where the PI is a PhD student. Institutional Board Review (IRB) approval was



also obtained from Jahangirnagar University in Bangladesh. A third and final approval was obtained from WHO Research Ethics Review Committee (WHO ERC).

Settings and Locations

DF implements their DS-TB program in 22 districts in the North-Western part of Bangladesh. The DS-TB patients detected in these areas are treated following same regimen but the delivery of the DOTS is done by the community members. Whereas, in rest of the 42 districts all over Bangladesh follows the treatment delivery method by community health workers. This study collected data from districts of all eight divisions of Bangladesh.

Study Perspective

The study will be conducted form the societal perspective, which will encompass all costs incurred by the health care providers, patients, and community.

Comparators

For the DS-TB patients standard 6 months regimen is followed by all participating NGOs. However, the mode of delivery is different for different NGOs. BRAC has employed Community Health Workers (CHWs) besides the DOTS centers to ensure patient compliance, while Damien Foundation (DF) trained and employed influential community members to help the patients to be adhered to the treatment protocol. Another NGO, Salvation Army Bangladesh, is using drug sellers at the pharmacies as the counselor and drug distributors for the TB patients. We treated drug sellers under the community model. This study will conduct economic evaluation between this two DS-TB treatment delivery models.



Time Horizon

The study will take a life-time horizon to capture all costs and outcomes according to WHO guideline (Edejer, 2003). Since this study includes patients 18 years and older, we assume they will live through to their life expectancy (72 years in 2015). Mean age of the patient interviewed was 45 years. Therefore, we repeated the cycle for 27 times to include their whole lifespan.

Discount Rate

The study will follow the WHO discount rate of 3% for both costs and outcomes (Edejer, 2003).

Choice of Outcome

Quality Adjusted Life Year (QALY) is the chosen outcome for this study.

Measurement of Effectiveness

QALY has been modeled using patient-level EuroQol-5 Dimensions-5 Levels (EQ-5D-5L) measure of health related quality of life (EuroQol Research Foundation, 2017). We used Zimbabwe score sets for EQ-5D, since Zimbabwe is another developing country like Bangladesh and we assume the EQ-5D score sets would have been similar between populations of these two countries. Finally, the QALY measure was estimated after controlling for patient's socio-economic factors. Lognormal (log value of QALY as dependent variable) has been found more suitable model. Recycled prediction has been used to find out the estimated QALY for each type of patient outcome (e.g., cure, failure, and default) for both regimens (Glick Book). For death state utility has been assumed zero.



Measurement of Costs

Health systems costs were collected from different sources. Prevention and promotion costs, training costs, meeting costs, human resources costs, capital costs, and other costs were collected from DS-TB treatment facilities under the purview of the study using a pre-set provider questionnaire. Heath facility managers, personnel responsible for financial transactions were interviewed using the questionnaire and relevant costs were collected. Capital costs were annuitized using 5% interest rate and 10-year lifetime for Microscopes and Gene Xpert machines, and vehicles. Costs for drugs and diagnostic tests were assumed to be equal across two treatment modalities since both follows the same drug regimen and treatment protocol. Per patient cost of delivering DS-TB treatment for both types of regimen have been estimated dividing costs among the number of treated patients in one year in those facilities using total number of patients got treatment in 2015. Patient costs were collected through interviews of DS-TB patients using a pre-set and pre-tested questionnaire prepared on the basis of Stop-TB questionnaire on patient's cost (Stop TB Partnership DOTS Expansion Working Group (TB and Poverty subgroup), 2008). The questionnaire was translated in Bengali and the retranslated in English to validate the translation. Both patient and provider questionnaires were pre-tested and changes were incorporated before using those in the survey. Total costs of treating a patient under each regimen was calculated by summing up the patient and provider level costs. Then the cost was estimated using a Generalized Linear Model with Gamma variance and log link using data from 1,000 interviewed DS-TB patients who had complete information on disease outcomes. GLM diagnostics were used to validate the GLM variance and link. Recycled prediction was used to estimate the costs for each type



of patient outcome (e.g., cure, failure, default) for both regimens (H. Glick, Doshi, Sonnad, & Polsky, 2007). For death state costs were assumed to be zero.

Models and Analysis

A dynamic, stochastic, Markov simulation model was used to model the costeffectiveness of the two comparison regimens. Markov models have unique
characteristics which fit the progression of TB well. For example, in a Markov model
states are mutually exclusive, states are complete (i.e. no people are lost) and people
remain in each state for a fixed period of time. Also, a Markov model is preferred over
the decision trees when health event repeats over time, or have longer term health effects,
effect of treatment either stops quickly after initial treatment or continue at an earlier
level, and the risk of different health events does not depend on patient's prior history.

We can represent the Markov model for DS-TB as Figure 2.4. A more simplistic decision tree model can be shown as in Figure 5.1:

From the Markov model and decision tree we can find that after starting of the treatment the DS-TB patient can move to either of the four states, e.g., cure/treatment complete, failure/relapse, default, and death. Cure state has been defined as those who had completed treatment protocol without any evidence of failure and had at least three consecutive negative cultures from samples collected at least 30 days apart in the final 12 month of treatment, or "Treatment Completed", those who completed treatment according to treatment protocol but did not meet the definition for cure because of lack of bacteriological results.



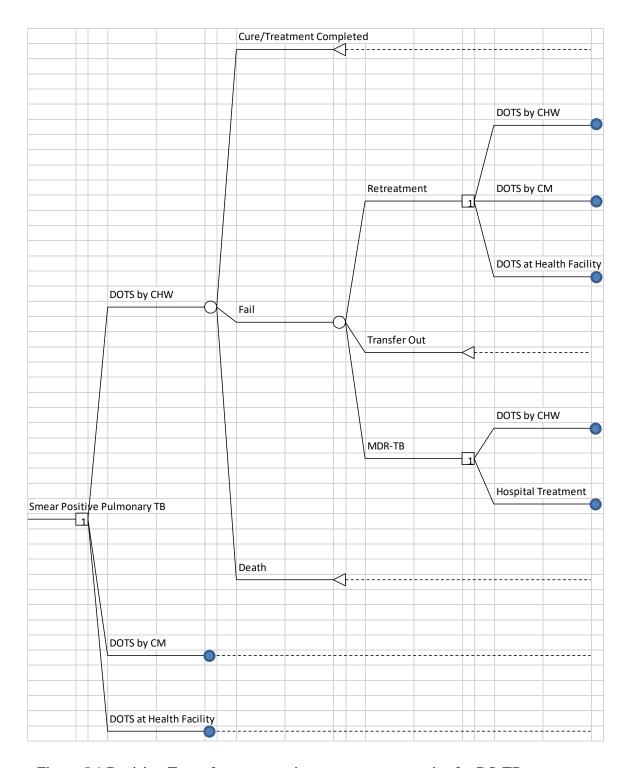


Figure 5.1 Decision Tree of two comparison treatment strategies for DS-TB



Failure/relapse cases were defined as those who had been treated for DS-TB, were declared cure or treatment completed at the end of their most recent course of treatment, and later diagnosed with a recurrent episode of DS-TB. Lost to follow up/ default cases were defined as those DS-TB patients whose treatment was interrupted for two or more consecutive months for any reason (National Tuberculosis Control Program (NTP), 2013, 2014).

Here death is the absorbing state, i.e., if a patient is dead he/she can move from that state to another. If any patient is cured he/she can remain cured, relapse/reinfection may occur, lost to follow-up (default) or can be dead. On the other hand, the failed/relapsed patients undergo another cycle of treatment and can culminate into cure, remain failed, can default, or can be dead as well. Similarly, from default state one can move over to other three states.

Cost-effectiveness evaluation was performed using *heemod* package in R (Filipović-Pierucci, Zarca, & Durand-Zaleski, 2017). The BCEA package was also used to validate the results form analysis with *heemod* package (Filipović-Pierucci et al., 2017). Cost and utility data was modeled using STATA 14.2 (StataCorp, 2015).

Measurement of Transition Probabilities

As mentioned earlier, cost-effectiveness analysis of these two regimens has not been undertaken yet. However, the programmatic outcome for the two treatment strategies has been derived from the quarterly reports on the health outcomes after 12-15 months of completion of treatment, which were collected from the health facilities. Using the reports from 2015, total number of patients and their transition between different



states have been calculated. Transitional probabilities were estimated from these numbers of DS-TB patients in different states using Markov simulation as an evidence synthesis technique (Sutton, Welton, & Cooper, 2012). R has been used for the analysis along with *r2jags* package for estimating the transitional probabilities for two treatment regimens (Su & Yajima, 2012).

From these two studies the transitional probabilities for first two cycles were estimated and furnished in Table 5.1.

Table 5.1 Transitional Probabilities of DS--TB Treatment Regimens

Input variable	CM	CHW
Cure to Cure (tpC2C)	0.609	0.529
Cure to Failure (tpC2F)	0.126	0.133
Cure to Default (tpC2Def)	0.127	0.170
Cure to Death (tpC2Death)	0.138	0.168
Failure to Cure (tpF2C)	0.354	0.268
Failure to Failure (tpF2F)	0.209	0.244
Failure to Default (tpF2Def)	0.209	0.240
Failure to Death (tpF2Death)	0.228	0.248
Default to Cure (tpDef2C)	0.250	0.245
Default to Failure (tpDef2F)	0.252	0.255
Default to Default (tpDef2Def)	0.250	0.250
Default to Death (tpDef2Death)	0.248	0.250

Parameters

Patient level cost for each regimen as well as the per patient provider costs are shown in Table 5.2.

Table 5.3 shows parameters for the cost-effectiveness analysis along with their distributions. Parameters mainly consist of transitional probabilities for transition between different states, costs for treating each type of states, and the utility of each states. Apart from this initial age was determined as the mean age of the interviewed



population (45 years). Time horizon was therefore fixed at 27 years, accounting for the rest of the general life expectancy of Bangladeshi people (life expectancy at birth in Bangladesh is 72 years (The World Bank, 2017b)). Both age and cycle parameters were kept fixed for the model.

Costs parameters follow gamma distribution as mentioned earlier and measured in 2015 Bangladeshi Taka (BDT). Utility values are measured in QALY and their distributions were lognormal.

Table 5.2 Patient, Provider and Total Costs for two regimens of DS-TB treatment in Bangladesh

Costs	CHW	CM
A. Patient Level Costs	Mean (BDT)	Mean (BDT)
Direct Costs		
Before Diagnosis Costs	10,894.24	14,895.78
TB Diagnosis Costs	777.37	1,328.08
Follow-up Costs	213.55	144.33
Drug Collection Costs	2.81	22.49
Hospital Costs	1,421.63	3,250.39
Additional Food Costs	1,926.06	2,457.74
Accompanying Person Costs	365.7	389.66
MDR-TB Relocation Costs		
Drug Side-effects Costs	438.33	432.11
Total Direct Costs	16,039.69	22,920.58
Indirect Costs		
Patient Opportunity Costs (Income Loss)	432.8	396.63
Accompanying Person Opportunity Costs	508.28	784.68
Total Indirect Costs	941.08	1,181.31
Total Patient Level Costs	16,980.77	24,101.89
B. Provider Level Costs		
Prevention and Promotion Costs	Not reported	Not reported
Diagnostic Costs	Same	Same
Drug Costs	Same	Same
Training Costs	6.23	44.02



Meeting Costs	0.70	5.67
Incentive Payment	100.99	13.21
Human Resources Costs	1,827.20	1,592.01
Capital Costs	36.56	94.41
Other Costs	2.29	11.26
Total Provider Level Costs	19,73.97	1,760.56
Total Costs	18,954.74	25,862.45

Table 5.3 Input Parameters for Cost-Effectiveness Analysis of DS-TB Treatments

Variable	Distribution ^a	Value	Low	High	Reference
Category					
Starting age of	Fixed	45 Years	-	-	Study Data
cohort (mean)					
Time horizon	Fixed	27 Years	-	-	World
					Bank
Cost of Cure	Gamma	BDT	BDT	BDT	Study Data
(CM)		25,095	20,076	30,114	
Cost of Failure	Gamma	BDT	BDT	BDT	Study Data
(CM)		31,849	25,479	38,219	
Cost of Default	Gamma	BDT	BDT	BDT	Study Data
(CM)		34,132	27,306	40,958	
Cost of Cure	Gamma	BDT	BDT	BDT	Study Data
(CHW)		17,719	14,175	21,263	
Cost of Failure	Gamma	BDT	BDT	BDT	Study Data
(CHW)		22,488	17,990	26,985	
Cost of Default	Gamma	BDT	BDT	BDT	Study Data
(CHW)		24,100	19,280	28,920	
Utility of Cure	Lognormal	0.783	0.626	0.940	Study Data
(CM)					
Utility of	Lognormal	0.738	0.590	0.886	Study Data
Failure (CM)					
Utility of	Lognormal	0.742	0.594	0.890	Study Data
Default (CM)					
Utility of Cure	Lognormal	0.776	0.621	0.931	Study Data
(CHW)					
Utility of	Lognormal	0.732	0.586	0.878	Study Data
Failure (CHW)					
Utility of	Lognormal	0.736	0.589	0.883	Study Data
Default (CHW)					
Discount Rate	Fixed	3%	0%	6%	Edejer,
					2003

^a In Probabilistic Sensitivity Analysis

Results



Base Case Results

Base case results show that over the 27 cycles each DS-TB patient under CM treatment model gains 3.61 QALYs with a cost of BDT 131,555. For the DS-TB patients under the CHW model the cost is 81,650 and the QALY gain is 3.12. The Incremental Cost-Effectiveness Ratio (ICER) is 103,454, i.e., the CM model is cost-effective if per QALY gain one can afford more than BDT 131,454 (Table 5.4).

Table 5.4 Base Case Results

Treatment Model	Cost Per Patient (BDT)	QALY Gained	ICER
CM	131,555	3.61	103,454
CHW	81,650	3.12	

Deterministic Sensitivity Analysis

Parameters were varied on different scales for high and low values and univariate sensitivity analysis was performed to assess the robustness of the findings. A tornado plot has been prepared to illustrate the effect of change in each variable. Based on the findings in one-way sensitivity analysis, input parameters were varied accordingly to see their impact in probabilistic sensitivity analysis.

In the tornado plot, shown in the Figure 5.2, we find that the most influential parameters were utility of cure state for CM model, followed by utility of cure state of CHW model, costs of cure state of CM model, utility of loss to follow-up state of CHW model, and utility of loss to follow-up state of CM model. according to the more effects on the results.



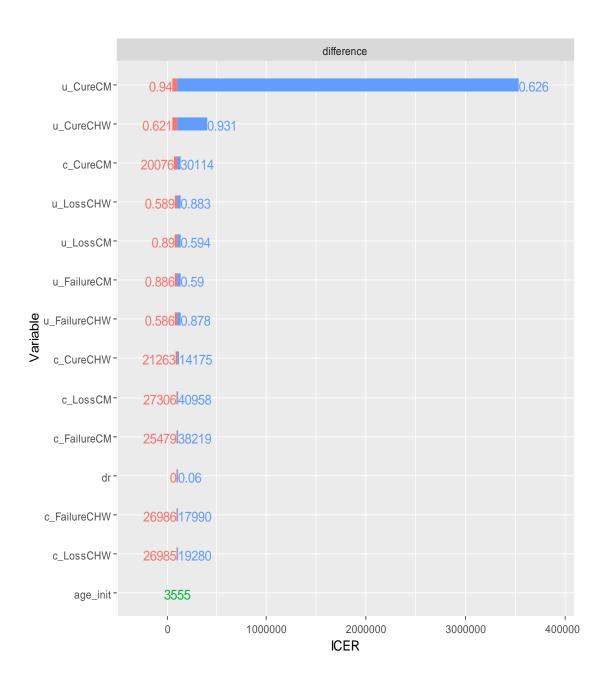


Figure 5.2 Tornado Plot of Deterministic Sensitivity Analysis



Probabilistic Sensitivity Analysis

In probabilistic sensitivity analysis, the costs values and utility values for each state were varied by 20%. Initially, the base case Markov model with all parameters with their values and distribution was run for 1000 iterations and the base case results were assessed using incremental cost effectiveness ratio (ICER), Cost-effectiveness Acceptability Curve (CEAC), and Expected Value of Perfect Information (EVPI).

Incremental Cost-effectiveness Ratio (ICER)

From the results reported in the Table 6 shows that over life-time the CM method costs BDT 197,680; while CHW model costs BDT 76,836. At the same time QALY gained by CM model is 6.13 whereas in CHW model QALY gain is 2.91. Therefore, the resultant ICER is BDT 37,487 per QALY gained. It shows that CM method is cost-effective if willingness to pay is more than BDT 37,487 (Table 5.5). Results shown in the cost effectiveness plane also shows that the ICER is in the North-East Quadrant (Figure 5.4).

Table 5.5 ICER from Probabilistic Model

Treatment Model	Cost Per Patient (BDT)	QALY Gained	ICER	EVPI
CM	197,680	6.13	37,487	18,388
CHW	76,836	2.91		

Cost-Effectiveness Acceptability Curve (CEAC)

Since the ICER for each QALY gain is BDT 37,480, cost-effectiveness acceptability curve for CM and CHW crosses over at that point. After the value CM becomes more cost-effective and thereby acceptable (Figure 5.5).



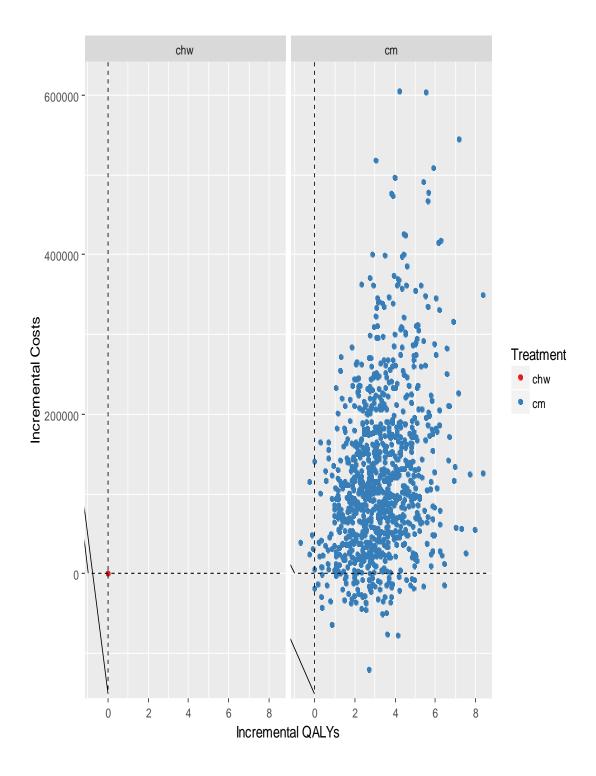


Figure 5.4 Cost-Effectiveness Plane



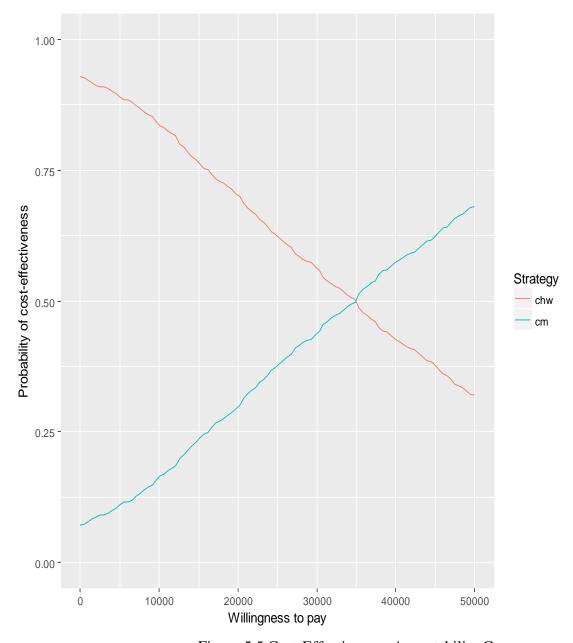


Figure 5.5 Cost-Effectiveness Acceptability Curve

Expected Value of Perfect Information (EVPI)

EVPI is the absolute limit of the value of further research that would completely eliminate the uncertainty around the parameters in the model. EVPI value of BDT 18,388 shows that with reducing uncertainty around the parameters would require only BDT 18,388. It is also evident from the figure 6 that with EVPI peaks between BDT 300,00



and BDT 400,000. It signifies that with increasing willingness-to-pay EVPI also increases till certain value, then it declines.

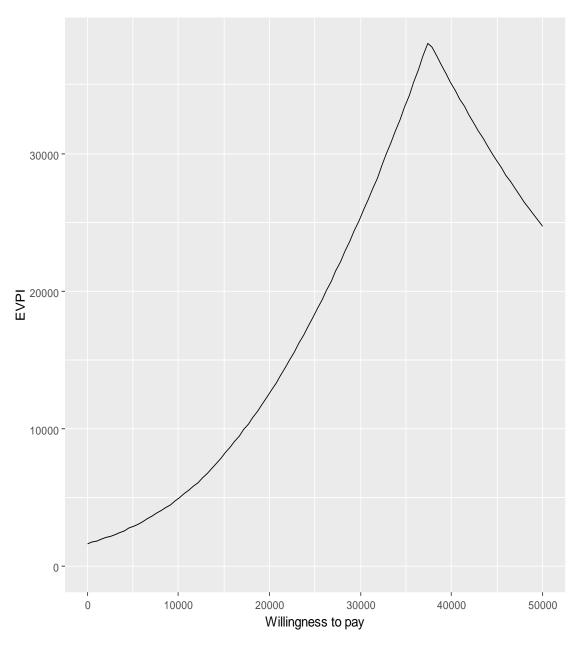


Figure 5.6 Expected Value of Perfect Information (EVPI) Curve



Discussions

The study results show that the CM method of DS-TB treatment is cost-effective in Bangladesh from a societal perspective. ICER form base case analysis is BDT 103,454 (USD 1293) which is even lower than per capita GDP of Bangladesh in 2015, which is USD 1342. The deterministic sensitivity analysis shows that the utility garnered from the cure status of CM method is the principal factor behind the result. Even when we changed the utility value by 20% (within a range of 0 and 1), we found that the CM method remains cost-effective.

Several studies have also been conducted to assess the cost-effectiveness of DOTS program itself. DOTS was found cost-effective in developing country settings, Thailand (Hunchangsith et al., 2012), Egypt and Syria (Vassall et al., 2002), Botswana (Moalosi et al., 2003), Haiti (Jacquet et al., 2006), Uganda (Okello et al., 2003), Brazil (Mohan et al., 2007). These studies invariably documented that the DOTS strategy or involving the communities in the care process is cost-effective over SAT. In Malawi it was shown that community based DOTS was cost-effective than the usual hospital-based model (Floyd, Skeva, Nyirenda, Gausi, & Salaniponi, 2003).

Very few studies compared between community member DOTS model and community health worker DOTS model. In a study conducted in Thailand, the results show that community member model is dominant, while ICER was USD 1,100 for each DALY gained in health worker model (Hunchangsith et al., 2012). In some countries, randomized controlled trials (RCT) were conducted to find the efficacy of DOTS model. In Nepal, both family based DOTS and Community based DOTS were found to be capable of attaining international targets for treatment success (Newell, Baral, Pande,



Bam, & Malla, 2006). In Senegal, it was found that the package based on improved patient counseling and communication, decentralization of treatment, patient choice of DOT supporter, and reinforcement of supervision led to improved patient outcomes (Thiam et al., 2007).

Community members can be anyone in the community. Neighbors are mostly selected by DF to cater drugs daily to the DS-TB patients. Salvation Army in Dhaka city deployed drug sellers in the vicinity to provide anti-TB drugs to the DS-TB patients on regular basis. This idea of involving community is not novel; The Union prefers the integration of community members in the treatment procedure (Aït-Khaled et al., 2010). Most important objective of the DOTS model was to enhance the adherence to the TB drug regimen (World Health Organization & Stop TB Initiative, 2010). In Bangladesh, it has been found that community based models works better than the hospital-based model (Islam et al., 2002). But comparison between community member model and community health workers have not been done to move forward with the best and cost-effective model.

From costs figures for the two methods, we see that patients incur more costs for DS-TB treatment in case of CM model. Patients' costs were higher for before diagnosis costs, diagnosis costs, hospital costs, and additional food costs. On the other hand, providers with CM model spend more on training, while CHW model spend more on incentives to the CHWS. Each CHW gets a remuneration of BDT 500 for successful completion of the DS-TB regimen. Interestingly, no facility reported any costs for prevention activities. Also note that, present treatment guidelines for DS-TB does not



require any hospitalization. These show that CM model lacks in early diagnosis of the patients which leads to undue hospitalization and more costs.

However, CM model is successful in reducing stigma related to TB disease. It has been documented in several studies that effective stigma reducing strategies are focused on individual and community levels (Heijnders & Van Der Meij, 2006). Stigma related to TB disease and its care is widespread in Bangladesh and women are the worst sufferer (Somma et al., 2008). Community member's involvement reduce the chance of spread of the news of TB infection and thereof avoid gossiping, undue fear of transmitting the disease and potential discrimination. If community health workers do not divulge the news of TB disease, her daily presence in a particular house is a telltale sign for neighbors which leads to discrimination for the TB patient. Therefore, CHW models are not adequate to ensure the adherence and proper treatment which is evident in the utility value of CM method.

This study has several limitations. Although for transitional probabilities we used quarterly reports from 18 facilities under the survey, the reported outcomes were not verified like controlled trial settings. This can make the results biased. Cost and effectiveness data from the patients may suffer from the recall bias, although we included only those patients who completed their treatment not more than six months.

Despite these limitations, this study tried to collect the patient and provider level data comprehensively using pre-set questionnaires. To our best knowledge, it is also the first study to conduct economic evaluation between CHW and CM model of DS-TB treatment in developing country setting.



Community based DOTS is getting acceptance worldwide due to its proven efficacy. However, early diagnosis and treatment of TB cases is still a far cry due to lack of knowledge about TB in general population and absence of any proper prevention programs. This indicates the importance of deploying community members in disseminating the knowledge about TB signs and symptoms which may facilitate early diagnosis of the disease as well as reduce the stigma associated with TB disease.

In conclusion, the evidence of cost-effectiveness of CM method encourages us to adopt this model all over Bangladesh. More involvement of the community members of all sort will help prevent the transmission of the disease, early diagnosis will lead to early cure at low costs, and treatment adherence will help us to attain the End TB strategy by 2035.

Acknowledgements

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CHAPTER 6

STUDY III

 $6.1\ Economic\ Evaluation\ of\ Multi\ Drug-Resistant\ Tuberculosis\\ (MDR-TB)\ Treatment\ Approaches\ in\ Bangladesh^3$

³ Haider, M. R. To be submitted.



Introduction: Worldwide Tuberculosis (TB) control has been halted by the emergence of multi-drug resistant TB (MDR-TB). Bangladesh has also experienced surge in the number of MDR-TB cases with a 29% of MDR-TB cases were found among the retreatment of pulmonary TB cases in 2015. In Bangladesh, two MDR-TB treatment regimens (9-month and 20-24 month) are practiced and this study intends to conduct economic evaluation between those two.

Methods: The incremental cost-effectiveness ratio (ICER) of treating MDR-TB patients, 35 years old on average, by the 9-month regimen versus the 20-24-month regimen was compared using a Markov model with life-time horizon (37 years). The measure of effectiveness, Quality adjusted life year (QALY) and cost of treatment was collected from 145 MDR-TB patients (58 undergone the 9-month treatment and 87 from the 20-24-month regimen) in Bangladesh. Transition probabilities between Markov states were estimated from two published studies and cost and QALY both were discounted at a rate of 3%. Deterministic and probabilistic sensitivity analyses were conducted in a Monte Carlo Simulation using R.

Results: Based on the study data, each patient under the 9-month regimen gained 6.21 QALY with a total cost of BDT 987,418. Whereas, each patient under CHW model gained 5.74 QALY by incurring costs of BDT 1,501,221. Therefore, the 9-month regimen is clearly dominating over the 20-24-month regimen because it costs less while it gains more QALY.

Conclusions: Our study results suggest that the shorter 9-month regimen remains costeffective in Bangladesh setting with changing costs and utility parameters varied in the



probabilistic sensitivity analysis. MDR-TB treatment is itself cost-effective in developed countries and with cost-effective shorter regimen both treatment adherence and efficacy of the treatment will be improved.

Keywords: Economic Evaluation, Multi Drug Resistant Tuberculosis, Costeffectiveness, Cost, QALY, Bangladesh Regimen, Bangladesh



Introduction

Tuberculosis (TB) is a deadly tropical disease caused by *Mycobacterium tuberculosis*, a bacillus which typically affects lungs (pulmonary tuberculosis) in addition to other parts of the human body. Tuberculosis is prevalent in the temperate region of the world and this tropical disease is endemic in South-East Asian and African countries. India and China, the two largest countries in terms of population, had the highest number of cases (26% and 12% respectively) in 2012. Bangladesh, a South Asian country with hot tropical weather, also harbors the disease in huge numbers of afflicted people. Bangladesh is a high burden TB country and its number of Multi-Drug Resistant TB (MDR-TB) patients is on the rise (World Health Organization (WHO), 2016a).

The world has experienced a slow gain in TB control in recent years and that progress has been halted by the emergence of MDR-TB and Extremely Drug Resistant TB (XDR-TB) strains. Bangladesh has also experienced a surge in the number of MDR-TB cases with a 29% of MDR-TB cases found among the re-treatment of pulmonary TB cases in 2015 (World Health Organization (WHO), 2016a).

Bangladesh follows a model involving community health workers or community members to provide drugs regularly to the patients undergoing drug sensitive TB (DSTB). Under the public private partnership models different Non-governmental organizations (NGOs) are responsible for directly observed treatment short-course (DOTs). Any lack of adherence may result in menacing drug resistant strains, MDR or XDR TB; treatment for both the conditions are costly and time consuming. The current WHO guidelines prescribed treatment regimen is of 20-24 months, while the shortest effective MDR-TB treatment regimen spans over nine months (Deun et al., 2010). The



prolonged treatment schedule may result in more incidence of treatment discontinuation. WHO in cooperation with STOP TB Partnership came up with a response plan in 2007-2008 and Bangladesh is one of the seven countries using shorter treatment regimens for MDR-TB in June 2013 (World Health Organization (WHO), 2013c).

National Tuberculosis Control Program (NTP) in Bangladesh follows the 20-24-month treatment regimen for MDR-TB patients. The NTP follows the Programmatic Management of Drug-resistant TB (PMDT) guideline (Falzon et al., 2011) and the treatment is supervised and administered by DOTS providers (National Tuberculosis Control Program (NTP), 2013)

Damien Foundation (DF) runs their own protocol of treatment for MDR-TB patients, which span over 9 months and differs in drug composition as well. DF generally admits the MDR-TB patients in one of their three hospitals located in Jalchatra of Madhupur in Tangail district, Shomvuganj in Mymensingh district and at Anantapur in Netrakona district for four months of intensive phase; which is followed by the continuation phase of five months for which drugs are administered at patients' home by DOTs providers (Damien Foundation Bangladesh, 2015). DF has initiated a shorter regimen treatment of 9 months in 1997 (Van Deun et al., 2010), which eventually came to known as "Bangladesh" regimen (Aung et al., 2014). In a recent publication in 2014, DF researchers presented their findings from their observation study that 84.4% of the patients undergone the shorter regimen had bacteriologically favorable outcomes up to two years after treatment completion (Aung et al., 2014). This success of Bangladesh regimen inspired United Sates Agency for International Development (USAID), the International Union Against Tuberculosis and Lung Disease (the Union), and Janssen



Research & Development, LLC to commission a clinical trial to find out the effectiveness of the regimen in other countries like Ethiopia, Mongolia, South Africa, and Vietnam (International Union Against Tuberculosis and Lung Disease (The Union), 2017).

Although several studies showed the cost-effectiveness of MDR-TB treatment as a whole in developed and developing country settings (Diel, Nienhaus, Lampenius, Rüsch-Gerdes, & Richter, 2014; Diel, Vandeputte, et al., 2014; Fitzpatrick & Floyd, 2012), no study was conducted to perform economic evaluation of two regimens, the 9-month regimen (shorter) and current WHO recommended the 20-24-month regimen (current). This study aims to fill the void in conducting the cost-effectiveness evaluation of these two treatment regimens from a societal perspective (including both patients and payer perspectives) using Quality Adjusted Life Years (QALYs) as the outcome for effectiveness, which qualifies this as a cost-utility study in health economics parlance (Drummond, Sculpher, Claxton, Stoddart, & Torrance, 2015).

Methods

Study Design and Data Sources

The study follows a purposive sampling method for interviewing MDR-TB patients. According to the recent estimates in 2014 number of laboratory-confirmed MDR-TB patients was 954 in Bangladesh and the prevalence of MDR-TB is 5,100 in 2015 (World Health Organization (WHO), 2016a). In our study area, it is understandable that the number will be significantly lower. Therefore, we collected the information of the MDR-TB patients from the TB control programs and reach those who were accessible (Brazier et al., 2002).



Institution-level data has been collected for assessing the provider level costs associated with TB treatment. Number of patients diagnosed and treated in 2015 has been collected from the secondary sources such as the World TB report published by WHO and the Annual TB report of NTP, Bangladesh. Drug costs have been collected from the STOP TB Global Drug Facility website. Costs for diagnostic tests have been estimated after consulting with local experts who are knowledgeable of the MDR-TB programs in Bangladesh and also know the market price of different tests.

Cost Of Illness (COI) includes direct, indirect and tangibles costs incurred by the patients (Centers for Disease Control and Prevention (CDC), 2013) and in this study the TB patients will be traced and interviewed for the detail cost descriptions. The variables of interest are the duration of illness, time elapsed before diagnosis, present status of the illness, how many healthcare providers has been consulted, the direct costs incurred in each encounter, duration of DOTS treatment, transportation cost to DOTS center, any friend/relative accompanying with and the opportunity cost of their time, and the lost work days and income of the patients among others. The programmatic cost has been collected using a pre-set questionnaire and using annuitization for capital costs per patient costs were calculated for each regimen.

The study will seek data on tuberculosis burden on the population of Bangladesh from different sources, e.g., published reports, program documentations and various surveys and the incidence of TB will be extracted from those sources. To find the QALYs gained through the program we interviewed patients using EuroQoL 5D-5L questionnaire and used the tariff provided by EuroQol.



Target Population and Study Sample

This study covers the adult (more than 18 years old) MDR-TB patients all over Bangladesh. 145 MDR-TB patients' data was used for patient level cost and outcomes (QALY) estimation for this study. Out of these 145 patients, 58 were under the 9-month regimen and 87 were under the 20-24-month regimen, who were under treatment in four MDR-TB treatment facilities (two for each regimens). Table 1 shows the number of patients interviewed under each regimen and the districts to which they belonged. In the Appendix Figure 1 the geographic location of the study districts is shown.

Glick (H. A. Glick, 2011) proposed a sample size formula for cost-effectiveness evaluation of clinical trials. Although our study is not a typical clinical trial, given the nature of the intervention and the study design we can apply the formula for calculating the required sample size for our study. The formula calculates the sample size for each of the two groups with similar standard deviation of costs and effect and same sample size:

$$n = \frac{2(Z_{\alpha} + Z_{\beta})^{2}[sd_{c}^{2} + (W * sd_{q}^{2})^{2} - (2W\rho * sd_{c} * sd_{q})}{(WQ - C)^{2}}$$

Where:

 Z_{α} is the standard normal quantile for the level of Type I error (set at 95%)

 Z_{β} is the standard normal quantile for the level of Type II error (set at 80%)

 sd_q , sd_c are the std deviations for each group for treatment effect and cost respectively

W is the Maximum Willingness to Pay



Q is the expected mean difference in treatment effectiveness

C is the expected mean difference in treatment cost

 ρ is the expected correlation of the difference in cost (C) and effect (Q)

This is a measure of the covariance of changes in effectiveness and changes in cost. Negative covariance, where cost decreases with increasing effectiveness result in a larger sample size. Positive covariance where cost increases with increasing effectiveness result in smaller sample sizes.

With 95% confidence interval and 80% power of the test, we assumed that the standard deviation of costs (sd_c) is 100 USD, standard deviation of effect (sd_q) is 0.25 QALY, ρ , correlation of difference in cost (C) and effect (Q) is 0.5. The expected mean difference in treatment effectiveness (Q) is 0.15 QALY and expected mean difference in treatment cost (C) is 1000 USD. We set the willingness-to-pay threshold (W) at the three times of GDP of Bangladesh which is 3942 USD (BDT 315,360) (Macroeconomics, 2001). We found the sample size for one group is 70 and another is 104 with a 2:1 sample size ratio.

Settings and Locations

DF implements their MDR program in 22 districts in the North-west part of Bangladesh. The MDR-TB patients detected in these areas are treated following the 9-month regimen. Whereas, in rest of the 42 districts all over Bangladesh follows the 20-24-month regimen. Thus this study covers the whole country.



Ethical consideration

The study has already got ethics approval from University of South Carolina in the USA where the PI is a PhD student. Institutional Board Review (IRB) approval will also be taken from Jahangirnagar University in Bangladesh. A third and final approval was obtained from WHO Research Ethics Review Committee (WHO ERC).

Study Perspective

The study will be conducted form the societal perspective, which will encompass all costs incurred by the health care providers, patients, and community.

Comparators

The two distinct programs carried out by NTP and DF will be the comparators in this study. DF runs a 9-month regimen which administers high-dose Gatifloxacin (GFX), Ethambutol (EMB), Pyrazinamide (PZA), and Clofazimine (CFZ) throughout, supplemented during the minimum 4-month intensive phase by Kanamycin (KM), Prothionamide (PTH), and Isoniazide (INH) (Aung et al., 2014).

NTP follows the 20-24-month regimen which includes Kanamycin (KM),
Ofloxacin (OFX), Pyrazinamide (PZA), Ethonamide (ETO), and Cycloserine (CS) in 610 months (on average 8 month) long intensive phase and Ofloxacin (OFX),
Pyrazinamide (PZA), Ethonamide (ETO), and Cycloserine (CS) in 13-18 months (on
average 12 months) of continuation phase of treatment (National Tuberculosis Control
Program (NTP), 2013).



Time Horizon

The study will take a life-time horizon to capture all costs and outcomes according to WHO guideline (Edejer, 2003). Since this study includes patients 18 years and older, we assume they will live through to their life expectancy (72 years in 2015). Mean age of the patient interviewed was 35 years. Therefore, we repeated the cycle for 37 times to include their whole lifespan.

Discount Rate

The study will follow the WHO discount rate of 3% for both costs and outcomes (Edejer, 2003).

Choice of Outcome

Quality Adjusted Life Year (QALY) is the chosen outcome for this study.

Measurement of Effectiveness

QALY has been modeled using patient-level EuroQol-5 Dimensions-5 Levels (EQ-5D-5L) measure of health related quality of life (EuroQol Research Foundation, 2017). We used Zimbabwe score sets for EQ-5D, since Zimbabwe is another developing country like Bangladesh and we assume the EQ-5D score sets would have been similar between populations of these two countries. Finally, the QALY measure was estimated after controlling for patient's socio-economic factors. Lognormal (log value of QALY as dependent variable) has been found more suitable model. Recycled prediction has been used to find out the estimated QALY for each type of patient outcome (e.g., cure, failure, and default) for both regimens (Glick Book). For death state utility has been assumed zero.



Measurement of Costs

Health systems costs were collected from different sources. Prevention and promotion costs, training costs, meeting costs, human resources costs, capital costs, and other costs were collected from MDR-TB treatment facilities under the purview of the study using a pre-set provider questionnaire. Heath facility managers, personnel responsible for financial transactions were interviewed using the questionnaire and relevant costs were collected. Capital costs were annuitized using 5% interest rate and 10year lifetime for Microscopes and Gene Xpert machines, and vehicles. Costs for drugs were derived from the Stop TB Global Drug Repository (Stop TB Partnership, 2017) and per patient drug costs were calculated for each regimen (Appendix Table 1 and Table 2). Diagnostic tests costs were derived from expert knowledge from the officials working in MDR-TB program in Bangladesh (Appendix Table 3 and Table 4). Per patient cost of delivering MDR-TB treatment for both types of regimen have been estimated dividing costs among the number of treated patients in one year in those facilities using Annual Tuberculosis Report by NTP, Bangladesh (National Tuberculosis Control Program (NTP), 2015). Patient costs were collected through interviews of MDR-TB patients using a pre-set questionnaire prepared on the basis of Stop-TB questionnaire on patient's cost (Stop TB Partnership DOTS Expansion Working Group (TB and Poverty subgroup), 2008). The questionnaire was translated English to Bengali and then retranslated back into English to validate the translation. Both patient and provider questionnaires were pre-tested and changes were incorporated before using those in the survey. Total costs of treating a patient under each regimen was calculated by summing up the patient and provider level costs. Then the cost was modeled using a Generalized Linear Model with



Gamma family and log link using data from 145 interviewed MDR-TB patients who had complete information on disease outcomes. GLM diagnostics were used to validate the GLM family and link. Recycled prediction was used to estimate the costs for each type of patient outcome (e.g., cure, failure, default) for both regimens (H. Glick et al., 2007). For death state costs were calculated as zero.

Models and Analysis

A dynamic, stochastic, Markov simulation model was used to estimate the costeffectiveness of the two comparison regimens. Markov model has some unique
characteristics which fit the progression of TB well. For example, in Markov model states
are mutually exclusive, states are complete (i.e. no people are lost) and people remain in
that state for a fixed period of time. Also, Markov model is preferred over the decision
trees when health event repeats over time, or have longer term health effects, effect of
treatment either stops quickly after initial treatment or continue at an earlier level, and the
risk of different health events does not depend on patient's prior history.

A simplistic decision tree model can be shown in Figure 6.1:

From the Markov model and decision tree we find that after starting of the treatment the MDR-TB patient can move to either of the four states, e.g., cure/treatment complete, failure/relapse, default, and death. Cure state has been defined as those who had completed treatment protocol without any evidence of failure and had at least three consecutive negative cultures from samples collected at least 30 days apart in the final 12 month of treatment, or "Treatment Completed", those who completed treatment according to treatment protocol but did not meet the definition for cure because of lack of bacteriological results.



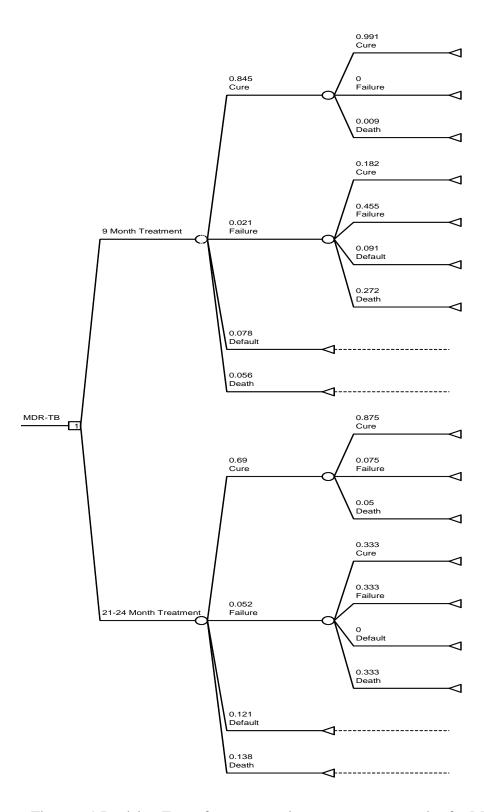


Figure 6.1 Decision Tree of two comparison treatment strategies for MDR-TB



Failure/relapse cases were defined as those whose treatment was needed to be terminated or at least change of two anti-TB drugs due to lack of conversion, bacteriological reversion in the continuation phase, evidence of additional acquired resistance, or adverse drug reaction. Lost to follow up/ default cases were defined as those MDR-TB patients whose treatment was interrupted for two or more consecutive months for any reason (National Tuberculosis Control Program (NTP), 2013).

Here death is the absorbing state, i.e., if a patient is dead he/she can move from that state to another. If any patient is cured he/she can remain cured, relapse/reinfection may occur, lost to follow-up (default) or can be dead. On the other hand, the failed/relapsed patients undergo another cycle of treatment and can culminate into cure, remain failed, can default, or can be dead as well. Similarly, from default state one can move over to other three states.

Cost-effectiveness evaluation was performed using *heemod* package in R (Filipović-Pierucci et al., 2017). BCEA package was also used to validate the results form analysis with *heemod* package (Filipović-Pierucci et al., 2017). Cost and utility data was modeled using STATA 14.2 (StataCorp, 2015).

Measurement of Transition Probabilities

As mentioned earlier, cost-effectiveness analysis of these two regimens has not been undertaken yet. However, the programmatic outcome for the two treatment strategies has been derived from two papers. One paper reported the results of the standardized treatment spanning 20-24 month (Van Deun et al., 2004). Another paper published recently reported the results of the alternative regimens of 9 month (Aung et



al., 2014). Both studies were conducted in Bangladesh. Transitional probabilities were estimated from these two studies using Markov simulation as an evidence synthesis technique (Sutton, Welton, & Cooper, 2012). R has been used for the analysis along with *r2jags* packages for estimating the transitional probabilities for two treatment regimens (Su & Yajima, 2012).

From these two studies the transitional probabilities for first two cycles were estimated and furnished in Table 6.1.

Table 6.1 Transition Probabilities of MDR-TB Treatment Regimens

Input variable	20-24-Month Regimen ^a	9-Month Regimen ^b
Cure to Cure (tpC2C)	0.813	0.846
Cure to Failure (tpC2F)	0.071	0.043
Cure to Default (tpC2Def)	0.024	0.080
Cure to Death (tpC2Death)	0.092	0.031
Failure to Cure (tpF2C)	0.143	0.119
Failure to Failure (tpF2F)	0.431	0.486
Failure to Default (tpF2Def)	0.139	0.040
Failure to Death (tpF2Death)	0.287	0.355
Default to Cure (tpDef2C)	0.088	0.023
Default to Failure (tpDef2F)	0.087	0.499
Default to Default (tpDef2Def)	0.637	0.365
Default to Death (tpDef2Death)	0.188	0.113

^a Source: (Aung et al., 2014)

Parameters

Patient level cost for each regimen as well as the per patient provider costs are shown in Table 6.2.

Table 6.3 shows parameters for the cost-effectiveness analysis along with their distributions. Parameters mainly consist of transitional probabilities for transition between different states, costs for treating each type of states, and the utility of each states. Apart from this initial age was determined as the mean age of the interviewed



^b Source: (Van Deun et al., 2004)

population (35 years). Time horizon was therefore fixed at 37 years, accounting for the rest of the general life expectancy of Bangladeshi people (life expectancy at birth in Bangladesh is 72 years (The World Bank, 2017b)). Both age and cycle parameters were kept fixed for the model. Transmission of secondary infection was assumed to be .003 annually based on the findings of a study that 0.03 new secondary cases may develop among the MDR-TB population (Sloot, Schim van der Loeff, Kouw, & Borgdorff, 2014).

Costs parameters follow a gamma distribution as mentioned earlier and are measured in 2015 Bangladeshi Taka (BDT). Utility values are measured in QALY and their distributions were lognormal.

Table 6.2 Patient, Provider and Total Costs for two regimens of MDR-TB treatment in Bangladesh

Costs	DS-TB Patients	MDR-TB Patients	
A. Patient Level Costs	Mean	Mean	
Direct Costs			
Before Diagnosis Costs	13287.16	14844.23	
TB Diagnosis Costs	1106.69	684.63	
Follow-up Costs	172.16	877.17	
Drug Collection Costs	14.6	1.17	
Hospital Costs	2515.23	7669.4	
Additional Food Costs	2244.01	2678.39	
Accompanying Person Costs	380.03	2114.7	
MDR-TB Relocation Costs	0	341.24	
Drug Side-effects Costs	434.58	1647.28	
Total Direct Costs	20154.46	30858.21	
Indirect Costs			
Patient Opportunity Costs (Income Loss)	407.07	1522.86	
Accompanying Person Opportunity Costs	673.57	2593.91	
Total Indirect Costs	1080.64	4116.77	
Total Patient Level Costs	21235.10	34974.98	
B. Provider Level Costs			
Prevention and Promotion Costs	652.17	202.02	



Diagnostic Costs	17825.00	38300.00
Drug Costs	42761.50	61833.27
Training Costs	2608.70	767.68
Meeting Costs	2608.70	808.08
Incentive Payment	3100.00	3100.00
Human Resources Costs	23728.70	28267.15
Capital Costs	22100.92	10101.01
Other Costs	345.73	101.01
Total Provider Level Costs	115731.41	143480.23
Total Costs	136966.51	178455.21

Table 6.3 Input Parameters for Cost-Effectiveness Analysis

Variable	Distribution ^a	Value	Low	High	Reference
Category					
Starting age of	Fixed	35 Years	-	-	Study Data
cohort (mean)					
Time horizon	Fixed	37 Years	-	-	(LE Data)
Cost of Cure (20-	Gamma	BDT	BDT	BDT	Study Data
24 Month)		194893	151740	238046	
Cost of Failure	Gamma	BDT	BDT	BDT	Study Data
(20-24 Month)		204155	149920	258390	
Cost of Default	Gamma	BDT	BDT	BDT	Study Data
(20-24 Month)		206983	162744	251222	
Cost of Cure (9	Gamma	BDT	BDT	BDT	Study Data
Month)		125977	98084	153870	
Cost of Failure (9	Gamma	BDT	BDT	BDT	Study Data
Month)		131964	96905	167023	
Cost of Default (9	Gamma	BDT	BDT	BDT	Study Data
Month)		133972	105376	162568	
Utility of Cure	Lognormal	0.777	0.524	1.00	Study Data
(20-24 Month)					
Utility of Failure	Lognormal	0.710	0.428	0.99	Study Data
(20-24 Month)					
Utility of Default	Lognormal	0.757	0.543	0.971	Study Data
(20-24 Month)					
Utility of Cure (9	Lognormal	0.813	0.560	1.00	Study Data
Month)					
Utility of Failure	Lognormal	0.746	0.464	1.00	Study Data
(9 Month)					
Utility of Default	Lognormal	0.793	0.579	1.00	Study Data
(9 Month)					



Discount Rate	Fixed	3%	0%	6%	Edejer,
					2003
Probability of	Binomial	0.03	0	0.06	Germany
Secondary					Paper
transmission (per					
year)					

^a In Probabilistic Sensitivity Analysis

Results

Base Case Results

The base case results show that after 37 cycles (years) each patient under the 9-month regimen gained 6.21 QALY with a total cost of BDT 987,418. Whereas, each patient under CHW model gained 5.74 QALY by incurring costs of BDT 1,501,221 (Table 5). Therefore, the 9-month regimen is clearly dominating the 20-24-month regimen because it costs less while it gains more QALY.

Table 6.5 Base Case Results

Regimen	Cost Per Patient (BDT)	QALY Gained	ICER
9 Month (DF)	987,418	6.21	-1,086,095
20-24 Month (NTP)	1,501,221	5.74	(Dominates)

Deterministic Sensitivity Analysis

Parameters were varied on different scales for high and low values and univariate sensitivity analysis was performed to see the robustness of the findings. A tornado plot has been prepared to see the effect of change in each variable. Based on the findings in one-way sensitivity analysis, input parameters were varied accordingly to see their impact in probabilistic sensitivity analysis.

In tornado plot, shown in the Figure 6.2, we find that the most influential parameters were costs of cure state of the 20-24-month regimen, cost of cure state of the



9-month regimen, costs of failure state of 9 month regimen, cost of failure state of the 20-24-month regimen according to the more effects on the results. Among the utility values utility of cure state of the 9-month regimen has more effect than utilities of other states.

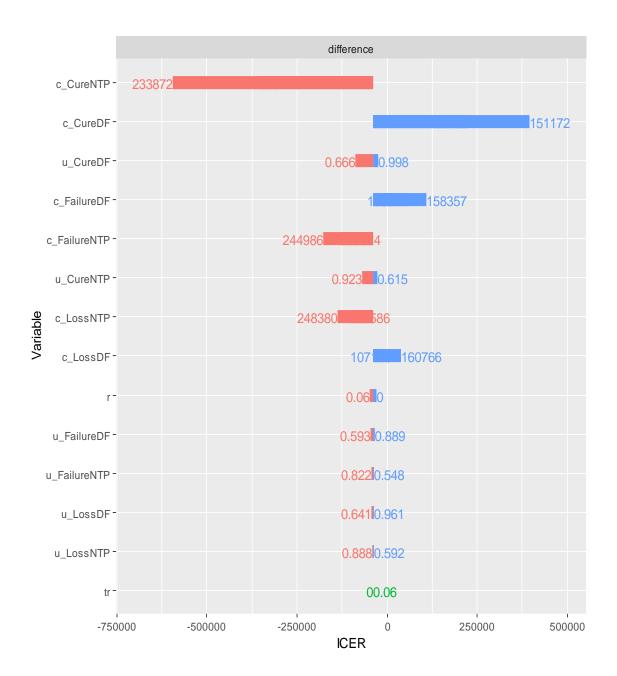


Figure 6.2 Tornado Plot of Deterministic Sensitivity Analysis



Probabilistic Sensitivity Analysis

In probabilistic sensitivity analysis, the costs values and utility values for each state were varied by 20%. At first the base case Markov model with all parameters with their values and distribution was run for 1000 iterations and the base case results were assessed in the form of incremental cost effectiveness ratio (ICER), Cost-effectiveness Acceptability Curve (CEAC), and Expected Value of Perfect Information (EVPI).

Incremental Cost-effectiveness Ratio (ICER)

From the results reported in the Table 5 shows that over life-time the 9-month regimen costs about BDT 942,315 while the 20-24-month standard regimen costs BDT 1,434,254. At the same time QALY gained by the 9-month regimen is 6.00 whereas in the 20-24-month regimen QALY is gained 5.51. Therefore, the resultant ICER is -BDT 997,257 per QALY gained. It shows that 9 month regimen dominates the 20-24-month regimen (Table 5). Results shown in the cost effectiveness plane also shows that the ICER is in the North-West Quadrant which makes the 9-month regimen dominant over the 20-24-month regimen (Figure 6.4).

Table 6.5 ICER from Probabilistic Model

Regimen	Cost Per	QALY Gained	ICER	EVPI
	Patient (BDT)			
9 Month (DF)	942,315	6.00	-997,257	13,719
20-24 Month	1,434,254	5.51	(Dominates)	
(NTP)				

Cost-Effectiveness Acceptability Curve (CEAC)

CEAC has been shown in Figure 6.3. Since the 9-month regimen dominates over the 20-24-month regimen, the CEAC for the 9-month regimen is over 0.75 form the beginning. With increasing willingness-to-pay CEAC value gets bigger. On the other



hand, it decreases for the 20-24-month regimen for higher values (Figure 6.4).

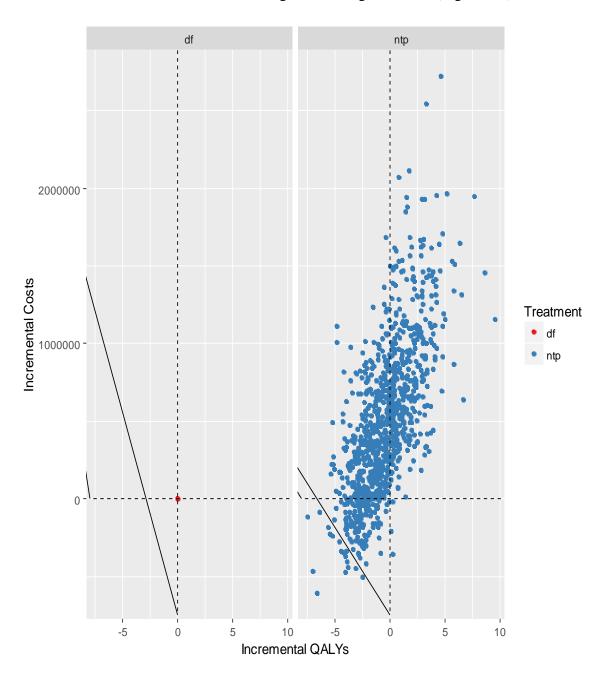


Figure 6.3 Cost-Effectiveness Plane



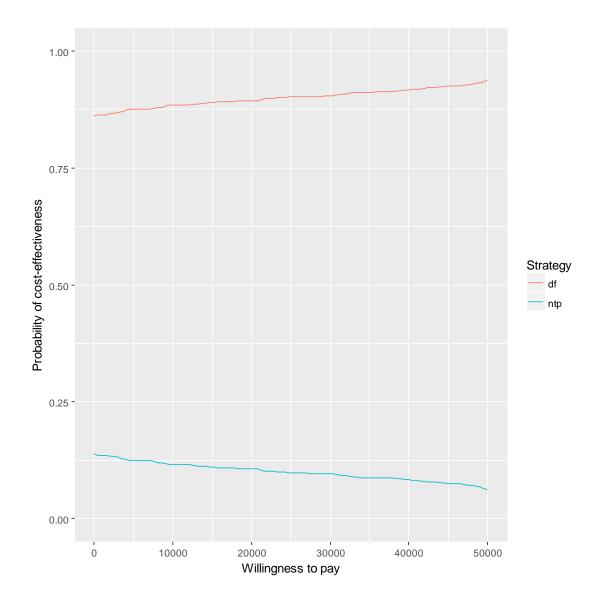


Figure 6.4 Cost-Effectiveness Acceptability Curve

Expected Value of Perfect Information (EVPI)

EVPI is the absolute limit of the value of further research that would completely eliminate the uncertainty around the parameters in the model. EVPI value of BDT 13, 719 shows that with reducing uncertainty around the parameters would require only BDT 13,719. It is also evident from the Figure 6 that with increasing willingness-to-pay the EVPI decreases and offsets the need of further research due to the small gains.



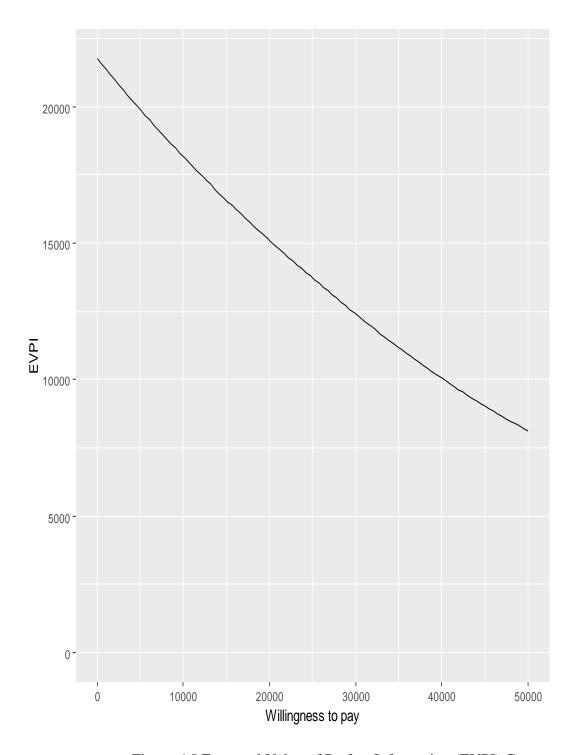


Figure 6.5 Expected Value of Perfect Information (EVPI) Curve



Discussions

The study results show that the 9-month "Bangladesh" regimen is cost-effective from a societal perspective. Bangladesh regimen has more cure rate and it provides more utility to the patient than those who undergone the 20-24-month regimen which WHO currently endorse. However, in 2012 a clinical trial named the STREAM (Standardized Treatment Regimen of Anti-Tuberculosis Drugs for Patients with MDR TB) in Ethiopia, Mongolia, South Africa, and Vietnam had been initiated (International Union Against Tuberculosis and Lung Disease (The Union), 2017). The primary objective of the first stage of the trial is to evaluate the effectiveness of the shorter "Bangladesh" regimen in other settings (Moodley & Godec, 2016). Damien Foundation initiated the Fluoroquinolone based shorter regimen and tested over 12-year period (Van Deun et al., 2010); but the study lacks in terms of it was only an observational study and more patients opted out from participation in the study; patients with HIV were not included; and cohorts were enrolled consecutively, i.e., various regimens were tested in various time periods, and cohort sizes were not predetermined (TBFACTS.ORG, 2017). Since in the published description of the trial does not mention any undertaking of costeffectiveness analysis alongside the clinical trial this study provides an important evidence of cost-effectiveness for the shorter regimen.

Several studies have also been conducted to assess the cost-effectiveness of different MDR-TB treatment regimens. Fitzpatrick et al. (2012) conducted a systemic review of studies which used primary data and outcome which eventually includes only four studies conducted in Estonia, Peru, the Philippines, and Tomsk, Russia. Cost per DALY averted with second line drugs were \$598, \$163, \$143, \$745 respectively. The



cost per DALY averted was lower than GDP per capita in all 14 WHO sub-regions considered. In other studies, Diel and colleagues showed that the treatment of MDR-TB is cost-effective in Germany (Diel, Nienhaus, et al., 2014) and European Union (EU) countries combined (Diel, Vandeputte, et al., 2014).

Our study results suggest that shorter regimen is cost-effective in Bangladesh setting. Since Bangladesh is developing country, the findings can be emulated in other developing countries. There is already evidence that MDR-TB treatment is itself cost-effective in developed countries; with shorter regimen treatment adherence will increase in efficacy of the treatment. Although STREAM trial will evaluate the incidence of adverse drug reactions among the shorter and current regimens, in this study we found that the adverse drug events were less among the patients undergoing shorter regimen. Both shorter duration of treatment and lesser adverse drug reactions translated into more QALY for the shorter regimen and eventually made the shorter regimen cost-effective.

This study has several limitations. For transitional probabilities we used one study for each treatment regimens due to lack of published study. This made the results biased. Results from the STREAM study will help to fill this void and the clinical trial results can be used for future studies. Cost and effectiveness data from the patients may suffer from the recall bias, especially for the patients under the 20-24-month regimen. Health care facility level cost data were also collected from four facilities (two for each regimen) may suffer from biased estimates.

Despite these limitations, this study tried to collect the patient and provider level data comprehensively using pre-set questionnaires. To our best knowledge, it is also the



first study to conduct economic evaluation between shorter and current regimens of MDR-TB treatment.

Growing evidence of efficacy compels the policy makers to adopt the shorter regimen as the approved regimen in near future. Results of this study makes the case for shorter regimen stronger with the evidence of cost-effectiveness which is often considered as the pivotal consideration for allocating scarce resources.

We can conclude that the evidence of cost-effectiveness of shorter regimen of MDR-TB and the efficacy of the regimen from other studies reflect that it is high time to adopt the shorter regimen as the prescribed treatment for MDR-TB treatment. This will prevent the deadly disease to spread among the vulnerable population worldwide and help us to reach the End TB strategy goals.

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APPENDIX A- STUDY III SUPPLEMENTAL INFORMATION

Table A.1 MDR-TB Drug Costs

	Short			Daily	
Drug	Form	Dose (mg)	Price	Dose (mg)	Daily dose Price
Kanamycin	Km	1000	79.25	500	0.79
Moxifloxacin	Mfx	400 mg	39	400	0.39
Prothionamide	Pto	250	13.39	500	0.27
Levofloxacin	Lfx	750	10.00	750	0.10
Cycloserine	Cs	250	28.80	500	0.58
Ethionamdie	Eto	250	7.77	500	0.16
Clofazimine	Cfz	100	103.86	100	1.04
Pyrazinamide	Z	500	17.305	1000	0.05
High dose Isoniazid (H)	Н	300	12.76	900	0.06
Ethambutol	Eto	400	20.29	400	0.03

Table A.2 Drug Costs for MDR-TB Treatment Regimes

			Duration	Price (Each	Price	
Phase	20-24-mont	h regimen	(month)	Day)	(Period)	BDT
Intensive	Z-KM-Eto-C	Cs-Ofx/Lvx	8	1.68	402.10	
Continu						
ation	Z-Eto-Cs-O	fx/Lvx	14	0.89	370.82	
				Total	772.916	61833
			Duration	Price (Each	Price	
Phase	9-month reg	gimen	(month)	Day)	(Period)	BDT
	Km-Mfx-Pto	o-Cfz-Z-				
Intensive	Hhigh-dose-	E	4	2.60	311.69	
Continu						
ation	Mfx-Cfz-Z-	E	5	1.49	222.83	
				Total	534.5187	42762



Table A.3 Diagnostic Test Costs for 20-24-Month Regimen

	Laboratory Investigations	At base line or before starting the treatment	Rate	Intensive Phase (Injectable Period- usually	in 8* Month (Investigation	During Continuation Phase (Oral Medication only-Usually 12 month***)	Total Amount in 12* Month (Investigation Rate X Frequency)
	Pure Tone Audiometry (PTA)	Must	750	Monthly	6000	No clear decision regarding this!	
2	S. Creatinine	Must	350	Monthly**	2800	Need based on symptoms / Clinical Decision**	1400
3	S. Electrolyte	Must	900	Monthly**	7200	Need based on symptoms / Clinical Decision**	3600
4	S. Bilirubin, SGPT, ALP	Must	950	Every 1-3 Monthly	2850	Need based on symptoms / Clinical Decision**	3800
5	Thyroid Function Test (TSH)	Must	900	Every 6 Monthly	900	Every 6 Monthly	1800
	Complete Blood Count (CBC)	Must	450	Need based on symptoms / Clinical Decision**	900	Need based on symptoms / Clinical Decision**	1800
7	S. Uric Acid	Must	400	Need based on symptoms / Clinical Decision**	800	Need based on symptoms / Clinical Decision**	1600
×	Random Blood Sugar (RBS)	Must	250	Need based on symptoms / Clinical Decision*	500	Need based on symptoms / Clinical Decision**	1000
9	Chest X Ray	Must	450	Every 6 Monthly	450	Every 6 Monthly	900
10	Pregnancy Test (Female at Child Bearing Age)	Must	300	Need based on symptoms / Clinical Decision**		Need based on symptoms / Clinical Decision**	
	Expenditure at Baseline/Patient		5700	Expenditure at Intensive Phase/Patient	22400	Expenditure in Continuation Phase/Patient	15900

Table A.4 Diagnostic Test cost for 9-Month Regimen

	4 Month		5 Month	Total
Intensive		Continuation		17825
Phase	(22400/8)*5	Phase	(15900/12)*5	



APPENDIX B- PATIENT QUESTIONNAIRE (DS-TB) Face Sheet for DRUG SENSITIVE TB Patient Interview

IDENTIFICATION	
DIVISION:	
DISTRICT:	
UPAZILA:	
NAME AND TYPE OF THE FACILITY:	
(Union Health Center =01, Upazila Health Complex =02, District	
Hospital =03)	
HEALTH FACILITY CODE:	
TYPE OF THE PROVIDER:	
(BRAC =01, Damien Foundation =02, Other (Please specify)=03)	
RESPONDENT:	
(Patient=01, Friend/Guardian=02, Other (Please specify)=03)	
SEX OF THE RESPONDENT:	
(Male=01, Female= 02)	
SEX OF THE PATIENT:	
(Male=01, Female= 02)	
NAME OF THE DOT PROVIDER:	



INTERVIEWER VISITS						
	1		2	3	FINAI	L VISIT
DATE						
INTERVIEWER'S NAME & CODE					RESULT CODE	
RESULT CODE*					RESULT CODE	
*RESULT CODES:						
01 COMPLETED	03 POSTPO	NED	05 PA	RTLY CO	MPLETED	
02 NOT AVAILABLE	04 REFUSE	D	96 OT	HER, SPE	CCIFY	
SUPERVISOR	FIELD EDI	ГОР	OFFICE ED	DITOR	KEYI	ED BY

Collect information from only those patients who have completed their treatment within last two months.



Section 1: Patient Information

(TO BE FILLED BY INTERVIEWER TRANSFERING INFORMATION FROM THE TB CARD)

		Options		
	ent obtained from	Yes1		
patie	nt or caregiver?	No2		
No.	Questions and filters	Coding categories	Response	Skip
101	Patient age			
			Years	
			Months	
102	Patient Sex	Male1		
102	1 attent Sex			
		Female2		
103	BCG Vaccine	No		
		Scar1		
		Scar Seen2		
104	Type of TB	Pulmonary smear positive1		
		Pulmonary smear negative2		
		Xpert MTB/RIF Positive3		
		Extra-Pulmonary4 Please Specify		

105	Type of Patient	New		1		
		Failure		2		
		Transfer in		3		
		Relapse		.4		
		Treatment afte	r loss to follo	w		
		up/default		5		
		Other		6		
		Specify				
106	Referred by which	Private Practit	ioner (Gradua	te)1		
	type of provider?	Private Practit	ioner (Non-			
		Graduate)		2		
		Govt. field sta	ff	3		
		Shasthya Sebil	_			
		field staff (NG	FS)	4		
		Village Doctor	r (VD)	5		
		Community V	olunteer	6		
		Govt. Hospital	l	7		
		Private Hospit	al	8		
		Community H				
		TB Patient				
		Other		10		
		Please specify				
107	Results of sputum	Month 0	Smear 1	Smear 2	Xpert Result	Weight (kg)
	Examination	2/3				
		3/4				
		6/8				
108	Treatment Regimen	Cat I		1		
		Cat II		2		



109	Treatment outcome	Cured1	
		Treatment completed2	
		Died3	
		Treatment failure4	
		Lost to follow	
		up/Default5	
		Transfer Out6	
		Not Evaluated7	
110	HIV Status	Positive1	
		Negative2	
		Not Tested3	
		Unknown4	
		Declined5	
111	Type of Drug		
	Reaction		
112	Date of starting		Day
	treatment		
			Month
			Year
113	Date of treatment completion		Day
			_
			Month
			Year

Section 2.0: Previous treatment

PATIENT INTERVIEW SECTION

No.	Questions and filters	Coding categories	Response	Skip
201	Have you ever had TB treatment before? CROSS-CHECK WITH INFORMATION FROM PATIENT CARD	Yes1 No2		If 2 ▶ 301
202	Have you completed your previous TB treatment? CROSS-CHECK WITH INFORMATION FROM PATENT CARD	Yes		If 1
203	Why did not you complete your previous treatment?	A. Distance to the facility B. Lack of money for treatment costs		
	[For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	C. Drug Side Effects D. Moved/Migrated E. Thought that no more treatment was necessary E. Other Please Specify		

Start time: Hours |__|_ | Minutes |__|_|

Section 3: Delay, Prediagnostic & Diagnostic Costs

No.	Questions and filters	Coding categories	Response	Skip
301	What symptoms did you experience that led you to	a. Cough		
	seek treatment for your	b. Evening rise of		
	most recent illness with TB?	temperature/low grade fever		
	ID:	c. Night sweats		
		d. Coughing up blood		
		e. Weight loss		-
	[For each option, record 1 if the option is mentioned	f. Other		_
	and record 2 if the option has not been mentioned.]	Specify		
302	How long did you experience these symptoms	a. Cough	Weeks	
	before you went to seek treatment?	b. Evening rise of temperature/low grade fever	Weeks	
	[For each option, record	c. Night sweats	Weeks	-
	number of weeks if mentioned yes in the previous question.]	d. Coughing up blood	Weeks	-
	previous question.	e. Weight loss	Weeks	
		f. Other	Weeks	
		Specify		



Sect	tion 3.1 First Visit		
803	After you experienced	Community Clinic1	
	the symptoms, which provider did you go to	Union Subcenter2	
	first?	Upazila Health Complex3	
		District Hospital4	
		Pharmacy & Drug Store5	
		Homoeopath6	
		Private Hospital7	
		Village Doctor8	
		Traditional Healer9	
		Other10	
		Specify	
304	What was the distance to the provider from your home?		Kilometer
305	What was the travel time to reach this provider?		Hour
	to reach and provider.		Minute
306	What was the waiting		Hour
	and consultation time with the provider?		Minute
307	What was the registration	IF not known put 99999	
307	(ticket) cost paid by you? In TAKA	IF no registration cost 000	
308	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000	
309	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000	



310	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
311	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
312	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
313	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
314	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	ion 3.2 Second Visit			
No.	Questions and filters	Coding categories	Response	Skip
315	Once you experience the symptoms to which provider did you go after you have seen the provider type	Community Clinic		
	(MENTION THE FIRST VISIT'S PROVIDER TYPE)	Pharmacy & Drug Store .5 Homoeopath .6 Private Hospital .7 Village Doctor .8 Traditional Healer .9 Other .10 Specify		
316	What was the distance to the provider from your home?		Kilometer	
317	What was the travel time to reach this provider?		Hour Minute	
318	What was the waiting and consultation time with the provider?		Hour Minute	
319	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
320	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
321	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



322	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
323	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
324	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
325	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
326	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	tion 3.3 Third Visit			
No.	Questions and filters	Coding categories	Response	Skip
327	Once you experience the symptoms to which provider did you go after you have seen the provider type?	Community Clinic		
	(Mention the second visit's provider type)	Pharmacy & Drug Store		
328	What was the distance to the provider from your home?		Kilometer	
329	What was the travel time to reach this provider?		Hour Minute	
330	What was the waiting and consultation time with the provider?		Hour Minute	
331	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
332	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
333	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



334	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
335	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
336	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
337	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
338	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	ion 3.4 Fourth Visit			
No.	Questions and filters	Coding categories	Response	Skip
339	Once you experience the symptoms to which provider did you go after you have seen the provider type?	Community Clinic		
	(Mention the third visit's provider type)	Pharmacy & Drug Store		
340	What was the distance to the provider from your home?		Kilometer	
341	What was the travel time to reach this provider?		Hour Minute	
342	What was the waiting and consultation time with the provider?		Hour Minute	
343	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
344	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
345	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



346	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
347	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
348	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
349	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
350	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	tion 3.5 Fifth Visit			
No.	Questions and filters	Coding categories	Response	Skip
351	Once you experience the symptoms to which provider did you go after you have seen the provider type?	Community Clinic		
	(Mention the fourth visit's provider type)	Pharmacy & Drug Store		
352	What was the distance to the provider from your home?		Kilometer	
353	What was the travel time to reach this provider?		Hour Minute	
354	What was the waiting and consultation time with the provider?		Hour Minute	
355	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
356	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
357	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



358	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
359	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
360	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
361	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
362	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



No.	Questions and filters	Coding categories	Response	Skip
363	Once you experience the symptoms to which provider did you go after you have seen the provider type? (Mention the fifth visit's provider type)	Community Clinic .1 Union Subcenter .2 Upazila Health Complex .3 District Hospital .4 Pharmacy & Drug Store .5 Homoeopath .6 Private Hospital .7 Village Doctor .8 Traditional Healer .9 Other .10 Specify		
364	What was the distance to the provider from your home?		Kilometer	
365	What was the travel time to reach this provider?		Hour Minute	
366	What was the waiting and consultation time with the provider?		Hour Minute	
367	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
368	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
369	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



370	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
371	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
372	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
373	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
374	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	tion 3.7 Seventh Visit			
No.	Questions and filters	Coding categories	Response	Skip
375	Once you experience the symptoms to which provider did you go after you have seen the provider type?	Community Clinic		
	(Mention the sixth visit's provider type)	Pharmacy & Drug Store		
376	What was the distance to the provider from your home?		Kilometer	
377	What was the travel time to reach this provider?		Hour Minute	
378	What was the waiting and consultation time with the provider?		Hour Minute	
379	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
380	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
381	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



382	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
383	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
384	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
385	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
386	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Section 3.8 Eighth Visit					
No.	Questions and filters	Coding categories	Response	Skip	
387	Once you experience the symptoms to which provider did you go after you have seen the provider type?	Community Clinic			
	(Mention the seventh visit's provider type)	Pharmacy & Drug Store			
388	What was the distance to the provider from your home?		Kilometer		
389	What was the travel time to reach this provider?		Hour Minute		
390	What was the waiting and consultation time with the provider?		Hour Minute		
391	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000			
392	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000			
393	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000			



394	What was the cost for x-ray? In TAKA	IF not known put 99999	
		IF no cost for x-ray 000	
395	What was the costs of	IF not known put 99999	
	drugs? In TAKA	IF no cost for drugs 000	
396	What was travel costs?	IF not known put 99999	
	(INCLUDE ALL	IF no travel cost 000	
	TRAVEL RELATED		
	COSTS: RETURN		
	TRAVEL, TRAVEL		
	FOR LABORATIRY		
	TESTS, DTUGS,		
	COST FOR		
	ACCOMPANYING		
	PERSONS) In TAKA		
397	What was the food	IF not known put 99999	
	costs? In TAKA	IF no food cost 000	
		ir no rood cost ooo	
398	What was cost for	IF not known put 99999	
	accommodation?	IF no accommodation cost 000	



Section 4.0: Treatment Costs

Sect	Section 4.1 Cost Related to DOT			
No.	Questions and filters	Coding categories	Response	Skip
401	From where did you get your TB drugs?	Health facility 1 Home 2 Community 3 Pharmacy 4 Workplace 5 Other 9 Specify		
402	How often do you travel to the health facility / hospital for picking up your TB drugs?		Times/month	
403	How long does it take to go to the place for picking up the drugs (ONE WAY)?	A. On Foot	Hour Minute	
		B. By Transport	Hour Minute	
404	How long does one of these visits take on average, including time on the road and waiting time (total turnaround time)?		Hour Minute	
405	From your home to the facility, how much does it cost if you take transport? (both ways)	IF not known put 99999		



406	If you go to a facility to pick up your drugs, how much do you spend on food on that day? (on the road, while waiting, lunch etc.)	IF not known put 99999		
407	Do you have to pay administration fees when picking up your TB drugs?	Yes1 No2		If 2 ▶ 409
408	What was the administration cost?	IF not known put 99999 IF no administration cost 000		
409	Do you have to pay for accommodation when picking up your TB drugs?	Yes1 No2		If 2 ▶ 411
410	What was the accommodation cost?	IF not known put 99999 IF no accommodation cost 000		
Sect	ion 4.2 Cost related t	to Follow Up Tests		
411	Did you ever have to go to the health facility in addition to your regular visits for follow up tests since the beginning of treatment?	Yes1 No2		If 2 ▶ 414
412	If yes, how many times?		Times	
413	If yes, did you have to pay any additional costs any time during the entire period?	Yes1 No2		If 2 ▶ 421
414	If so, what kind of costs			



	pay last time (In		
	TAKA)? IF not known put	B. Sputum Test	
		C. X -ray	
	99999 IF no cost put 000	D. TB Drugs	
		E. Other Drugs	
		F. Others	
415	How long does one of these follow-up visits take on average, including time on the		Hour Minute
	road, waiting time and tests (total turnaround time)?		

Section 5.0: Cost related to accompanied persons (Friends/Guardian)

No.	Questions and filters	Coding categories	Response	Skip
501	Does any family/friend/DOT supporter accompany you on any visits before diagnosis and/or during diagnosis?	Yes		If 2 ▶ 507
502	If YES, on how many visits has your family/friend/DOT supporter accompanied you or gone with you before diagnosis and/or during diagnosis?		Times	
503	What was the cost for pre- diagnosis/diagnosis visits of	A. Transport Cost		



		1		
	your accompanying person in Taka)?	B. Food Cost		
		C. Accommodation Cost		
	IF not known put 99999			
	IF no cost put 000			
504	Does your accompanying person earn?	Yes1		If 2
		No2		•
				506
505	If earn, how much the person earn per day (in			
	TAKA)?			
	IF not known put 99999			
506	Why did someone	A. Distance		
	accompany you?	B. Security		
		C. Administrative barrier		
	[For each option, record 1 if the option is mentioned	D. Too ill to travel alone		
	and record 2 if the option has not been mentioned.]	E. Was required for treatment		
		F. Other		
		Please		
		Specify		
507	Does any family/friend/DOT	Yes1		If 2
	supporter accompany you	No2		•
	on any visits during treatment (taking drugs)?			601
508	If YES, on how many visits		Times	
2 3 3	has your family/friend/DOT		Times	
	supporter accompanied you or gone with you during			
	treatment?			
509	What was the cost for visits of your accompanying	A. Transport Cost		
	or your accompanying			



	person during your treatment in TAKA)?	B. Food Cost	
		C. Accommodation Cost	
	IF not known put 99999		
	IF no cost put 000		
510	Does your accompanying person earn?	Yes	If 2
F11	TC 1. (1.		512
511	If earn, how much the person earn per day (In TAKA)?		
512	Why did someone accompany you?	A. Distance	
	1	B. Security concern	
	[For each option, record 1	C. Administrative barrier	
	if the option is mentioned and record 2 if the option	D. Too ill to travel alone	
	has not been mentioned.]	E. Was required for treatment	
		F. Other	
	IF not known put 99999	Please Specify	

Section 6.0: Hospitalization Costs

No.	Questions and filters	Coding categories	Response	Skip
601	Have you been hospitalized before (but due to TB) or	Yes1		If 2
	during your TB treatment?	No2		•
				701
602	If YES, how many times were you hospitalized for this illness?		Times	
603		A. Hospital Administration Fees		



	How much did you pay during your last stay at the	B. Hospital stay charges		
	hospital (In TAKA)?	C. Food (Not provided by the hospital)		
	IF not known put 99999	D. Transport (Both Ways)		
	IF no cost put 000	E. Drugs		
		F. Diagnostic Tests		
		G. Others		
604	Did any attendant/caregiver stay with you at the	Yes1		If 2
	hospital?	No2		•
				701
605	If YES, how many days he/she stay with you (sleep there)?		Days	
606	Were there any extra costs for your relative/friend for staying at the hospital?	Yes		If 2 ► 506
607	What was the cost for the accompanying person during your hospitalization (stayed at night)? In TAKA	A. Transport Cost B. Food Cost		
		C. Accommodation Cost		
	IF not known put 99999	D. Other Cost		
608	Does your accompanying person earn?	Yes1		If 2 /3
		No		610



609	If earn, how much the accompanying person earn per day?			
610	Did any other family/friend visit you while in hospital?	Yes1	I	If 2
	visit you willie ili ilospitar:	No2		•
				701
611	If YES, How many people visited you?		Days	
612	On an average, how many times did each of these person visit you?		Times	
613	What was the cost for EACH VISIT of them (IN	A. Transport Cost		
	TAKA)?	B. Food Cost		
	IF not known put 99999	C. Accommodation Cost		
		D. Other Cost		
614	How long were the visits including traveling time?		Hour	
			Minute	

Section 7.0: Other Costs, Other Illnesses and Coping Costs

Sect	Section 7.1 Other Costs					
No.	Questions and filters	Coding categories	Response	Skip		
701	Did you buy any supplements for your diet because of the TB illness, for example vitamins, meat, energy drinks, soft drinks, fruits or medicines?	Yes		If 2 ▶ 703		
702	If YES, how much did you spend approximately on	A. Meat				



	each of these items each month (in TAKA)?	B. Fish				
		C. Fruits				
	IF not known put 99999	D. Drinks				
		E. Vegetables				
		F. Vitamins/Herbs				
		G. Others				
Sect	tion 7.2 Other illnesses					
703	Do you have any chronic	Yes1	Т			If 2
	illness for which you are receiving treatment?	No2				•
						707
704	If YES, which disease do you have?	A. Diabetes				
	you have.	B. Heart Disease				
		C. High Blood Pressure				
	[For each option, record 1] if the option is mentioned	D. Cancer				
	and record 2 if the option has not been mentioned.]	E. Arthritis				
		F. Other				
		Specify				
		G. Other				
		Specify				
705	Are there any additional	Yes1				If 2
	costs for you because of this other illness besides	No2]	•
	the costs that you have already mentioned?					707
706	If YES, how much are	A. Drugs				
	these additional costs on average per month? In	D. Diagnostic tests	$\downarrow \downarrow$			
	TAKA	B. Diagnostic tests				



		C. Transport	
	IF not known put 99999	D. Fees	
		E. Others	
707	How much did you spend	IF not known put 99999	
707	on healthcare on average per month BEFORE the TB illness? In TAKA	If not known pat 3333	
708	How much did you spend on healthcare on average per month AFTER the TB illness? In TAKA	IF not known put 99999	
Sect	tion 7.3 Coping Costs		
709	Has your illness with TB	Yes1	
	resulted in a financial burden?	No2	
710	Did you borrow any	Yes1	If 2
	money to cover costs due to the TB illness?	No2	▶ 715
			/13
711	If YES, how much did you borrow? In TAKA	IF not known put 99999	
712	From whom did you borrow?	A. Family	
		B. Neighbor	
	[For each option, record 1 if the option is mentioned	C. Friend	
	and record 2 if the option has not been mentioned.]	D. Bank	
		E. Cooperative	
		F. NGO	
		G. Money lender	
		H. Others	
		Specify	



=10		T **	T	700
713	Have you already paid back the borrowed	Yes1		If 2
	money?	No2		>
				716
				710
714	How are you planning to	In		If 2
	pay back the money?	Full1		•
		In Installment2		716
				716
715	If you are paying in	IF not known put 99999		
	installment, what is the monthly installment? In			
	TAKA			
	**	***		¥0.0
716	Have you sold any of your property to finance the	Yes1		If 2
	cost of the TB illness?	No2		•
				718
				710
717	How much money did you get from the sale of each	A. Land		
	of these properties?	D.Y.I.		
	1 1	B. Livestock		
		C. Transport/Vehicle		
		C. Transport/venicle		
	IF not known put 99999	D. Household item		
	1			
		E. Farm produce		
		1		
		F. Jewelry		
		G. Savings (FDR)		
		H. Other		
		Specify		
=10				10.0
718	Have you employed any household help for your	Yes1		If 2
	illness?	No2		•
				801
				501



7	19	How much do you pay	IF not known put 99999				
		monthly to the household					
		help? In TAKA					
		_					

Section 8.0 Patient Income and Patient's Household Income

Secti	ion 8.1 Personal Income	2		
No.	Questions and filters	Coding categories	Response	Skip
801	Who is the primary income earner of the household?	Patient		
802	What is the highest level of Education (In years) of these individuals? [For each option, record the number of years spent studying. if the person is illiterate/did not go to school record "0"]	A. Patient B. Primary Income Earner (If Other than patient) C. Household Head (If Other than patient) B. Spouse of Household Head (If Other than patient)		
803	Are you involved in income earning activities?	Yes, formal work		If 4 ▶ 809

804	If No, why are you not involved in any income earning activities?	Cannot work due to illness1 Stopped working after contracting TB		
805	Have you left your job due to your TB illness?	Yes1		If 2
		No2		•
				809
806	If YES, how many months ago did you leave your job?		Months	
807	What was your monthly income when you were working? In TAKA	IF not known put 99999		
808	How regularly did you work before you became ill with TB?	Throughout the year		

809	What was your main occupation before your illness with TB?	Service	
810	Did you have to change jobs when you became ill with TB?	Yes1 No2	
811	What is your main occupation after your illness with TB?	Service	
812	How many hours did you work on average per day BEFORE you became ill with TB?		Hours
813	How many hours do you work on average NOW per day?		Hours



814	If answer to 812 differs	Yes1	If 2
	from answer to 813:Is the		
	change related to the TB	No2	
	illness?		818
			010
815	What was your estimated	IF not known put 99999	
	personal income per	-	
	month BEFORE the TB		
	illness? In TAKA		
816	What is your estimated	IF not known put 99999	
	personal income per		
	month NOW? In TAKA		
817	Is someone doing the	Chausa	
017	Is someone doing the work you used to do after	Spouse1	
	your illness?	Son2	
		Daughter3	
		Friend4	
		Nobody5	
		Other9	
		Specify	
818	Do you have children of	Yes1	If 2
	or below school age?		_
		No2	•
			822
			622
819	Are all of your children	Yes1	If 1
	attending school		
	regularly?	No2	•
			922
			822
820	Did your children go to	Yes1	
	school regularly before		
	your recent illness with	No2	
	TB?		
821	If your children do not go	A. Needs to help around the house	
	to school, indicate the	B. No money for school fess	
	reasons for not attending	B. No money for school less	
	school regularly?	C. Also sick	
		D. Has to work to earn	



	[For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	E. Take care of patient F. Other Specify	_
822	Has the TB illness affected your social or private life in any way?	Yes1 No2	If 2 • 823
823	If YES, how was your social life affected? [For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	A. Divorce B. Loss of job C. Dropped out of school D. Separated from spouse E. Disruption of sexual life F. Sick child G. Other Specify	
8824	What is your religion?	Islam	
Secti	on 8.2 Household Incor	me	
825	How much do you estimate was the average income of your household per month BEFORE the TB illness? (for all persons in the house, including patient; includes welfare payments, government assistance or	A. Patient's income B. Income of rest of the household C. Govt. assistance	



	other social support)? In TAKA	D. Other	
	IF not known put 99999		
826	How much do you estimate was the average income of your household per month NOW? (for all persons in the house, including patient; includes	A. Patient's income B. Income of rest of the household D. Govt. assistance	
	welfare payments, government assistance or other social support)? In TAKA	E. Other	
	IF not known put 99999		
827	How many people regularly sleep in your house? (including patient)		
828	How many members of the household are employed for wage/salary? (including patient)		
829	Besides yourself, does anyone else of your household receive treatment for TB?	Yes1 No2	If 2 ▶ 823
830	If YES, how many household members are suffering from TB?		

Section 9.0 Socioeconomic Indicators

Secti	ion 9.0 Questions ab	out the household	RESPONSES	Skip
901	What is the main source of lighting for the household?	Electrical mains 01 Solar electricity 02 Generator 03 Gas 04 Kerosine/Oil lamp 05 Candles / Torch 06 Coleman (Kerosene/Pressure lanterns) 07 Improvised lamp 08 Others 96 Specify 96		
902	What is the main source of water for drinking and food preparation for the household?	No source of lighting 98 Piped into household 01 Piped to yard/plot 02 Piped into neighborhood 03 Protected well 04 Unprotected well 05 Protected Spring 06 Unprotected Spring 07 River/stream 08 Pond/lake/dam 10 Communal tank 11 Rainwater 12 Tank truck / Water cart 13 Bottled water 14 Other 96 (Specify) Not reported 99		
903	What type of fuel is mainly used in your household for cooking?	Gas		

		Electricity06	
		Biogas07	
		Straw/shrubs/grass	
		Saw dust09	
		Others96	
		Specify	
904	What kind of	Own flush toilet01	
	toilet facility do	Shared flush toilet02	
	people in your	Ventilated improved pit latrine03	
	house mainly	Pit latrine with slab04	
	use?	Pit latrine without slab / open pit05	
	use.	Bowl/Bucket system06	
		Closet oversea/river	
		No facility/bush/seashore	
		Other	
		Not reported	
905	Main	Natural Floor:	
700	material of the	Earth	
	floor for the	Sand12	
	principal	Rudimentary:	
	residence	Wood planks21	
	structure	Palm/bamboo22	
		Finished:	
		Polished wood31	
		Vinyl/asphalt strips32	
		Marble/Ceramic tiles33	
		Floor tile	
		Cement / Concrete/35	
		Brick	
		Unpolished 38	
		Other96	
		(Specify)	
005		Not reported 99	
906	Main	Natural roofing	
	material of the roof of the	No roof	
	principal	Rudimentary roofing	
	residence	Bamboo21	
		Wood planks	
		Cardboard23	
		Finished roofing	
		Tin / Metal31	
1			Į.
		Wood	



		Cement / Concrete		
		(Specify)		
007	Main	Notarial Walls		1
907	material of the	Natural Walls No walls11		
	exterior walls of	Cane/Palm/Trunks		
	principal	Dirt		
	residence	Rudimentary walls		
		Bamboo with mud / matting21		
		Stone with mud22		
		Mud23		
		Fibro24		
		Plywood25		
		Cardboard26		
		Finished walls		
		Tin / Metal sheets31		
		Cement / Concrete32		
		Brick31		
		Stone with lime/cement		
		Bricks		
		Wood planks/shingles35		
		Other96		
908	What type of fuel	(Specify) ELECTRICITY		If 9:
900		ELECTRICITY		11 9,
	does your household mainly	LPG02		
		LI G		
	use for cooking?	NATURAL GAS		911
		BIOGAS		
		KEROSENE		
		COAL, LIGNITE		
		CHARCOAL07		
		WOOD		
		STRAW/SHRUBS/GRASS 09		
		AGRICULTURAL CROP		
		ANIMAL DUNG		
		NO FOOD COOKED		
		IN HOUSEHOLD95		
		OTHER 96		
		(SPECIFY)		
	i .	·	1	



909	Is the cooking usually done in the house, in a separate building, or outdoors?	IN THE HOUSE	
910	Do you have a separate room which is used as a kitchen?	Yes1 No2	
911	Does your household own any homestead?	Yes1 No2	

912	Does your household or any members	ber of the household own the items?	Yes	No	Responses
	Electricity connection?	A. Electricity	1	2	
	Solar Electricity?	A. Electricity	1	2	
	A radio?	C. Radio	1	2	
	A television?	D. Television	1	2	
	A mobile phone?	E. Mobile phone	1	2	
	A non-mobile phone?	F. Non-mobile phone	1	2	
	A refrigerator?	G. Refrigerator	1	2	
	A DVD/VCD player?	H. DVD/VCD player	1	2	
	An electric fan?	I. Fan	1	2	
	An Almirah/wardrobe?	J. Almirah	1	2	
	A water pump	K. Water Pump	1	2	
	An IPS?Generator	L. Generator/IPS	1	2	
	An air conditioner	M. AC	1	2	
	A computer/laptop?	N. Computer/Laptop	1	2	
913	In your opinion, is your	High income		1	
	household a high income household, a middle income	Middle income		2	
	household or a poor household?	Low income	3		
914	How many people live in your household?	Total household size			
915	How many adult members (15 years or older)	Number of adults in the household			
916	How many children (less than 15 years)	Number of children in the household			



917	How many separate sleeping rooms are there for the use of your household members in your residence?	Number of rooms	
918	What is your current place of residence?	Urban. 1 Urban slum. 2 Rural. 3 Other. 4 Specify	
919	If the government could provide you with some service to ease the burden of TB on you and your household, what would you prefer to have? DO NOT READ. INDICATE THE ITEMS MENTIONED. IF NOT IN THE LIST, ADD ANY NEW GOVT. SERVICE MENTIONED.	Transport vouchers	
920	How much would you be willing to pay for not becoming ill with TB in the first place? In TAKA		



Section 10.0 Performance of the Facility

The next part of the survey is about the quality of TB care that you received during your visits to this facility. Please answer the questions in this part of the survey about this facility only. Do not include any other facilities in your answer.

No.	Questions and filters	Coding categories	Response	Skip
Secti	on 10.1 Availabilit	y of TB Services		
1001	Are the waiting time(s) before being served by health providers of this facility acceptable to you?	Never		
1002	How often are you attended to by the same health providers in this facility?	Never		
1003	How often are the service hours of this facility inconvenient for you to get your TB treatment?	Never		
1004	How often are drugs not available when you require them?	Never		
1005	How often do you experience difficulties in obtaining TB services in this facility because of language barriers?	Never		



1	.006	How often do you have to go to	Never1	
		another health unit	Sometimes2	
		for TB services or treatment?	Usually3	
			Always4	
1	007	Is this health	Never1	
		facility easy to reach (distance)?	Sometimes2	
			Usually3	
			Always4	
1	008	How often are TB	Never1	
		services available during the working	Sometimes2	
		hours of this	Usually3	
		facility?	-	
			Always4	
1	.009	How often are the	Never1	
		relevant health providers you come	Sometimes2	
		to see in this facility	Usually3	
		available?	-	
			Always4	
S	Secti	on 10.2 Communi	cation and Information	
1	.010	Do the health	Yes1	
		providers in this facility tell you	No2	
		when you stop		
		spreading TB to others?		
L				
1	011	Do the health providers in this	Yes1	
		facility tell you that	No2	
		TB can be cured?		
1	012	Do the health	Yes1	
		providers in this facility tell you	No2	
		about the		
		importance of		
- 1		observed treatment?		



1013	Do the health providers in this facility tell you about the side effects of TB drugs?	Yes		
1014	Do the health providers in this facility tell you about the need for sputum tests at given points during your treatment schedule?	Yes1 No2		
1015	Do the health providers in this facility tell you about the duration of the TB treatment?	Yes1 No2		
1016	During your visits to this facility, do health providers tell you about how to store your drugs obtained for your treatment?	Yes		
1017	Does the health provider in this facility tell you when next to come back for TB services?	Yes1 No2		
Secti	on 10.3 Patient – I	Provider interaction and coun	nselling	
1018	During your visits to this facility, how often does the health provider treat you with respect?	Never		



		1	
1019	During your visits to this facility, how	Never1	
	often does the	Sometimes2	
	health provider listen carefully to	Usually3	
	you?	Always4	
1020	0 3	Never1	
	to this facility, how often do health	Sometimes2	
	providers explain things in a way you	Usually3	
	can understand?	Always4	
1021	8 7	Never1	
	to this facility, how often do you have	Sometimes2	
	sufficient time to discuss your	Usually3	
	problems?	Always4	
1022	0,	Never1	
	to this facility, how often do health	Sometimes2	
	providers discuss with you how to	Usually3	
	deal with your problems?	Always4	
1023	0,	Never1	
	to this facility, how often do you	Sometimes2	
	experience discrimination	Usually3	
	because you have TB?	Always4	
1024	0,	Never1	
	to this facility, how often is your	Sometimes2	
	privacy respected during	Usually3	
	examination?	Always4	



1025	Do health providers at this facility tell	Never	
	you how TB can affect your every day life?	Usually	
		Always4	
Secti	on 10.4 Infrastruc	ture	
1026	How often is this facility clean?	Never1	
		Sometimes	
		Always4	
1027	How often is there	Never1	
	safe drinking water in this facility?	Sometimes2	
		Usually3	
		Always4	
1028	How often are the toilets in this	Never1	
	facility usable?	Sometimes2	
		Usually3	
		Always4	
1029	How often do you find enough	Never1	
	comfortable places	Sometimes2	
	to sit on in this facility?	Usually3	
		Always4	
Secti	on 10.5 Profession	al Competence	
1030	Does this facility offer services to	Yes1	
	examine your sputum?	No2	



1031	Does this facility offer home based	Yes1	
	TB treatment?	No2	
1032	Were you physically examined during your first visit to this health facility?	Yes	
1033	Was your sputum examined when you were diagnosed with TB?	Yes	
1034	How many working days were there between your first sputum submission and the time you got your results?	0-2 Working days	
1035	In case of germs in your sputum that cause TB, were your close contacts examined by the TB facility?	Yes	
1036	How often is there a treatment observer checking on your daily intake of TB drugs?	Never	
Section	on 10.6 Affordabil	iity	

1037	How often do you have to pay for your	Never1		
	regular TB services	Sometimes2		
	(e.g. sputum tests, TB-drugs, X-rays,	Usually3		
	etc.)?	Always4		
1038	How often do you have to pay a tip in	Never1		
	order to receive TB services?	Sometimes2		
	services?	Usually3		
		Always4		
1039	How often do costs (e.g. transport)	Never1		
	prevent you from	Sometimes2		
	getting to the health facility?	Usually3		
		Always4		
Secti	on 10.7 Support			
	l ==	3.7	1	
1040	How often do you	Never1		
1040	receive transport support from the	Sometimes		
1040	receive transport			
1040	receive transport support from the	Sometimes2		
1040	receive transport support from the health facility? How often do you	Sometimes		
	receive transport support from the health facility? How often do you receive food support from the	Sometimes		
	receive transport support from the health facility? How often do you receive food	Sometimes .2 Usually .3 Always .4 Never .1		
	receive transport support from the health facility? How often do you receive food support from the	Sometimes .2 Usually .3 Always .4 Never .1 Sometimes .2		
	receive transport support from the health facility? How often do you receive food support from the health facility? How often do you	Sometimes .2 Usually .3 Always .4 Never .1 Sometimes .2 Usually .3		
1041	receive transport support from the health facility? How often do you receive food support from the health facility? How often do you receive financial assistance from the	Sometimes .2 Usually .3 Always .4 Never .1 Sometimes .2 Usually .3 Always .4		
1041	receive transport support from the health facility? How often do you receive food support from the health facility? How often do you receive financial	Sometimes .2 Usually .3 Always .4 Never .1 Sometimes .2 Usually .3 Always .4 Never .1		
1041	receive transport support from the health facility? How often do you receive food support from the health facility? How often do you receive financial assistance from the	Sometimes .2 Usually .3 Always .4 Never .1 Sometimes .2 Usually .3 Always .4 Never .1 Sometimes .2		



1043	Does the health provider talk to you the same way you are spoken to when you receive services other than TB?	Never	
1044	Does the health provider welcome you into the health facility when you visit for TB services?	Never	
1045	Does the health provider turn his/her face away when speaking with you?	Never	
1046	Do you feel that you are treated with dignity when you visit the health facility?	Never .1 Sometimes .2 Usually .3 Always .4	

Section 11.0 Quality of Life

Section 11.1 Health Related Quality of Life (FACIT-TB Questionnaire)

How would you describe your level of feeling for the following statements: (record the number in the right column)		No t At All	Slight ly	Moderately	Quite A Bit	Extrem ely	Response
A. Physical well-being							
1101	I feel ill	1	2	3	4	5	
1102	I get tired easily	1	2	3	4	5	



1103	I have a lack of energy	1	2	3	4	5	
1104	I have pain	1	2	3	4	5	
1105	I feel weak all over	1	2	3	4	5	
1106	I feel fatigued	1	2	3	4	5	
1107	I have been short of breath	1	2	3	4	5	
1108	I have nausea-"a sense of vomiting outbreak"	1	2	3	4	5	
1109	Because of my physical condition, I have trouble meeting the needs of my family	1	2	3	4	5	
1110	I am bothered by fever (episode of high body temperature)	1	2	3	4	5	
1111	I am forced to spend time in bed	1	2	3	4	5	
1112	I have discomfort or pain in my stomach area	1	2	3	4	5	
1113	I have had itching	1	2	3	4	5	
1114	I have a loss of appetite	1	2	3	4	5	
1115	I have been coughing	1	2	3	4	5	
1116	I am bothered by side effects of treatment	1	2	3	4	5	
1117	Dusts Worsen my symptoms	1	2	3	4	5	
В.	Social and economic well-being						
1118	I feel close to my friends	1	2	3	4	5	
1119	I get emotional support from my family	1	2	3	4	5	
1120	I am satisfied with my family communication about my illness	1	2	3	4	5	
1121	My family has accepted my illness	1	2	3	4	5	
1122	I feel close to my partner (or the person	1	2	3	4	5	



	· · ·	I	ı	T		-	T
	who is my main						
	support)						
1123	I get support from my friends	1	2	3	4	5	
1124	My physical condition and/or medical treatment cause me financial difficulties	1	2	3	4	5	
C.	Emotional well-						
	being/Stigma of having						
	ТВ						
1125	I worry that my condition will get worse	1	2	3	4	5	
1126	I worry about dying	1	2	3	4	5	
1127	I am concerned about what the future holds for me	1	2	3	4	5	
1128	I am embarrassed by my illness	1	2	3	4	5	
1129	It is hard to tell other people about my infection	1	2	3	4	5	
1130	I am losing hope in the fight against my illness	1	2	3	4	5	
1131	I am bothered by the change in weight	1	2	3	4	5	
1132	I worry about spreading my infection	1	2	3	4	5	
1133	I feel nervous	1	2	3	4	5	
1134	I feel sad	1	2	3	4	5	
1135	I am satisfied with how I am coping with my illness	1	2	3	4	5	
D.	Functional well-being						
1136	I am content with the quality of my life right now	1	2	3	4	5	
1137	My work (include work at home) is fulfilling	1	2	3	4	5	
1138	I am able to work (include work at home)	1	2	3	4	5	
1139	I am able to enjoy life	1	2	3	4	5	
1140	I am enjoying the things I usually do for fun	1	2	3	4	5	



1141	I have accepted my illness	1	2	3	4	5	
1142	I am sleeping well	1	2	3	4	5	
E. Spiritual well-being							
1143	I find strength in my faith or spiritual belief	1	2	3	4	5	
1144	My illness has strengthened my faith or spiritual belief	1	2	3	4	5	
1145	My life is still productive	1	2	3	4	5	

Section 11.2 EQ-5D-5L Questionnaire

Under each heading, please tick the ONE box that best describes your health TODAY.

Sl	Category	Options	Response
1146	Mobility	I have no problems in walking about	
1147	Self-Care	I have no problems washing or dressing myself	



1148	Usual Activities (e.g., work, study, housework, family or leisure activities)	I have no problems doing my usual activities	
1149	Pain/Discomfor t	I have no pain or discomfort	
1150	Anxiety/Depres sion	I am not anxious/depressed	



Section 11.3 SF-6D Questionnaire Under each heading, please tick the ONE box that best describes your health TODAY. SI Category **Options** Respon se 115 Physical My health does not limit me in vigorous Functioning activities.....1 My health limits me a little in vigorous activities.....2 My health limits me a little in moderate activities.....3 My health limits me a lot in moderate activities.....4 My health limits me a little in bathing and dressing.....5 My health limits me a lot in bathing and dressing.....6 115 Role I have no problems with your work or other regular daily 2 Limitation activities as a result of your physical health or any emotional problems.....1 I am limited in the kind of work or other activities as a result of your physical health.....2 I accomplish less than you would like as a result of emotional problems.....3 You are limited in the kind of work or other activities as a result of your physical health and accomplish less than you would like as a result of emotional problems......4 Social My health limits my social activities none of the 115 Functioning time.....1 3 My health limits my social activities a little of the time.....2



		My health limits my social activities some of the	
		time3	
		My health limits my social activities most of the	
		time4	
		My health limits my social activities all of the	
		time5	
115	Pain	I have no	
4		pain1	
		I have pain, but it does not interfere with my normal work (both	
		outside the home and	
		housework)2	
		I have pain that interferes with your normal work (both outside the	
		home and housework) a little	
		bit3	
		I have note that interfered with your normal youls (both outside the	
		I have pain that interferes with your normal work (both outside the home and housework)	
		moderately4	
		I have pain that interferes with your normal work (both outside the	
		home and housework) quite a bit5	
		I have pain that interferes with your normal work (both outside the	
		home and housework)	
		extremely6	
115 5	Mental Health	I feel tense or downhearted and low none of the time1	
3		ume1	
		I feel tense or downhearted and low a little of the	
		time2	
		I feel tense or downhearted and low some of the	
		time3	
		I feel tense or downhearted and low most of the time4	
		unic4	
		I feel tense or downhearted and all of the	
		time5	
115	Vitality	I have a lot of energy all of the	
6		time1	
		I have a lot of energy most of the	



	time2	
	I have a lot of energy some of the time	
	I have a lot of energy a little of the time4	
	I have a lot of energy none of the time5	

Section 11.4 Visual Analog Scale

1157 Visual Analog Scale Score

1158	Comments by interviewer on the interview
1159	INTERVIEWERS: CHECK YOUR FILLED IN QUESTIONNAIRE CAREFULLY
110)	BEFORE LEAVING THE RESPONDENTS AND END YOUR INTERVIEW BY
	GIVING THANKS TO THE RESPONDENT.
	RECORD THE END TIME OF THE Hour
	INTERVIEW: Minutes



APPENDIX C – PATIENT QUESTIONNAIRE (MDR-TB) Face Sheet for MULTI-DRUG RESISTANT TB(MDR-TB) Patient Interview

IDENTIFICATION		
DIVISION:		
DISTRICT:		
UPAZILA:		
NAME AND TYPE OF THE FACILITY:		
(NIDCH =01, District Chest Hospital =02, Damien Foundation Hospital		
=03, Other=04, Please Specify)		
HEALTH FACILITY CODE:		
DRUG REGIMEN FOLLOWED:		
(21 MONTH REGIMEN =01, 9 MONTH REGIMEN =02)		
RESPONDENT:		
(Patient=01, Friend/Guardian=02, Other (Please specify)=03)		
SEX OF THE RESPONDENT:		
(Male=01, Female= 02)		
SEX OF THE PATIENT:		
(Male=01, Female= 02)		
NAME OF THE DOT PROVIDER:		

	INTERVI	EWER VISITS			
	1	2	3 FINAL VISIT		
DATE					
INTERVIEWER'S NAME & CODE			RESULT CODE		
RESULT CODE*			RESULT CODE		
*RESULT CODES:	·				
01 COMPLETED 03 POSTPON		NED 05	05 PARTLY COMPLETED		
02 NOT AVAILABLE 04 REFUSED 96			OTHER, SPECIFY		
SUPERVISOR	FIELD EDITO	R OFFICE EDITOR	KEYED BY		

Take information for only those patients (>18 years of age) who are undergoing the treatment or have completed their treatment within last two months.



Section 1: Patient Information

(TO BE FILLED IN BY THE INTERVIEWER WITH HELP OF PATIENT TB CARD; FILL IN ALSO IF INTERVIEW IS REFUSED FRO NON-RESPONSE ANALYSIS)

		Options	
Consent obtained from patient or caregiver?		Yes1 No2	
No .	Questions and filters	Coding categories	Response
10	Patient age		Years
10 2	Patient Sex	Male1 Female2	
10 3	Site	Pulmonary	
10 4	History of contact with TB/ DR TB Patient	Yes1 No2	
10 5	IF YES, Relation and duration	WRITE DOWN FROM PATIENT CARD	
10 6	Medical Diagnosis other than TB	WRITE DOWN FROM PATIENT CARD	



10	Registration Status	CAT I	Non converter	1	
7		CAT I Failure2			
		Treatment After loss to follow-up- CAT			
		CAT I Relapse4			
		CAT I Non converter5			
		CAT II Failure6			
		Treatment After loss to follow-up- CAT			
			Relapse		
		Close contact of DR TB with			
		S/S			
		Transfer In			
			nary-clinically diag		
			eviously treated ulmonary new/pre		
		Extra pulmonary, new/previously treated13			
			Unknown TB treatment history14		
			ulmonary), bacter	•	
			confirmed15		
10	Type of Resistance	MDR T	TB/XDR		
8		TB/Pol	yresistance	1	
			•		
		Monoresistance2			
10	Previous TB Treatment	No.	Start Date	Regimen	Outcome
9	Episode including DR TB				
		Drugs:			
		Direct I	Line Drugs:	Casand Lina Day	~~.
				Second Line Drug Km=Kanamycin	gs.
		H=Isoniazid Km=Kanamycin R=Rifampicin Ofx=Ofloxacin			
		E=Ethambutol Lfx=Levofloxacii		n	
			rizinamide	Eto=Ethionamide	
				Cs=Cycloserine	
			eptomycin	Cs=Cycloserine PAS=Para-amino	osalicyclic Acid
				Cs=Cycloserine PAS=Para-amino Cm=Capreomyci	
				PAS=Para-amino	
				PAS=Para-amino Cm=Capreomyci Clf=Clofazimine Lzd=Linezoli	
				PAS=Para-amino Cm=Capreomyci Clf=Clofazimine Lzd=Linezoli Trd=Terizidone	n
				PAS=Para-amino Cm=Capreomyci Clf=Clofazimine Lzd=Linezoli Trd=Terizidone Amx/Clv=Amxic	
				PAS=Para-amino Cm=Capreomyci Clf=Clofazimine Lzd=Linezoli Trd=Terizidone	n cillin+Clavulanate



11 0	Regimen and Drug Doses *Date: Date treatment	Date *	Z (mg	Km (gm)	Ofx/Lf x (mg)	Eto (mg	Cs (m		Cm (gm)	PAS (gm)
)			,	,			
	started and doses, Change									
	of doses (if any)	Date	Clf	Amx/Cl	Trd	Lzd	M	fx	Othe	Comment
		*	(mg	v (gm)	(mg)	(mg	(m		r	S
)	, ,	, 0,))			
11	Results of sputum	Month	Wee	k Date of	sample co	llection	1	Res	sult	
l	Examination	0								
		1	1							
			2							
			3							
	[Notation method:	2	4					-		
		2	2							
	No AFB =0		3					-		
			4							
	1-9 AFB per 100 HPF=	3	1							
	Scanty (report number of		2							
	AFB)		3							
			4							
	10-99 AFB per 100 HPF=	4	1							
	+	5								
		6								
	1-10 AFB per HPF= ++	7								
		8								
	>10 AFB=+++]	9								
		10								
		11								
		12								
		13								
		14								
		15								
		16								
		17 18								
		19								
		20								
		21								
		22								
		23								
		24						1		
										



11 2	Adverse Drug Reaction	Date	Adve	erse Drug tion	Sust Drug	pected g	Measure Taken
11 3	Results of Culture	Month		Date of sample collection		Result	
	[Notation Method:	0 1 2					
	No growth reported=0	3					
	Fewer than 10 colonies=report number of colonies (1-9) 10-100 Colonies=+ More than 100 colonies=++ Innumerable or confluent growth=+++ Non-tuberculous mycobacteria= NTM Contaminated=contaminat ed]	5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24					
		24					



11	Drug Susceptibility	Method	Date	S	Н	R	Е	Km	
4	Testing (DST) Results					İ	<u> </u>		
	[Notation:						1		
	[Notation:	Method	Date	Ofx/Lfx	Eto	Other	Other	Other	
	R=Resistant	Method	Date	OIX/LIX	1210	Other	Other	Other	
	S=Susceptible								
	C=Contaminated								
	Unk=Unknown]								
	Method:								
	Xpert MTB/RIF1								
	Line Probe Assay (LPA2								
	Liquid Culture3								
	Solid Culture (L-J)4								
11	In which phase of the	Injectable	phase of	21 months i	regimen	1			
5	treatment are you currently on?	Continuati	on phase	of 21 mont	hs			7	
		regimen	.2						
		Injectable	njectable phase of 9 months regimen3						
		injectuere	phase of	9 months re	egimen	3			
				9 months re					
11	How long are you on this								
11 6	How long are you on this phase of treatments?								
							Months		
11			on phase	of 9 month			Months		
6	phase of treatments?	Continuati	on phase	of 9 month			Months		
11	phase of treatments?	Continuati Positive	on phase	of 9 month12			Months		
11	phase of treatments?	Positive Negative Not Tested	on phase	of 9 month12			Months		



11	Final outcome (If	Cured1			1
8	treatment is completed)	Treatment completed2			
		Died3			
		Treatment failure4			
		Default5			
		Transfer Out6			
11 9	Date of starting treatment		Day		
			Month		
			Year		
12	Date of completion of		Day		-
0	treatment		2 11)		
			Month		
			Year		
					ı

Section 2.0: Previous treatment

PATIENT INTERVIEW SECTION	Start time: Hours _
Minutes	

No.	Questions and filters	Coding categories	Response	Skip
201	Have you ever had TB treatment before? CROSS-CHECK WITH INFORMATION FROM PATENT CARD	Yes		If 2 ▶ 301
202	What is your TB treatment history CROSS-CHECK WITH INFORMATION FROM PATENT CARD	Cat I Treatment failure		If 1,2,5 ▶ 301 If 3 ▶ 204
203	Your treatment card indicated that you had default treatment. Why was previous treatment not completed? [For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	A. Distance to the facility B. Lack of money for treatment costs C. Drug Side Effects D. Moved/Migrated E. Other Please Specify		
204	If on Cat IV treatment, how long have you been on TB treatment before you were diagnosed with MDR-TB?		Months	



Section 3: Delay, Prediagnostic & Diagnostic Costs

No.	Questions and filters	Coding categories	Response	Skip
301	What symptoms did you experience that led you to	a. Cough		
	seek treatment for your most recent illness with TB?	b. Evening rise of temperature/low grade fever		
		c. Coughing up blood		
		d. Weight loss		
	[For each option, record 1 if	e. Other		
	the option is mentioned and record 2 if the option has not been mentioned.]	Specify		
302	How long did you experience these symptoms before you	a. Cough	Weeks	
	went to seek treatment?	b. Evening rise of temperature/low grade fever	Weeks	
	[For each option, record number of weeks if mentioned	c. Coughing up blood	Weeks	
	yes in the previous question.]	d. Weight loss	Weeks	
		e. Other	Weeks	
		Specify]

	tion 3.1 First Visit		<u>, </u>
303	After you experienced the symptoms, which provider did you go to first?	Community Clinic1 Union Subcenter2	
		Upazila Health Complex3	
		District Hospital4	
		Pharmacy & Drug Store5	
		Homoeopath6	
		Private Hospital7	
		Traditional Healer8	
		Other9	
		Specify	
304	What was the distance to the provider from your home?		Kilometer
305	What was the travel time to reach this provider?		Hour
	-		Minute
306	What was the waiting and consultation time with the provider?		Hour
	provider:		Minute
307	What was the registration (ticket) cost paid by you? In	IF not known put 99999	
	TAKA	IF no registration cost 000	
308	What was the consultation fee you have paid? In	IF not known put 99999	
	TAKA	IF no consultation fee 000	
309	What was the cost you paid for diagnostic tests? In	IF not known put 99999	
	TAKA	IF no cost for tests 000	



310	What was the cost for x-ray? In TAKA	IF not known put 99999	
		IF no cost for x-ray 000	
311	What was the costs of	IF not known put 99999	
	drugs? In TAKA	IF no cost for drugs 000	
312	What was travel costs?	IF not known put 99999	
	(INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF no travel cost 000	
313	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
314	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Secti	ion 3.2 Second Visit			
No.	Questions and filters	Coding categories	Response	Skip
315	Once you experience the symptoms to which provider did you go after you have seen the provider type? (MENTION THE FIRST	Community Clinic		
	VISIT'S PROVIDER TYPE)	Pharmacy & Drug Store		
316	What was the distance to the provider from your home?		Kilometer	
317	What was the travel time to reach this provider?		Hour Minute	
318	What was the waiting and consultation time with the provider?		Hour Minute	
319	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
320	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
321	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



3322	What was the cost for x-ray? In TAKA	IF not known put 99999	
		IF no cost for x-ray 000	
323	What was the costs of	IF not known put 99999	
	drugs? In TAKA	IF no cost for drugs 000	
324	What was travel costs?	IF not known put 99999	
	(INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF no travel cost 000	
325	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
326	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	

No.	Questions and filters	Coding categories	Response	Skip
339	Once you experience the symptoms to which provider did you go after you have seen the provider type	Community Clinic		
	(MENTION THE THIRD VISIT'S PROVIDER TYPE)	District Hospital		
340	What was the distance to the provider from your home?		Kilometer	
341	What was the travel time to reach this provider?		Hour Minute	
342	What was the waiting and consultation time with the provider?		Hour Minute	
343	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
344	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
345	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



346	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
347	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
348	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
349	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
350	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	

Sect	Section 3.5 Fifth Visit			
No.	Questions and filters	Coding categories	Response	Skip
351	Once you experience the symptoms to which provider did you go after you have seen the provider type? (MENTION THE FOURTH VISIT'S PROVIDER TYPE)	Community Clinic		
352	What was the distance to the provider from your home?		Kilometer	
353	What was the travel time to reach this provider?		Hour Minute	
354	What was the waiting and consultation time with the provider?		Hour Minute	
355	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
356	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
357	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



358	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
359	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
360	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
361	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
362	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



	on 3.6 Sixth Visit			
No.	Questions and filters	Coding categories	Response	Skip
363	Once you experience the symptoms to which provider did you go after you have seen the provider type	Community Clinic1 Union Subcenter2		
	?	Upazila Health Complex3 District Hospital4		
	(MENTION THE FIFTH VISIT'S PROVIDER	Pharmacy & Drug Store5		
	TYPE)	Homoeopath6		
		Private Hospital7		
		Traditional Healer		
		Specify		
364	What was the distance to the provider from your home?		Kilometer	
365	What was the travel time to reach this provider?		Hour	
			Minute	
366	What was the waiting and consultation time with the provider?		Hour Minute	
367	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
3368	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
369	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



370	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
371	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
372	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
373	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
374	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	Section 3.7 Seventh Visit			
No.	Questions and filters	Coding categories	Response	Skip
375	Once you experience the symptoms to which provider did you go after you have seen the provider type	Community Clinic		
	(MENTION THE SIXTH VISIT'S PROVIDER TYPE)	District Hospital		
376	What was the distance to the provider from your home?		Kilometer	
377	What was the travel time to reach this provider?		Hour Minute	
378	What was the waiting and consultation time with the provider?		Hour Minute	
379	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
380	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
381	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



382	What was the cost for x-ray? In TAKA	IF not known put 99999 IF no cost for x-ray 000	
383	What was the costs of drugs? In TAKA	IF not known put 99999 IF no cost for drugs 000	
384	What was travel costs? (INCLUDE ALL TRAVEL RELATED COSTS: RETURN TRAVEL, TRAVEL FOR LABORATIRY TESTS, DTUGS, COST FOR ACCOMPANYING PERSONS) In TAKA	IF not known put 99999 IF no travel cost 000	
385	What was the food costs? In TAKA	IF not known put 99999 IF no food cost 000	
386	What was cost for accommodation?	IF not known put 99999 IF no accommodation cost 000	



Sect	Section 3.8 Eighth Visit			
No.	Questions and filters	Coding categories	Response	Skip
387	Once you experience the symptoms to which provider did you go after you have seen the provider type? (MENTION THE SEVENH VISIT'S PROVIDER TYPE)	Community Clinic		
388	What was the distance to the provider from your home?	Other9 Specify	Kilometer	
389	What was the travel time to reach this provider?		Hour Minute	
390	What was the waiting and consultation time with the provider?		Hour Minute	
391	What was the registration (ticket) cost paid by you? In TAKA	IF not known put 99999 IF no registration cost 000		
392	What was the consultation fee you have paid? In TAKA	IF not known put 99999 IF no consultation fee 000		
393	What was the cost you paid for diagnostic tests? In TAKA	IF not known put 99999 IF no cost for tests 000		



394	What was the cost for x-ray? In TAKA	IF not known put 99999	
	149 : 111 1111111	IF no cost for x-ray 000	
395	What was the costs of	IF not known put 99999	
	drugs? In TAKA	IF no cost for drugs 000	
396	What was travel costs?	IF not known put 99999	
	(INCLUDE ALL	IF no travel cost 000	
	TRAVEL RELATED		
	COSTS: RETURN		
	TRAVEL, TRAVEL		
	FOR LABORATIRY		
	TESTS, DTUGS, COST		
	FOR		
	ACCOMPANYING		
	PERSONS) In TAKA		
397	What was the food	IF not known put 99999	
	costs? In TAKA	IF f	
		IF no food cost 000	
398	What was cost for	IF not known put 99999	
	accommodation?	IF no accommodation cost 000	



Section 4.0: Treatment Costs

Sect	Section 4.1 Cost Related to DOT			
No.	Questions and filters	Coding categories	Response	Skip
401	From where did you get your TB drugs?	Health facility 1 Home 2 Community 3 Pharmacy 4 Workplace 5 Other 9 Specify		
402	How often do you travel to the health facility / hospital for picking up your TB drugs?		Times/ month	
403	How long does it take to go to the place for picking up the drugs (ONE WAY)?	A. On Foot	Hour Minute	
		B. By Transport	Hour Minute	
404	How long does one of these visits take on average, including time on the road and waiting time (total turnaround time)?		Hour Minute	
405	From your home to the facility, how much does it cost if you take transport? (BOTH WAYS)	IF not known put 99999		



406	If you go to a facility to pick up your drugs, how much do you spend on food on that day? (on the road, while waiting, lunch etc.)	IF not known put 99999		
407	Do you have to pay administration fees when picking up your TB drugs?	Yes		If 2 ▶ 409
408	What was the administration cost?	IF not known put 99999 IF no administration cost 000		
409	Do you have to pay for accommodation when picking up your TB drugs?	Yes1 No2		If 2 ▶ 411
410	What was the accommodation cost?	IF not known put 99999 IF no accommodation cost 000		
Sect	ion 4.2 Cost related t	to Follow Up Tests		
411	Did you ever have to go to the health facility in addition to your regular visits for follow up tests since the beginning of treatment?	Yes		If 2 ▶ 414
412	If yes, how many times?		Times	
413	If yes, did you have to pay any additional costs any time during the entire period?	Yes1 No2		If 2 ▶ 421
414	If so, what kind of costs and how much did you	A. Fees		



	pay last time (In		
	TAKA)?	B. Sputum Test	
	IF not known put 99999	C. X -ray	
	IF no cost put 000	D. TB Drugs	
		E. Other Drugs	
		F. Others	
415	How long does one of these follow-up visits take on average, including time on the		Hour Minute
	road, waiting time and tests (total turnaround time)?		

Section 5.0: Cost related to accompanied persons (Friends/Guardian)

No.	Questions and filters	Coding categories	Response	Skip
501	Does any family/friend/DOT supporter accompany you on any visits before diagnosis and/or during diagnosis?	Yes		If 2 ▶ 507
502	If YES, on how many visits has your family/friend/DOT supporter accompanied you or gone with you before diagnosis and/or during diagnosis?		Times	
503	What was the cost for pre- diagnosis/diagnosis visits of	A. Transport Cost		



	your accompanying person in Taka)?	B. Food Cost		
		C. Accommodation Cost		
	IF not known put 99999			
	IF no cost put 000			
504	Does your accompanying person earn?	Yes1		If 2
		No2		•
				506
505	If earn, how much the person earn per day (in TAKA)?	IF not known put 99999		
506	Why did someone accompany you?	A. Distance		
	accompany you:	B. Security		
	[For each option, record 1	C. Administrative barrier		
	if the option is mentioned and record 2 if the option	D. Too ill to travel alone		
	has not been mentioned.]	E. Was required for treatment		
		F. Other		
		Please Specify		
507	Does any	Yes1		If 2
	family/friend/DOT supporter accompany you	No2		•
	on any visits during treatment (taking drugs)?			601
508	If YES, on how many visits has your family/friend/DOT supporter accompanied you		Times	
	or gone with you during treatment?			
509	What was the cost for visits of your accompanying	A. Transport Cost		
	person during your treatment in TAKA)?	B. Food Cost		



	IF not known put 99999	C. Accommodation Cost	
510	IF no cost put 000 Does your accompanying person earn?	Yes1 No2	If 2 • 512
511	If earn, how much the person earn per day (In TAKA)?		
512	Why did someone accompany you? [For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	A. Distance B. Security C. Administrative barrier D. Too ill to travel alone E. Was required for treatment F. Other Please Specify	

Section 6.0: Hospitalization Costs

No.	Questions and filters	Coding categories	Response	Skip
601	Have you been hospitalized	Yes1		If 2
	before (but due to TB) or	No2		•
	during your TB treatment?			•
				701
602	TOTATION 1			
602	If YES, how many times		Times	
	were you hospitalized for this illness?			
	uns miess:			
603	How much did you pay	A. Hospital Administration Fees		
	during your last stay at the			
	hospital (In TAKA)?	B. Hospital stay charges		



604	IF not known put 99999 IF no cost put 000	C. Food (Not provided by the hospital) D. Transport (Both Ways) E. Drugs F. Diagnostic Tests G. Others		
604	Did any attendant/caregiver stay with you at the hospital?	Yes		If 2 ▶ 701
605	If YES, how many days he/she stay with you (sleep there)?		Days	
606	Were there any extra costs for your relative/friend for staying at the hospital?	Yes		If 2 ▶ 506
607	What was the cost for the accompanying person during your hospitalization (stayed at night)? In TAKA IF not known put 99999 IF no cost put 000	A. Transport Cost B. Food Cost C. Accommodation Cost D. Other Cost		
608	Does your accompanying person earn?	Yes		If 2 /3 ► 610



609	If earn, how much the accompanying person earn per day? IF not known put 99999		
610	Did any other family/friend visit you while in hospital?	Yes	If 2
611	If YES, How many people visited you?		Days
612	On an average, how many times did each of these person visit you?		Times
613	What was the cost for EACH VISIT of them (IN TAKA)?	A. Transport Cost B. Food Cost	
	IF not known put 99999 IF no cost put 000	C. Accommodation Cost D. Other Cost	
614	How long were the visits		Hour
	including traveling time?		Minute

Section 7.0: Other Costs, Other Illnesses and Coping Costs

Sect	tion 7.1 Other Costs			
No.	Questions and filters	Coding categories	Response	Skip
701	Did you buy any supplements for your diet because of the TB illness, for example vitamins, meat, energy drinks, soft drinks, fruits or medicines?	Yes		If 2 ▶ 703
702	If YES, how much did you spend approximately on each of these items each month (in TAKA)?	A. Meat B. Fish C. Fruits		
	IF not known put 99999	D. Drinks E. Vegetables F. Vitamins/Herbs G. Others		
703	Did you have to move to be able to receive (MDR) TB treatment?	Yes		If 2 ▶ 705
704	If YES: how much did you pay for relocation? (In TAKA) IF not known put 99999			
	•			

705	Did you experience any adverse events during the treatment of (MDR-) TB? (Adverse events are any additional health problems that occur during(MDR-) TB treatment and that may be related to the treatment) If YES: Was treatment	Yes	If 2 ▶ 708
	required of these events? This includes changes in TB drug regimen!	No2	
707	IF YES, How much did you spend on treatment of adverse events and/or changes in the TB drug regimen approximately?	A. Drugs B. Fees C. Transport D. Accommodation E. Costs borne by guardian/friends F. Others	
Sect	tion 7.2 Other illnesses		
708	Do you have any chronic illness for which you are receiving treatment?	Yes1 No2	If 2 ▶ 712
709	If YES, which disease do you have? [For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	A. Diabetes B. Heart Disease C. High Blood Pressure D. Cancer E. Arthritis F. Other Specify	



		G. Other	
		Specify	
710	Are there any additional	Yes1	If 2
	costs for you because of this other illness besides	No2	•
	the costs that you have		710
	already mentioned?		712
711	If YES, how much are	A. Drugs	
	these additional costs on		
	average per month? In TAKA	B. Diagnostic tests	
	171101		
		C. Transport	
	IF not known put 99999		
		D. Fees	
		E. Others	
		L. Others	
712	How much did you spend		
	on healthcare on average		
	per month BEFORE the TB illness? In TAKA		
713	How much did you spend on healthcare on average		
	per month AFTER the TB		
	illness? In TAKA		
Sect	ion 7.3 Coping Costs		
714	Has your illness with TB	Yes1	
	resulted in a financial	No2	
	burden?	1102	
715	Did you borrow any	Yes1	If 2
	money to cover costs due		
	to the TB illness?	No2	
			720
716	If YES, how much did you	IF not known put 99999	
	borrow? In TAKA		
		A. Family	



717	From whom did you borrow? [For each option, record 1 if the option is mentioned and record 2 if the option has not been mentioned.]	B. Neighbor C. Friend D. Bank E. Cooperative F. NGO G. Money lender H. Others Specify	
718	Have you already paid back the borrowed money?	Yes	If 2 ▶ 721
719	How are you planning to pay back the money?	In Full	If 2 ▶ 721
720	If you are paying in installment, what is the monthly installment? In TAKA	IF not known put 99999	
721	Have you sold any of your property to finance the cost of the TB illness?	Yes	If 2 ▶ 723
722	How much money did you get from the sale of each of these properties? IF not known put 99999	A. Land B. Livestock C. Transport/Vehicle D. Household item	
		E. Farm produce	



		F. Jewelry G. Savings (FDR) H. Other				
723	Have you employed any household help for your illness?	Specify Yes				If 2
724	How much do you pay monthly to the household help? In TAKA	IF not known put 99999				801

Section 8.0 Patient Income and Patient's Household Income

No.	Questions and filters	Coding categories	Response	Skip
801	Who is the primary income	Patient1		
	earner of the household?	Patient's Wife2		
		Patient's Husband3		
		Patient's Mother4		
		Patient's Father5		
		Patient's Son6		
		Patient's Daughter7		
		Other8		
		Specify		
802	What is the highest level of Education (In years) of these individuals?	A. Patient		
	[For each option, record	B. Primary Income Earner (If Other than patient)		-
	the number of years spent studying. if the person is illiterate/did not go to school record "0"]	C. Household Head (If Other than patient)		_
	sensor record o 1	B. Spouse of Household Head (If Other than patient)		-
803	Are you involved in	Yes, formal work1		If 4
	income earning activities?	Yes, agricultural and other household work2		▶ 809
		Yes, informal work3		
		No4		



804	If No, why are you not involved in any income earning activities?	Cannot work due to illness1 Stopped working after contracting TB		
805	Have you left your job due	Yes1		If 2
	to your TB illness?	No2		•
				809
906	If VEC 1			
806	If YES, how many months ago did you leave your		Months	
	job?			
807	What was your monthly	IF not known put 99999		
	income when you were working? In TAKA			
808	How regularly did you	Throughout the year1		
	work before you became ill with TB?	Seasonal/part of the year2		
		Day Labor3		
		Other4		
		Specify		

809	What was your main occupation before your illness with TB?	Service		
810	Did you have to change jobs when you became ill with TB?	Yes		
811	What is your main occupation after your illness with TB?	Service. 1 Agriculture. 2 Household work. 3 Construction. 4 Garments worker. 5 Transport. 6 Student. 7 Retired. 8 Other. 9 Specify		
812	How many hours did you work on average per day BEFORE you became ill with TB?		Hours	
813	How many hours do you work on average NOW per day?		Hours	



814	If answer to 812 differs from answer to 813:Is the change related to the TB illness?	Yes1 No2	If 2 ▶ 818
815	What was your estimated personal income per month BEFORE the TB illness? In TAKA	IF not known put 99999	
816	What is your estimated personal income per month NOW? In TAKA	IF not known put 99999	
817	Is someone doing the work you used to do after your ilnness?	Spouse 1 Son 2 Daughter 3 Friend 4 Nobody 5 Other 9 Specify 9	
818	Do you have children of or below school age?	Yes1 No2	If 2 ▶ 822
819	Are all of your children attending school regularly?	Yes1 No2	If 1 ▶ 822
820	Did your children go to school regularly before your recent illness with TB?	Yes1 No2	
821	If your children do not go to school, indicate the reasons for not attending school regularly?	A. Needs to help around the house B. No money for school fess C. Also sick D. Has to work to earn	



	[For each option, record 1 if the option is mentioned and record 2 if the option	E. Take care of patient E. Other		
	has not been mentioned.]	Specify		
822	Has the TB illness affected	Yes1		If 2
	your social or private life in any way?	No2		•
	in any way:			922
				823
823	If YES, how was your social life affected?	A. Divorce		
	social file affected?	B. Loss of job		
		C. Dropped out of school		
	[For each option, record 1			
	if the option is mentioned and record 2 if the option	D. Separated from spouse		
	has not been mentioned.]	E. Disruption of sexual life		
		F. Sick child		
		G. Other		
		Specify		
824	What is your religion?	Islam1		
		Hindu2		
		Christian3		
		Buddhism4		
		Other5		
		Specify		
Sect	ion 8.2 Household Inco	me	,	
825	How much do you	A. Patient's income		
	estimate was the average			
	income of your household per month BEFORE the	B. Income of rest of the household		
	TB illness? (for all	C Court assistance		
	persons in the house, including patient; includes	C. Govt. assistance		



	welfare payments, government assistance or other social support)? In TAKA	D. Other	
	IF not known put 99999		
826	How much do you estimate was the average income of your household per month NOW? (for all	A. Patient's income B. Income of rest of the household	
	persons in the house, including patient; includes welfare payments, government assistance or	c. Govt. assistance D. Other	
	other social support)? In TAKA		
	IF not known put 99999		
827	How many people regularly sleep in your house? (including patient)		
828	How many members of the household are employed for wage/salary? (including patient)		
829	Besides yourself, does anyone else of your household receive treatment for TB?	Yes1 No2	If 2 ▶ 823
830	If YES, how many household members are suffering from TB?		



Section 9.0 Socioeconomic Indicators

Secti	Section 9.0 Questions about the household		RESPONSES	Skip
901	What is the main source of lighting for the household?	Electrical mains 01 Solar electricity 02 Generator 03 Gas 04 Kerosine/Oil lamp 05 Candles / Torch 06 Coleman (Kerosene/Pressure lanterns) 07 Improvised lamp 08 Others 96 Specify 96		
902	What is the main source of water for drinking and food preparation for the household?	No source of lighting 98 Piped into household 01 Piped to yard/plot 02 Piped into neighborhood 03 Protected well 04 Unprotected well 05 Protected Spring 06 Unprotected Spring 07 River/stream 08 Pond/lake/dam 10 Communal tank 11 Rainwater 12 Tank truck / Water cart 13 Bottled water 14 Other 96 (Specify) Not reported 99		
903	What type of fuel is mainly used in your household for cooking?	Gas		

		Electricity06		
		25		
		Biogas07		
		Straw/shrubs/grass08		
		Straw/shruos/grass00		
		Saw dust		
		Others96		
		Specify		
904	What kind of	Own flush toilet01		
904		Shared flush toilet		
	toilet facility do	Ventilated improved pit latrine03		
	people in your	Pit latrine with slab04		
	house mainly	Pit latrine without slab / open pit05		
	use?	Bowl/Bucket system06		
		Closet oversea/river07		
		No facility/bush/seashore08		
		Other 96		
		(Specify)		
		Not reported 99		
905	Main	Natural Floor:		
	material of the	Earth		
	floor for the	Sand		
	principal residence	Rudimentary: Wood planks21		
	structure	Palm/bamboo		
	Structure	Finished:		
		Polished wood31		
		Vinyl/asphalt strips32		
		Marble/Ceramic tiles33		
		Floor tile		
		Cement / Concrete/35		
		Brick36		
		Carpet		
		Unpolished		
		Other		
		(Specify)		
		Not reported		
906	Main	Natural roofing		
	material of the	No roof11		
	roof of the	Thatch/palm leaf12		
	principal	Rudimentary roofing		
	residence	Bamboo21		
		Wood planks22		
		Cardboard23		
		Finished roofing		
		Tin / Metal		
		Ceramic Tiles		
Ī	1	Coramic rico	i l	



	Cement / Concrete 34		
	(Specify)		
Main	Natural Walls		
	No walls11		
residence			
	Cardboard26		
	Finished walls		
	Tin / Metal sheets31		
	Cement / Concrete32		
	Brick31		
What type of fuel			If 95
* *	322011101111111111111111111111111111111		11 / 0
•	LPG02		•
•			
use for cooking.	NATURAL GAS		911
	Progra		
	BIOGAS		
	KEROSENE 05		
	KEROSENE		
	COAL, LIGNITE		
	CHARCOAL		
	WOOD		
	STRAW/SHRUBS/GRASS09		
	ACDICULTUDAL CDOD 10		
	AGRICULTURAL CROT		
	ANIMAL DUNG		
	NO FOOD COOKED		
	IN HOUSEHOLD95		
	OTHER OC		
	OTHER 96		
	(SPECIFY)		
		i e	
	Main material of the exterior walls of principal residence What type of fuel does your household mainly use for cooking?	Material of the exterior walls of principal residence	Main material of the exterior walls of principal residence



909	Is the cooking usually done in the house, in a separate building, or outdoors?	IN THE HOUSE	
910	Do you have a separate room which is used as a kitchen?	Yes1 No2	
911	Does your household own any homestead?	Yes1 No2	

	ber of the household own the items?	Yes	No	Respon
Flectricity connection?	A Flectricity	1	2	ses
	·	-		
			2	
-	•		2	
	•	1	2	
		1	2	
An electric fan?		1	2	
An Almirah/wardrobe?		1	2	
A water pump	K. Water Pump	1	2	
An IPS?Generator	L. Generator/IPS	1	2	
An air conditioner		1	2	
A computer/laptop?	N. Computer/Laptop	1	2	
In your opinion, is your	High income		1	
household a high income				
	Middle income		2	
· ·				
nousenoru or a poor nousenoru.				
	income	3		
How many people live in your	Total household size			
household?				
Horrimony adult members (15	Number of edults in the household			
years or older)	Number of adults in the household			
How many children (less than	Number of children in the household			
15 years)				
	An Almirah/wardrobe? A water pump An IPS?Generator An air conditioner A computer/laptop? In your opinion, is your household a high income household, a middle income household or a poor household? How many people live in your household? How many adult members (15 years or older)	Solar Electricity? A radio? C. Radio	Solar Electricity? A radio? C. Radio	Solar Electricity? B. Solar electricity. 1 2 A radio? C. Radio



917	How many separate sleeping rooms are there for the use of your household members in your residence?	Number of rooms	
918	What is your current place of residence?	Urban .1 Urban slum .2 Rural .3 Other .4 Specify	
919	If the government could provide you with some service to ease the burden of TB on you and your household, what would you prefer to have? DO NOT READ. INDICATE THE ITEMS MENTIONED. IF NOT IN THE LIST, ADD ANY NEW GOVT. SERVICE MENTIONED.	Transport vouchers	
920	How much would you be willing to pay for not becoming ill with TB in the first place? In TAKA		



Section 10.0 Performance of the Facility

The next part of the survey is about the quality of TB care that you received during your visits to this facility. Please answer the questions in this part of the survey about this facility only. Do not include any other facilities in your answer.

No.	Questions and filters	Coding categories	Response	Skip
Secti	on 10.1 Availability of T	B Services		
1001	Are the waiting time(s) before being served by health providers of this facility acceptable to you?	Never		
1002	How often are you attended to by the same health providers in this facility?	Never		
1003	How often are the service hours of this facility inconvenient for you to get your TB treatment?	Never		
1004	How often are drugs not available when you require them?	Never		
1005	How often do you experience difficulties in obtaining TB services in this facility because of language barriers?	Never		



1006	How often do you have to go to another health unit for TB services or treatment?	Never	
1007	Is this health facility easy to reach (distance)?	Never	
1008	How often are TB services available during the working hours of this facility?	Never	
1009	How often are the relevant health providers you come to see in this facility available?	Never	
Secti	on 10.2 Communication	and Information	
1010	Do the health providers in this facility tell you when you stop spreading TB to others?	Yes1 No2	
1011	Do the health providers in this facility tell you that TB can be cured?	Yes1 No2	
1012	Do the health providers in this facility tell you about the importance of observed treatment?	Yes1 No2	
1013	Do the health providers in this facility tell you about the side effects of TB drugs?	Yes1 No2	



1014	Do the health providers in this facility tell you about the need for sputum tests at given points during your treatment schedule? Do the health providers in	Yes		
	this facility tell you about the duration of the TB treatment?	No2		
1016	During your visits to this facility, do health providers tell you about how to store your drugs obtained for your treatment?	Yes1 No2		
1017	Does the health provider in this facility tell you when next to come back for TB services?	Yes1 No2		
Coati	on 10 3 Patient – Provide	er interaction and counselling	gr .	
Secu			9	
1018	During your visits to this facility, how often does the health provider treat you with respect?	Never		
	During your visits to this facility, how often does the health provider treat you	Never		



1021	During your visits to this facility, how often do you have sufficient time to discuss your problems? During your visits to this	Never	
1022	facility, how often do health providers discuss with you how to deal with your problems?	Sometimes	
1023	During your visits to this facility, how often do you experience discrimination because you have TB?	Never	
1024	During your visits to this facility, how often is your privacy respected during examination?	Never	
1025	Do health providers at this facility tell you how TB can affect your every day life?	Never	
Secti	on 10.4 Infrastructure		
1026	How often is this facility clean?	Never	



1027	Is there safe drinking water	Never1						
	in this facility?	Sometimes2						
		Usually3						
		Always4						
1028	How often are the toilets in this facility usable?	Never1						
	this facility usable:	Sometimes2						
		Usually3						
		Always4						
1029	How often do you find enough comfortable places	Never1						
	to sit on in this facility?	Sometimes2						
		Usually3						
		Always4						
Section 10.5 Professional Competence								
1030	Does this facility offer	Yes1						
	services to examine your sputum?	No2						
1021	Donathia facilitae affac	Yes1						
1031	Does this facility offer home based TB treatment?							
		No2						
1032	Were you physically	Yes1						
	examined during your first visit to this health facility?	No2						
	visit to this ileater facility.							
1033	Was your sputum examined when you were diagnosed	Yes1						
	with TB?	No2						
1034	How many working days	0-2 Working days1						
	were there between your first sputum submission and	3-4 Working days2						
	the time you got your results?	More than 5 Working days3						



1035	In case of germs in your sputum that cause TB, were your close contacts examined by the TB facility?	Yes1 No2				
1036	How often is there a treatment observer checking on your daily intake of TB drugs?	Never				
		Always4				
Secti	on 10.6 Affordability					
1037	How often do you have to pay for your regular TB services (e.g. sputum tests, TB-drugs, X-rays, etc.)?	Never 1 Sometimes 2 Usually 3 Always 4				
1038	How often do you have to pay a tip in order to receive TB services?	Never				
1039	How often do costs (e.g. transport) prevent you from getting to the health facility?	Never				
Section 10.7 Support						
1040	How often do you receive transport support from the health facility?	Never 1 Sometimes 2 Usually 3 Always 4				



1041	How often do you receive food support from the health	Never1	
	facility?	Sometimes2	
		Usually3	
		Always4	
1042	How often do you receive financial assistance from the	Never1	
	health facility?	Sometimes2	
		Usually3	
		Always4	
Secti	on 10.8 Stigma		
1043	Does the health provider	Never1	
	talk to you the same way you are spoken to when you	Sometimes2	
	receive services other than TB?	Usually3	
		Always4	
1044	Does the health provider	Never1	
	welcome you into the health facility when you visit for	Sometimes2	
	TB services?	Usually3	
		Always4	
1045	Does the health provider turn his/her face away when	Never1	
	speaking with you?	Sometimes2	
		Usually3	
		Always4	
1046	Do you feel that you are	Never1	
	treated with dignity when you visit the health facility?	Sometimes2	
		Usually3	
		Always4	



Section 11.0 Quality of Life

Section 11.1 Health Related Quality of Life (FACIT-TB Questionnaire)

feelir	would you describe your level of ag for the following statements: rd the number in the right column)	Not At All	Slight ly	Moderat ely	Quite A Bit	Extrem ely	Respo nse
F	A. Physical well-being						
110 1	I feel ill	1	2	3	4	5	
110 2	I get tired easily	1	2	3	4	5	
110 3	I have a lack of energy	1	2	3	4	5	
110 4	I have pain	1	2	3	4	5	
110 5	I feel weak all over	1	2	3	4	5	
110 6	I feel fatigued	1	2	3	4	5	
110 7	I have been short of breath	1	2	3	4	5	
110 8	I have nausea-"a sense of vomiting outbreak"	1	2	3	4	5	
110 9	Because of my physical condition, I have trouble meeting the needs of my family	1	2	3	4	5	
111 0	I am bothered by fever (episode of high body temperature)	1	2	3	4	5	
111 1	I am forced to spend time in bed	1	2	3	4	5	
111 2	I have discomfort or pain in my stomach area	1	2	3	4	5	
111 3	I have had itching	1	2	3	4	5	



		1	1	Г		ı	
111 4	I have a loss of appetite	1	2	3	4	5	
111 5	I have been coughing	1	2	3	4	5	
111 6	I am bothered by side effects of treatment	1	2	3	4	5	
111 7	Dusts Worsen my symptoms	1	2	3	4	5	
I	3. Social and economic well-being						
111 8	I feel close to my friends	1	2	3	4	5	
111 9	I get emotional support from my family	1	2	3	4	5	
112 0	I am satisfied with my family communication about my illness	1	2	3	4	5	
112 1	My family has accepted my illness	1	2	3	4	5	
112 2	I feel close to my partner (or the person who is my main support)	1	2	3	4	5	
112 3	I get support from my friends	1	2	3	4	5	
112 4	My physical condition and/or medical treatment cause me financial difficulties	1	2	3	4	5	
	C. Emotional well-being/Stigma of having TB						
112 5	I worry that my condition will get worse	1	2	3	4	5	
112 6	I worry about dying	1	2	3	4	5	
112 7	I am concerned about what the future holds for me	1	2	3	4	5	
112 8	I am embarrassed by my illness	1	2	3	4	5	



			1	1		1	
112 9	It is hard to tell other people about my infection	1	2	3	4	5	
113 0	I am losing hope in the fight against my illness	1	2	3	4	5	
113 1	I am bothered by the change in weight	1	2	3	4	5	
113 2	I worry about spreading my infection	1	2	3	4	5	
113 3	I feel nervous	1	2	3	4	5	
113 4	I feel sad	1	2	3	4	5	
113 5	I am satisfied with how I am coping with my illness	1	2	3	4	5	
I	D. Functional well-being						
113 6	I am content with the quality of my life right now	1	2	3	4	5	
113 7	My work (include work at home) is fulfilling	1	2	3	4	5	
113 8	I am able to work (include work at home)	1	2	3	4	5	
113 9	I am able to enjoy life	1	2	3	4	5	
114 0	I am enjoying the things I usually do for fun	1	2	3	4	5	
114 1	I have accepted my illness	1	2	3	4	5	
114 2	I am sleeping well	1	2	3	4	5	
I	E. Spiritual well-being						
114 3	I find strength in my faith or spiritual belief	1	2	3	4	5	
114 4	My illness has strengthened my faith or spiritual belief	1	2	3	4	5	



114 5	My life is still productive		1	2	3	4	5	
Secti	on 11.2 EQ-5D-5	L Questionnaire						
Unde	er each heading, p	please tick the ONI	E box tha	at best de	escribes you	ır health	TODAY.	
Sl	Category	Options					Respon	nse
114	Mobility	I have no problems about I have slight proble about I have moderate prabout I have severe problems about I am unable to wal about	ems in wareness2 roblems in3 lems in wareness4	alking n walking valking				
114 7	Self-Care	I have no problems myself	ems wash roblems v lems was	ning or dr washing o hing or d	essing or dressing			
114 8	Usual Activities (e.g., work, study, housework, family or leisure activities)	I have no problems activities I have slight proble activities I have moderate practivities3	1 ems doing 2 roblems d	g my usu: loing my	usual			



		activities4				
		I am unable to do my usual				
		activities5				
	D : 7D: 6					
114	Pain/Discomfo	I have no pain or				
9	rt	discomfort1				
		I have slight pain or				
		discomfort2				
		Thoras madagata main an				
		I have moderate pain or discomfort3				
		disconnort				
		I have severe pain or				
		discomfort4				
		I have extreme pain or				
		discomfort5				
115	Anxiety/Depre	I am not				
0	ssion	anxious/depressed1				
		I am slightly				
		anxious/depressed2				
		Lower moderately				
		I am moderately anxious/depressed3				
		unatous/depressed				
		I am severely				
		anxious/depressed4				
		I am extremely				
		anxious/depressed5				
G	11 2 CE CD C					
Secti	on 11.3 SF-6D Q	uesuonnaire				
Under each heading, please tick the ONE box that best describes your health TODAY.						
Sl	Category	Options	Response			
115	Physical	My health does not limit me in vigorous				
1	Functioning	activities1				
		Markarda Parkaran a Ref. 1				
		My health limits me a little in vigorous				
		activities2				
		My health limits me a little in moderate				
		activities3				
l		1				



		My health limits me a lot in moderate activities	
115 2	Role Limitation	I have no problems with your work or other regular daily activities as a result of your physical health or any emotional problems	
3	Social Functioning	My health limits my social activities none of the time	
115 4	Pain	I have no pain	



		work (both outside the home and housework)	
		2	
		I have pain that interferes with your normal work (both outside the home and housework) a little bit	
		I have pain that interferes with your normal work (both outside the home and housework) moderately	
		I have pain that interferes with your normal work (both outside the home and housework) quite a bit	
		I have pain that interferes with your normal work (both outside the home and housework) extremely	
115 5	Mental Health	I feel tense or downhearted and low none of the time1	
		I feel tense or downhearted and low a little of the time2	
		I feel tense or downhearted and low some of the time3	
		I feel tense or downhearted and low most of the time4	
		I feel tense or downhearted and all of the time5	
115	Vitality	I have a lot of energy all of the time1	
6		I have a lot of energy most of the time2	
		I have a lot of energy some of the time3	
		I have a lot of energy a little of the time4	
		1	



Section 11.4 Visual Analog Scale

1157 Visual Analog Scale Score

1158	Comments by interviewer on the interview
1159	INTERVIEWERS: CHECK YOUR FILLED IN QUESTIONNAIRE CAREFULLY
	BEFORE LEAVING THE RESPONDENTS AND END YOUR INTERVIEW BY
	GIVING THANKS TO THE RESPONDENT.
	RECORD THE END TIME OF THE Hour
	INTERVIEW: Minutes

APPENDIX D – PROVIDER QUESTIONNAIRE (DS-TB)

IDENTIFICATION	
DIVISION	
DISTRICT	
UPAZILA	
NAME OF THE FACILITY	
TYPE OF THE FACILITY	
(Union Health Center =01, Upazila Health Complex =02, District Hospital =03)	
TYPE OF THE PROVIDER	
(BRAC =01, Damien Foundation =02, Other =03, Please specify)	
NAME OF THE MANAGER	
LOCATION OF FACILITY: RURAL=1, URBAN=2	



		INTERVIEWER VISITS		
	1	2	3	FINAL VISIT
DATE				
INTERVIEWER'S NAME & CODE			CC	ODE
RESULT CODE*			RE	ESULT CODE
NEXT VISIT: DATE TIME			ТО	OTAL NO. OF VISITS
*RESULT CODES: 01 COMPLETED 02 NOT AVAILABLE 03 POSTPONED 04 REFUSED		05 PARTLY COME 96 OTHER(SPI		
SUPERVISOR	F	FIELD EDITOR	OFFICE EDITOR	KEYED BY
NAME	NAME		NAME	NAME
DATE	DATE		DATE	DATE

Section 1: Number of Personnel, their salary, and contribution to the TB Control Program

Collect data based on each staff working on the day of data collection during day-shift (9:00am- 5:00pm). This table is for collecting information on TB clinics open on day of the interview.

USE DECIMAL POINTS TO INDICATE PART-TIME WORK. FOR EXAMPLE, IF AN INDIVIDUAL WORKS IN TWO CLINICS, RECORD 0.5 FOR THIS PERSON IN BOTH

	A		В	C	D	E	F	G	Н
SL#		Number Employed	Monthly Salary			Overtime	Incentive Payment	Total	% involvement with TB
	DESIGNATION								program
101	Civil Surgeon								
102	Junior Consultant (Chest Clinic)								

	Upazila Health				
103	and Family Planning Officer (UHFPO)				
104	Medical Officer (TB/Leprosy) Designated				
105	Medical Officer Chest Disease Clinic				
106	Medical Officer Disease Control (MODC)				
107	Medical Officer, NGOs				
108	Program Organizer				
109	Medical Technologist (Laboratory)				
110	Health Inspector				
111	Assistant Health Inspector				
112	Family Planning Inspector				
113	Health Assistant (HA)				
114	Medical Assistant (MA)				
115	NGO Community Health Workers				



	Leprosy and TB				
116	Control				
	Assistant			 	
117	Statistical				
	Assistant				
110	Gene Xpert Technician				
	Community				
119	Health Worker				
	(CHW)				
120					
121					
122					
122					
123					
123					
104					
124					
125					
126					
127					
128					
129					



130									
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Section 2.0 Caseload (Number of New Patients Diagnosed, Number of Patients treated)

Please collect these information from Quarterly TB case finding reporting form (TB-10)

SL.	Indicator		Number Ouarter 1 Ouarter 2 Ouarter 3 Ouarter 4												
		(Oct	Quarter1 2015-Dec		(Jan 2	Quarter 2 2016-March		(Apr	Quarter : il 2016-Jur		(July	4 tember			
		Male	Female	Total	Male	Female	Total	Male	Female	Total	Male	Female	Total		
	TB Case Detection Rate	A	В	С	D	Е	F	G	Н	I	J	K	L		
201	Total new pulmonary smear-positive cases notified														
202	Total new (relapses) smear positive cases notified														
203	Total new (failures) smear positive cases notified														
204	Total new (loss to follow up/after default) smear positive cases notified														
205	Xpert MTB/RIF positive RIF sensitive new cases														
206	Xpert MTB/RIF positive RIF sensitive previously treated cases														
207	Total new pulmonary smear-negative cases notified														
208	Total new extrapulmonary cases notified														
209	Total others previously treated* cases notified														
210	Total cases registered														



1	١
	٨
)

		 	T	1	1	1	T		1	1
	Laboratory Activity									
	(Sputum Smear									
	Microscopy)									
211	Number of presumptive									
	TB cases/suspects									
	examined for diagnosis									
	by sputum smear									
	microscopy									
212	Number of presumptive									
	TB cases/suspects with									
	positive sputum smear									
	microscopy									
	Laboratory Activity									
	(GeneXpert Test)									
213	Number of presumptive									
	TB cases/suspects									
	examined for diagnosis									
	by Xpert MTB/RIF									
214	Number of presumptive									
	TB cases/suspects with									
	positive Xpert MTB/RIF									
	result									
	HIV Activities									
214	Number of People living									
	with HIV/AIDS									
	(PLHWA) tested for									
	Acid Fast Bacilli (AFB)									
215	Number of AFB positive									
	result among tested									
	PLWHA									

Section 3.0 TB Patient Referral

Please enumerate the number of TB patients referred by different providers using TB Form-10

Sl.	TB Patient Referral		Num	ber	
		Quarter 1	Quarter 2	Quarter 3	Quarter 4
		A	В	С	D
301	Private Practitioner (Graduate)				
302	Private Practitioner (Non-Graduate)				
303	Govt. field staff				
304	Shasthya Sebika (SS)/ Non-govt. field staff (NGFS)				
305	Village Doctor (VD)				
306	Community Volunteer (CV)				
307	Govt. Hospital				
308	Private Hospital				
309	Community Health Care Provider (CHCP)				
310	TB Patient				
311	Others (Please specify)				
312	Total				



Section 4.0 Treatment outcomes (TB Par	tients Registered 3-6 Months Earlier)
--	---------------------------------------

Quarter |__|_| **Year** |__|_|

PLEASE COMPLETE THIS TABLE FOR PULMONARY TB PATIENTS REGISTERED 3-6 MONTHS EARLIER (USING TB-12 FORM)

Sl.	Type of Patients	Smo	ear	Sm	ear	Die	-d	Fail	iire	Lost	t to	Transf	erred	No	nf	G	rand	Total
	Type of Tutients	Nega			Positive		-	1 4411	ui c	Follow-up		Out		Evaluated			- 4114	10001
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	Н	Ι	J	K	L	M	N	0	P	Q
401	Smear Positive																	
402	Xpert MTB/RIF																	
	Positive																	
403	Smear Negative																	

Sl.	Type of Patients	Sm Nega	ear ative		ear itive	Die	ed	Fail	ure		t to w-up	Transf Oı				Grand Total		
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	Retreatment	A	В	C	D	\mathbf{E}	F	G	H	I	J	K	L	M	N	О	P	Q
404	Relapses																	
405	Treatment after																	
	failure																	
406	Treatment after																	
	loss to follow																	
	up/Default																	
407	Others																	
408	Total																	

Quarter		Year	_
---------	--	------	---

PLEASE COMPLETE THIS TABLE FOR PULMONARY TB PATIENTS REGISTERED 3-6 MONTHS EARLIER (USING TB-12 FORM)

Sl.	Type of Patients	Smo	ear	Sm	ear	Die	ed	Fail	ure	Lost	t to	Transf	erred	No	ot	G	rand	Total
		Nega	tive	Posi	tive					Follov	w-up	Oı	ıt	Evalu	ated			
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	Н	Ι	J	K	L	M	N	0	P	Q
409	Smear Positive																	
410	Xpert MTB/RIF																	
	Positive																	
411	Smear Negative																	

Sl.	Type of Patients	Sme Nega			ear itive	Die	ed	Fail	ure	Los Follo		Transi		No Evalu		G	rand	Total
	M= Male F=Female	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	Retreatment	A	В	С	D	Е	F	G	Н	I	J	K	L	M	N	0	P	Q
412	Relapses																	
413	Treatment after failure																	
414	Treatment after loss to follow up/Default																	
415	Others																	
416	Total																	



Ouarter		Year		
Z mar tor	 	1001	 	

PLEASE COMPLETE THIS TABLE FOR PULMONARY TB PATIENTS REGISTERED 3-6 MONTHS EARLIER (USING TB-12 FORM)

CI	T	C		G		D:	1	T7 - 21.		T		T	P 1	N.T.	- 4	•		T-4-1
Sl.	Type of Patients	Smo		Sm	ear	Die	ea	Fail	ure	Lost	το	Transf	errea	No	-	G	rana	Total
		Nega	ntive	Posi	tive					Follov	w-up	Oı	ıt	Evalu	ıated			
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	\mathbf{F}	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	H	Ι	J	K	L	M	N	0	P	Q
417	Smear Positive																	
418	Xpert MTB/RIF																	
	Positive																	
419	Smear Negative							•									•	

Sl.	Type of Patients	Sme Nega			ear itive	Die	ed	Fail	ure	Los Follo		Transi		No Evalu		G	rand	Total
	M= Male F=Female	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	Retreatment	A	В	C	D	Е	F	G	Н	I	J	K	L	M	N	0	P	Q
420	Relapses																	_
421	Treatment after failure																	
422	Treatment after loss to follow up/Default																	
423	Others																	
424	Total																	



Ouarter		Year	

PLEASE COMPLETE THIS TABLE FOR PULMONARY TB PATIENTS REGISTERED 3-6 MONTHS EARLIER (USING TB-12 FORM)

Sl.	Type of Patients	Smo		Sm		Die	ed	Fail	ure	Lost		Transf		No		G	rand	Total
		Nega	itive	Posi	tive					Follor	w-up	Oı	ıt	Evalu	ıated			
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	H	Ι	J	K	L	M	N	О	P	Q
425	Smear Positive																	
426	Xpert MTB/RIF																	
	Positive																	
427	Smear Negative																	

Sl.	Type of Patients	Sm Nega			ear itive	Die	ed	Fail	ure		t to w-up	Transf		No Evalu		G	rand	Total
	M= Male F=Female	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	Retreatment	A	В	C	D	Е	F	G	Н	I	J	K	L	M	N	0	P	Q
428	Relapses																	
429	Treatment after failure																	
430	Treatment after loss to follow up/Default																	
431	Others																	
432	Total																	



Section 5.0	Treatment	outcomes ((TB Patients	Registered	12-15 Mo	nths Earlier)
Quarter	Year _	_ _				

PLEASE COMPLETE THIS TABLE FOR PULMONRY TB PATIENTS REGISTERED 12-15 MONTHS EARLIER (USING TB-11 FORM)

Sl.	Type of Patients	Cui	red		ment pleted	Die	ed	Fail	ure	Los Follo up/Def	ow-	Transi O		No Evalu		G	rand	Total
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	H	I	J	K	L	M	N	О	P	Q
501	Smear Positive																	
502	Xpert MTB/RIF Positive																	
503	Smear Negative																	
504	EP																	
506	Total															·	_	

Sl.	Type of Patients	Sme Nega			ear itive	Die	ed	Fail	ure	Los Follo		Transf Oı		No Evalu		G	rand	Total
	M= Male F=Female	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	Retreatment	A	В	С	D	E	F	G	Н	I	J	K	L	M	N	О	P	Q
507	Relapses																	
508	Failures																	
509	Treatment after loss to follow up/default																	
510	Others																	
511	Total																	

Quarter	_	Year	
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PLEASE COMPLETE THIS TABLE FOR PULMONRY TB PATIENTS REGISTERED 12-15 MONTHS EARLIER (USING TB-11 FORM)

Sl.	Type of Patients	Cured		Treatment Completed				Failure		Lost to Follow- up/Defaulted		Transferred Out		Not Evaluated		Grand Total		
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	H	I	J	K	L	M	N	0	P	Q
501	Smear Positive																	
502	Xpert MTB/RIF																	
	Positive																	
503	Smear Negative																	
504	EP																	·
506	Total																	

Sl.	Type of Patients	Smear Negative		Smear Positive		Died		Failure		Lost to Follow-up		Transferred Out		Not Evaluated		Grand Total		
	M= Male F=Female	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	Retreatment	A	В	С	D	E	F	G	Н	I	J	K	L	M	N	О	P	Q
507	Relapses																	
508	Failures																	
509	Treatment after loss to follow up/default																	
510	Others																	
511	Total																	



Ouarter	Y	ear		ĺ

PLEASE COMPLETE THIS TABLE FOR PULMONRY TB PATIENTS REGISTERED 12-15 MONTHS EARLIER (USING TB-11 FORM)

Sl.	Type of Patients	Cured		Treatment Completed				Failure		Lost to Follow- up/Defaulted		Transferred Out		Not Evaluated		Grand Total		
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	H	Ι	J	K	L	M	N	0	P	Q
501	Smear Positive																	
502	Xpert MTB/RIF																	
	Positive																	
503	Smear Negative																	
504	EP																	
506	Total																	

Sl.	Type of Patients	Smo Nega		Smear Positive		Died		Failure		Lost to Follow-up		Transferred Out		Not Evaluated		Grand Total		
	M= Male F=Female	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	Retreatment	A	В	С	D	E	F	G	Н	I	J	K	L	M	N	0	P	Q
507	Relapses																	
508	Failures																	
509	Treatment after loss to follow up/default																	
510	Others																	
511	Total																	

Quarter	 _	Year	_
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PLEASE COMPLETE THIS TABLE FOR PULMONRY TB PATIENTS REGISTERED 12-15 MONTHS EARLIER (USING TB-11 FORM)

Sl.	Type of Patients	Cui	red		ment pleted	Die	ed	Fail	ure	Los Follo up/Def	ow-	Transf Oı		No Evalu		G	rand	Total
	M= Male	M	F	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	New Cases	A	В	C	D	E	F	G	Н	I	J	K	L	M	N	0	P	Q
501	Smear Positive																	
502	Xpert MTB/RIF																	
	Positive																	
503	Smear Negative																	
504	EP																	·
506	Total																	

PLEASE COMPLETE THIS TABLE FOR PULMONARY TB PATIENTS REGISTERED 12-15 MONTHS EARLIER (USING TB-11 FORM)

Sl.	Type of Patients	Smo Nega		Sm	ear itive	Die	ed	Fail	ure	Los Follo		Transf O		No Evalu	ot	G	rand	Total
		Nega	itive	1 051	uve					T OHO	w-up	U	ıι	Evan	iaieu			
	M= Male	M	\mathbf{F}	M	F	M	F	M	F	M	F	M	F	M	F	M	F	Total
	F=Female																	
	Retreatment	A	В	C	D	E	F	G	H	I	J	K	L	M	N	О	P	Q
507	Relapses																	
508	Failures																	
509	Treatment after																	
	loss to follow																	
	up/default																	
510	Others																	
511	Total																	



Section 6.0 Quantity and Cost of Drugs

Number of TB Drugs received by the facility last year (January 2016- December 2016-PLEASE CONSULT LAST TWO REQUISITION FORM TB-08)

Ouarter	- 1	Year		١
Qual tel	 	I Cui	 	

SL#	Drug name	Quantity measures	Quantity received from National TB Control Program (NTP) (units) CAT I Patients	Quantity received from National TB Control Program (NTP) (units) CAT II Patients	Total received in this quarter	Market price of drugs per unit
		A	В	C	D	E
601	4FDC					
602	3FDC (R150/H75/E275)					
603	2 FDC (R150/H75)					
604	3 FDC (R60/H30/Z150) (Dispersible)					
605	2FDC (R60/H60) (Dispersible)					
606	2FDC (R60/H30) (Dispersible)					
607	Z 400 mg (Dispersible)					
608	H 100 mg (Dispersible)/For IPT					
609	R 150 mg					
610	H 300 mg					
611	R 450 mg					
612	Z 500 mg					
613	E 400 mg					

ſ	\
-	-
ť	

614	E 100 mg												•	
615	S 1g												•	
616	Inj. Water, 5 ml												•	
617	D/ Syringe 5 cc												•	
618	DST Liquid culture												•	
619	DST Solid Culture												•	
620													•	
621													•	
622													•	
623													•	
624													•	
625													•	

SL#	Drug name	Quantity received from National TB Control Program (NTP) (units) for CAT I Patients	Quantity received from National TB Control Program (NTP) (units) for CAT II Patients	Total received in this quarter
		${f E}$	\mathbf{F}	G
601	4FDC			
602	3FDC (R150/H75/E275)			
603	2 FDC (R150/H75)			
604	3 FDC (R60/H30/Z150) (Dispersible)			

Quarter |__|_| Year |__|_|

	AED C (D (0/H(0))	1					1	1	П	Т	- 1		
605	2FDC (R60/H60) (Dispersible)								L				
606	2FDC (R60/H30)	T				Т			ÌΓ				
606	(Dispersible)				L			<u> </u>	L				
607	Z 400 mg (Dispersible)												
608	H 100 mg (Dispersible)/For IPT												
609	R 150 mg												
610	H 300 mg												
611	R 450 mg												
612	Z 500 mg												
613	E 400 mg												
614	E 100 mg												
615	S 1g												
616	Inj. Water, 5 ml												
617	D/ Syringe 5 cc												
618	DST Liquid culture												
619	DST Solid Culture												
620													
621													
622													
623													
624													
625													
		1 1			_		 						

Quarter |__|_| **Year** |__|_|

SL#	Drug name	fro Co (N	om ont VTI	Na rol P) (y re atio Pr uni Pa	nal ogr ts) i	am for	Na Pro	atio gra	nal ım (TB	Cor P) (1	from ntrol units)	,	 		ived rtei	
					H						Ι					J		
601	4FDC																	
602	3FDC (R150/H75/E275)																	
603	2 FDC (R150/H75)																	
604	3 FDC (R60/H30/Z150) (Dispersible)																	
605	2FDC (R60/H60) (Dispersible)													L				
606	2FDC (R60/H30) (Dispersible)																	
607	Z 400 mg (Dispersible)																	
608	H 100 mg (Dispersible)/For IPT													Ĺ				
609	R 150 mg																	
610	H 300 mg																	
611	R 450 mg																	
612	Z 500 mg																	
613	E 400 mg																	
614	E 100 mg																	
615	S 1g																	
616	Inj. Water, 5 ml																	-

617	D/ Syringe 5 cc										
618	DST Liquid culture										
619	DST Solid Culture										
620											
621											
622											
623											
624											
625											

Quarter |__|_| Year |__|_|

SL#	Drug name	fi (ron Con (NT	n Na tro P)	ty ro atio l Pr (uni l Pa	nal ogr its) i	TB am for	N	atio ogr	onal am (TB (NT) for	Coi P) (l from ntrol units) nts	 tal 1		 d in r	_
					K						L				M		
601	4FDC																
602	3FDC (R150/H75/E275)																
603	2 FDC (R150/H75)																
604	3 FDC (R60/H30/Z150) (Dispersible)																
605	2FDC (R60/H60) (Dispersible)																
606	2FDC (R60/H30) (Dispersible)																



		 						1			
607	Z 400 mg (Dispersible)										
608	H 100 mg (Dispersible)/For IPT				L						
609	R 150 mg										
610	H 300 mg										
611	R 450 mg										
612	Z 500 mg										
613	E 400 mg										
614	E 100 mg										
615	S 1g										
616	Inj. Water, 5 ml										
617	D/ Syringe 5 cc										
618	DST Liquid culture										
619	DST Solid Culture										
620											
621											
622											
623											
624											
625											

Section 7.0 Quantity and Cost of Laboratory Reagents/Supplies/Equipment

Number of Laboratory reagents/ supplies received by the facility last year (October 2015-September 2016- Laboratory Request Form)

Quarter 4 (July 2016- September 2016)

SL#	Reagent/Lab supplies' name	Quantity measures per patient	Quantity received from National TB Control Program (NTP) (units)	Quantity purchased from market (units)	Total received in this quarter	Market price of reagents per unit
		A	В	C	D	E
701	Carbol fuchsin (1%) Solution	3.0 ml /sm+				
702	Methylene Blue (0.1%)	3.0 ml/sm+				
703	Sulphuric Acid (25%)	6.0 ml/sm+				•
704	Burning Spirit	1.5 ml/sm+				
705	Slides	1 pc/sm+				
706	Sputum Containers	1 pc/sm+				
707	Immersion Oil	0.05 ml/sm+				
711	Filter Paper Pieces	100 pc/ clinic				
712	Request Form (TB 05)	1 pc/person				
713	Lab Register (TB 04)					
714	Diamond Pencil					
715	Slide Box					
716						
717						
718						
719						



720												•	
721												•	
722												•	

Quarter 3 (April 2016- June 2016)

SL#	Reagent/Lab supplies' name	Quantity received from National TB Control Program (NTP) (units)	Quantity purchased from market (units)	Total received in this quarter
		F	G	H
701	Carbol fuchsin (1%) Solution			
702	Methylene Blue (0.1%)			
703	Sulphuric Acid (25%)			
704	Burning Spirit			
705	Slides			
706	Sputum Containers			
707	Immersion Oil			
708	Filter Paper Pieces			
709	Request Form (TB 05)			
710	Lab Register (TB 04)			
711	Diamond Pencil			
712	Slide Box			
713	Filter Paper Pieces			
714				
715				

716									
717									
718									
719									
720									
721									
722									

Quarter 2 (January 2016- March 2016)

SL#	Reagent/Lab supplies' name	Quantity received from National TB Control Program (NTP) (units)	Quantity purchased from market (units)	Total received in this quarter
		I	J	K
701	Carbol fuchsin (1%) Solution			
702	Methylene Blue (0.1%)			
703	Sulphuric Acid (25%)			
704	Burning Spirit			
705	Slides			
706	Sputum Containers			
707	Immersion Oil			
708	Filter Paper Pieces			
709	Request Form (TB 05)			
710	Lab Register (TB 04)			



711	Diamond Pencil										
712	Slide Box										
713	Filter Paper Pieces										
714											
715											
716											
717											
718											
719											
720											
721											
722											

Quarter 1 (October 2016- December 2016)

SL#	Reagent/Lab supplies' name	from Nat Control	received tional TB Program (units)	_	uantity om mai	_	Total received in this quarter
		I				M	N
701	Carbol fuchsin (1%) Solution						
702	Methylene Blue (0.1%)						
703	Sulphuric Acid (25%)						
704	Burning Spirit						
705	Slides						

706	Sputum Containers	П							
707	Immersion Oil								
708	Filter Paper Pieces								
709	Request Form (TB 05)								
710	Lab Register (TB 04)								
711	Diamond Pencil								
712	Slide Box								
713	Filter Paper Pieces								
714									
715									
716									
717									
718									
719									
720									
721									
722									



Section 8.0 Cost of Other Supplies

Number of Supplies received by the facility last year (October 2015-September 2016)

Quarter 4 (July 2016- September 2016)

SL#	Supplies' name	Quantity measures	fì	rom Cont	Na rol	rec tion: Prog (un	al T grai	B		mai	rket	nsed nits)	otal i this			M	arke	per	ce of r unit	olies
		A				В					C			D					E	
801	TB Register																		•	
802	Treatment Card																			
803	Pen																			
804	Paper																		•	
805	Box																		•	
806	Soap																		•	
807	Towel																		•	
808	Boxes																		•	
809	Tape																		•	
810	Raincoat																		•	
811	Torch light																		•	
812	Umbrella																		•	
813	Drug Baskets																		•	
814	Kit Bag																			
815	Poster																			
816	Sticker																			
817	Leaflet																			
818	Flip chart																			



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819	Flash Chart													•	
820	Brochure													•	
821														•	
822														•	
823														•	
824														•	
825														•	
826														•	
827														•	
828														•	
829														•	
830														•	

Ouarter 3 (April 2016- June 2016)

SL#	Supplies' name	fro Co	om Na ontrol	y recei ational l Progr) (unit	TB am				hased units)	T	otal i		
				F			(T T				Н	
801	TB Register												
802	Treatment Card												
803	Pen												
804	Paper												
805	Box												
806	Soap												
807	Towel												



808	Boxes												
		┢				H				Н			\dashv
809	Tape									Ц			_
810	Raincoat												
811	Torch light												
812	Umbrella												
813	Drug Baskets												
814	Kit Bag												
815	Poster												
816	Sticker												
817	Leaflet												
818	Flip chart												
819	Flash Chart												
820	Brochure												
821													
822													
823													
824													
825													
826													
827													
828													
829													
830													



Ouarter 2 (January 2016- March 2016)

	2 (January 2010- Mar	Quantity received		
G T //		from National TB	Quantity purchased	Total received in
SL#	Supplies' name	Control Program	from market (units)	this quarter
		(NTP) (units)		_
		I	J	K
801	TB Register			
802	Treatment Card			
803	Pen			
804	Paper			
805	Box			
806	Soap			
807	Towel			
808	Boxes			
809	Tape			
810	Raincoat			
811	Torch light			
812	Umbrella			
813	Drug Baskets			
814	Kit Bag			
815	Poster			
816	Sticker			
817	Leaflet			
818	Flip chart			
819	Flash Chart			
820	Brochure			



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821									
822									
823									
824									
825									
826									
827									
828									
829									
830									

Quarter 4 (October 2015- December 2015)

SL#	Supplies' name	f	Quar rom Cont (N'	Na trol	tion	al I gra	ΓB im		mar		sed nits)		rece qua	
					L			 1		M			N	
801	TB Register													
802	Treatment Card													
803	Pen													
804	Paper													
805	Box													
806	Soap													
807	Towel													
808	Boxes													
809	Tape													



810	Raincoat											
811		+							H			
-	Torch light	+							L			
812	Umbrella											
813	Drug Baskets											
814	Kit Bag											
815	Poster											
816	Sticker											
817	Leaflet											
818	Flip chart											
819	Flash Chart											
820	Brochure											
821												
822												
823												
824												
825												
826												
827												
828												
829												
830												

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Section 9.0 Other Capital Items

Number of Items used by the facility

SL#	Capital Items	Quantity (Unit)	Co	ece Nat ontr	iona ol P	tity from al TB rogram units)	p n	urc	hase	ntity ed from (units)	7	Γota	al is		ing	Life In '			M	arke	et pr	ice o	f iter	n per unit
		A			В				C	,			Ι)			E					ŀ	1	
901	Microscope																							
902	GeneXpert																							
903	Weight Scale																							
904	Signboard																							
905	Computer																							
906	Printer																							
907	Smart Phone																							
908	App development																							
909	Motor Cycle																							
910	Bicycle																							
911	Car																							
912	Ambulance																							
913																								
914																								
915																								
916																								
917																								
918																								
919																								



Section 10. Other Costs

Section 10.1 Cost of Supervision (facility supervisory visits conducted in last year (October 2015-September 2016)

SL#	Title of person conducting supervisory visit	Number of visits in last year (Oct 2015- Sep 2016)	Duration of visit (total for all visits in days)	Per Diem (total)	Travel expenses (total)	Other expenses	What % of costs paid by facility?
	A	В	C	D	E	F	Н
1001							
1002							
1003							
1004							

Section 10.2 Travel cost for Drug /Attending Meeting/Training

SL#	Title of person conducting travel	Number of travel in last year (Oct 2015-Sep 2016)	Purpose of the travel (Use Code from below)^	Duration of visit (total for all visits in days)	Per Diem (total)	Travel expenses (total)	Other expenses	What % of costs paid by facility?
	A	В	C	D	E	F	H	I
1005								
1006								
1007								
1008								

[^]Code: 1. Receiving drug 2. Attending Meetings 3. Training 4. Others (Please specify)



Section 10.3 Cost of Other Activities

No	Questions and Filters	Coding categories	Record Response	Skip
1009	Was Health education on TB organized by the DOTS center?	Yes1 No2		If 2 →1011
1010	If Health education on TB was organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
1011	Was DOTS committee meeting organized by the DOTS center?	Yes1 No2		If 2 →1013
1012	If DOTS committee meeting was organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
1013	Were Contact Tracing activities organized by the DOTS center?	Yes1 No2		If 2 →1015
1014	If Contact Tracing activities were organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka Taka	
1015	Were Refresher training activities organized by the DOTS center?	Yes		If 2 →1017
1016	If Refresher training activities were organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
1017	Were Other meetings organized by the DOTS center?	Yes		If 2 →1019
1018	If Other meetings were organized, How much money was paid for this activity in last year (October 2015- September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
1019	Were Incentive payments were provided to the providers by the DOTS center?	Yes1 No2		If 2 →1021



1020	If incentive payments were given to the providers, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS PAYMENT WAS NOT MADE AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
1021	Were Incentive payments were provided to the patients by the DOTS center?	Yes1 No2		If 2 →1023
1022	If incentive payments were given to the patients, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS PAYMENT WAS NOT MADE AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
1023	Were Incentive payments were provided to the community members by the DOTS center?	Yes1 No2		If 2 →END
1024	If incentive payments were given to the community members, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS PAYMENT WAS NOT MADE AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka Taka	

Thank you for your cooperation! Is there anything you would like to ask or say?

Comments by Interviewer:

Date, Signature by Interviewer:

APPENDIX E – PROVIDER QUESTIONNAIRE (MDR-TB)

IDENTIFICATION	
DIVISION	
DISTRICT	
UPAZILA	
NAME OF THE FACILITY	
TYPE OF THE FACILITY	
(NIDCH =01, District Chest Hospital =02, Damien Foundation Hospital =03, Other=04, Please Specify)	
DRUG REGIMEN FOLLOWED	
(21 Month Regimen=1, 9 Month Regimen=02)	
NAME OF THE MANAGER	
LOCATION OF FACILITY: RURAL=1, URBAN=2	



INTERVIEWER VISITS											
	1		2		3	FINAL VISIT					
DATE											
INTERVIEWER'S NAME & CODE						CODE					
RESULT CODE*					RESULT				CODE		
NEXT VISIT: DATE		TOTAL NO. OF				F VISITS					
TIME						101711	2110. 0	1 VISITS			
*RESULT CODES: 01 COMPLETED 02 NOT AVAILABLE 03 POSTPONED			04 REFUSED 05 PARTLY COMP 96 OTHER(
SUPERVISOR			FIELD EDITOR		OFFICE EDITO	R		KE	YED BY	Y	
NAME		NAME		NAME_			N.	AME			
DATE		DATE		DATE_			D	ATE			

Section 1: Number of Personnel, their salary, and contribution to the MDR-TB Control Program

Collect data based on each staff working on the day of data collection during day-shift (9:00am- 5:00pm). This table is for collecting information on TB clinics open on day of the interview.

USE DECIMAL POINTS TO INDICATE PART-TIME WORK. FOR EXAMPLE, IF AN INDIVIDUAL WORKS IN TWO CLINICS, RECORD 0.5 FOR THIS PERSON IN BOTH

	A		В	С	D	E	F	G	Н
SL#		Number Employed	Monthly Salary	Monthly Benefits	TA/DA	Overtime	Incentive Payment	Total	% involvement with TB
	DESIGNATION								program
101	Professor								
102	Associate Professor								



		1	1	1	T	1	1	1	ı
103	Assistant Professor								
104	Consultant								
105	Registrar								
106	Assistant Registrar								
107	Medical Officer								
108	НМО								
109	Staff Nurse								
110	Program Organizer								
111	Medical Technologist (Laboratory)								
112	Health Inspector								
113	Assistant Health Inspector								
114	Family Planning Inspector								
115	Health Assistant (HA)								
116	Medical Assistant (MA)								
117	NGO Community Health Workers								
118	Leprosy and TB Conrol Assistant								
119	Statistical Assistant								
120	Gene Xpert Technician								



121	Community Health Worker (CHW)				
122					
123					
124					
125					
126					
127					
128					
129					
130					

Section 2.0 Caseload (Number of New Patients Diagnosed, Number of Patients treated) last year (October 2015-September 2016)

PLEASE COMPLETE THIS TABLE FROM QUARTERLY REPORT ON DR TB CASE REGISTRATION (FORM DR TB 08)

Sl.			Quarter l			Quarter I			Quarter II			Quarter IV		
	M=Male	M	F	Total	M	F	Total	M	F	Total	M	F	Total	
	F=Female													
		A	В	C	D	E	F	G	H	I	J	K	L	
201	MDR													
202	XDR													
203	RR													
204	Other DR													
205	Total (Confirmed													
	DR TB)													
206	Presumptive DR													
	TB													
207	Grand Total													

PLEASE COMPLETE THIS TABLE FROM QUARTERLY REPORT ON DR TB CASE REGISTRATION (FORM DR TB 08)

Sl.	Patient registered in DR TB Register	Quarter I	Quarter II	Quarter III	Quarter IV
	Confirmed DR TB	A	В	С	D
208	New				
209	Failure after CAT I				
210	Failure after CAT II				
211	Relapse after CAT II				
212	Relapse after CAT I				
213	Treatment after lost to follow up CAT II				
214	Treatment after lost to follow up CAT I				
215	Delayed Converters CAT II				
216	Delayed Converters CAT I				
217	Close Contact of DR TB with S/S				



218	Total		
	Pulmonary Diagnosed		
219	New		
220	Previously treated		
221	Unknown TB treatment history		
	Extrapulmonary		
222	New		
223	Previously treated		
224	Unknown TB treatment history		
225	Total		
	Grand Total		

Section 3.0 Xpert MTB/RIF Result

Please enumerate the number of MDR-TB patients detected last year (October 2015-September 2016) USINF DR TB FORM 10A

Sl. MDR-TB Patient Referral Month													
		1	2	3	4	5	6	7	8	9	10	11	12
		A	В	C	D	Е	F	G	Н	I	J	K	L
301	Number of total presumptive DR VTB cases tested												
302	Number of MTB detected Rif resistance not detected (T)												
303	Number of MTB detected Rif resistance detected (RR)												
304	Number of MTB detected Rif resistance indeterminate (TI)												
305	Number of MTB not detected (N)												
306	Number of invalid/no result/ error (I)												

Section 4.0 Treatment outcomes (MDR-TB Patients got treatment 24 to 36 months earlier)

PLEASE COMPLETE THIS TABLE FOR MDR TB PATIENTS GOT TREATMENT 24 TO 36 MONTHS EARLIER USING DR TB FORM 09

Quarter |__|_| Year |__|_|

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed					up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	I	J	K
401	MDR											
402	XDR											
403	RR											
404	Other											
405	Presumptive											
	DR TB											
406	Total											

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed		_			up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
		_	quarter									
		A	В	С	D	E	F	G	H	I	J	K



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407	New						
408	Failure after CAT						
	Ι						
409	Failure after CAT						
	II						
410	Relapse after						
	CAT II						
411	Relapse after						
	CAT I						
412	Treatment after						
	lost to follow up						
	CAT II						
413	Treatment after						
	lost to follow up						
41.4	CAT I						
414	Delayed						
	Converters CAT II						
415	Delayed						
415	Converters CAT						
	I						
416	Close Contact of						
110	DR TB with S/S						
	Pulmonary						
	Diagnosed						
417	New						
418	Previously						
	treated						
419	Unknown TB	 				 	
	treatment history						
	Extrapulmonary						
420	New						
421	Previously						
	treated						
422	Unknown TB						
	treatment history						



PLEASE COMPLETE THIS TABLE FOR MDR TB PATIENTS GOT TREATMENT 24 TO 36 MONTHS EARLIER USING DR TB FORM 09

Quarter |__|_| Year |__|_|

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed					up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	I	J	K
401	MDR											
402	XDR											
403	RR											
404	Other											
405	Presumptive											
	DR TB											
406	Total											

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed					up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	Ι	J	K
407	New											



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100		1			1		1
408	Failure after CAT						
	I			 		 	
409	Failure after CAT		 	 		 	
	II						
410	Relapse after						
	CAT II						
411	Relapse after						
	CATI						
412	Treatment after						
	lost to follow up						
	CAT II						
413	Treatment after						
	lost to follow up						
	CAT I						
414	Delayed						
	Converters CAT						
	II						
415							
	Converters CAT						
	I						
416							
	DR TB with S/S						
	Pulmonary						
	Diagnosed						
417	New						
418	Previously						
	treated						
419	Unknown TB						
	treatment history						
	Extrapulmonary						
420	New						
421	Previously						
	treated						
422	Unknown TB						
	treatment history						
	a cament moory						



PLEASE COMPLETE THIS TABLE FOR MDR TB PATIENTS GOT TREATMENT 24 TO 36 MONTHS EARLIER USING DR TB FORM 09

Quarter |__|_| Year |__|_|

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed					up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	I	J	K
401	MDR											
402	XDR											
403	RR											
404	Other											
405	Presumptive											
	DR TB											
406	Total											

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed					up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	Ι	J	K
407	New											



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	T		_	ı				1
408	Failure after CAT							
	I							
409	Failure after CAT							
	II							
410	Relapse after							
	CAT II							
411	Relapse after							
	CAT I							
412	Treatment after							
	lost to follow up							
	CAT II							
413	Treatment after							
	lost to follow up							
	CAT I							
414	Delayed							
	Converters CAT							
	II							
415	Delayed							
	Converters CAT							
	I							
416	Close Contact of							
	DR TB with S/S							
	Pulmonary							
	Diagnosed							
417	New							
418	Previously							
	treated							
419	Unknown TB							
	treatment history							
	Extrapulmonary							
420	New							
421	Previously							
	treated							
422	Unknown TB							
	treatment history							
<u> </u>	a cathlent mistory		I					

PLEASE COMPLETE THIS TABLE FOR MDR TB PATIENTS GOT TREATMENT 24 TO 36 MONTHS EARLIER USING DR TB FORM 09

Quarter |__|_| Year |__|_|

Sl.	Patient Group	Total number of DR TB	Number of confirmed	Cured	Treatment Completed	Failure	Died	Lost to follow up	Transferred Out	Still on Treatment	Not Evaluated	Total
		patients registered	DR TB patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	I	J	K
401	MDR											
402	XDR											
403	RR											
404	Other											
405	Presumptive											
	DR TB											
406	Total											

Sl.	Patient Group	Total	Number	Cured	Treatment	Failure	Died	Lost to	Transferred	Still on	Not	Total
		number	of		Completed			follow	Out	Treatment	Evaluated	
		of DR TB	confirmed					up				
		patients	DR TB									
		registered	patients									
		during	registered									
		the	during									
		quarter	the									
			quarter									
		A	В	C	D	E	F	G	H	Ι	J	K
407	New											



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	T		_	ı	1		ı	ı	1
408	Failure after CAT								
	I								
409	Failure after CAT								
	II								
410	Relapse after								
	CAT II								
411	Relapse after								
	CAT I								
412	Treatment after								
	lost to follow up								
	CAT II								
413	Treatment after								
	lost to follow up								
	CAT I								
414	Delayed								
	Converters CAT								
	II								
415	Delayed								
	Converters CAT								
	I								
416	Close Contact of								
	DR TB with S/S								
	Pulmonary								
	Diagnosed								
417	New								
418	Previously								
	treated								
419	Unknown TB								
	treatment history								
	Extrapulmonary								
420	New								
421	Previously								
	treated								
422	Unknown TB								
	treatment history								
<u> </u>	a cathlent mistory		I		i			l .	

Section 5.0 Quantity and Cost of Drugs

Number of TB Drugs received by the facility last year (January 2016- December 2016-PLEASE CONSULT LAST TWO REQUISITION FORM DR TB-09)

Quarter |__|_| **Year** |__|_|

SL#	Drug name	Quantity measures	Quantity received from National TB Control Program (NTP) (units) CAT I Patients	Quantity received from National TB Control Program (NTP) (units) CAT II Patients	Total received in this quarter	Market price of drugs per unit
		A	В	C	D	E
501	Pyrazinamide 500 mg Tab					
502	Kanamycin 1 gm Vial (Only for IP)					
503	Ethionamide 250 mg Tab					
504	Cycloserine 250 mg Tab					
505	Ofloxacin 400 mg Tab					
506	Levofloxacin 250 mg Tab					
507	Moxifloxacin 400 mg Tab					
508	Clofazimine (Cfz 50 mg Tab)					
509	Amox/Clav 500/125 mg Tab					
510	Linezolid (Lzd)					
511	Capreomycin 1 gm Vial (Only for IP)					
512	PAS 4 gm sachet					
513	Omeprazole 20 mg Tab					



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514	Domperidone 10 mg Tab												•	I
515	Pyridoxine 25 mg Tab												•	
516	Multivitamin												•	
517	Alprazolam 0.5 mg													
518													•	
519													•	
520													•	
521														
522													•	
523	Others												•	
524	Syringe, 5 cc												•	
525	Distilled Water, 3 cc												•	
526													•	
527													•	
528													•	
529													•	
530													•	
531													•	
532													•	
533													•	
534													•	
535													•	

Quarter |__|_| Year |__|



SL#	Drug name	Quantity received from National TB Control Program (NTP) (units) for CAT I Patients	Quantity received from National TB Control Program (NTP) (units) for CAT II Patients	Total received in this quarter
		E	F	G
501	Pyrazinamide 500 mg Tab			
502	Kanamycin 1 gm Vial (Only for IP)			
503	Ethionamide 250 mg Tab			
504	Cycloserine 250 mg Tab			
505	Ofloxacin 400 mg Tab			
506	Levofloxacin 250 mg Tab			
507	Moxifloxacin 400 mg Tab			
508	Clofazimine (Cfz 50 mg Tab)			
509	Amox/Clav 500/125 mg Tab			
510	Linezolid (Lzd)			
511	Capreomycin 1 gm Vial (Only for IP)			
512	PAS 4 gm sachet			
513	Omeprazole 20 mg Tab			
514	Domperidone 10 mg Tab			
515	Pyridoxine 25 mg Tab			
516	Multivitamin			



517	Alprazolam 0.5 mg										
518											
519											
520											
521											
522											
523	Others										
524	Syringe, 5 cc										
525	Distilled Water, 3 cc										
526											
527											
528											
529											
530											
531											
532											
533											
534											
535											

Quarter |__|_| **Year** |__|_|

SL#	Drug name	Quantity received from National TB Control Program (NTP) (units) for CAT I Patients	Quantity received from National TB Control Program (NTP) (units) for CAT II Patients	Total received in this quarter
		Н	I	J
501	Pyrazinamide 500 mg Tab			
502	Kanamycin 1 gm Vial (Only for IP)			
503	Ethionamide 250 mg Tab			
504	Cycloserine 250 mg Tab			
505	Ofloxacin 400 mg Tab			
506	Levofloxacin 250 mg Tab			
507	Moxifloxacin 400 mg Tab			
508	Clofazimine (Cfz 50 mg Tab)			
509	Amox/Clav 500/125 mg Tab			
510	Linezolid (Lzd)			
511	Capreomycin 1 gm Vial (Only for IP)			
512	PAS 4 gm sachet			
513	Omeprazole 20 mg Tab			
514	Domperidone 10 mg Tab			
515	Pyridoxine 25 mg Tab			



516	Multivitamin										
517	Alprazolam 0.5 mg										
518											
519											
520											
521											
522											
523	Others										
524	Syringe, 5 cc										
525	Distilled Water, 3 cc										
526											
527											
528											
529											
530											
531											
532											
533											
534											
535											

Quarter |__|_| **Year** |__|_|

SL#	Drug name	Quantity received from National TB Control Program (NTP) (units) for CAT I Patients	Quantity received from National TB Control Program (NTP) (units) for CAT II Patients	Total received in this quarter
		K	L	M
501	Pyrazinamide 500 mg Tab			
502	Kanamycin 1 gm Vial (Only for IP)			
503	Ethionamide 250 mg Tab			
504	Cycloserine 250 mg Tab			
505	Ofloxacin 400 mg Tab			
506	Levofloxacin 250 mg Tab			
507	Moxifloxacin 400 mg Tab			
508	Clofazimine (Cfz 50 mg Tab)			
509	Amox/Clav 500/125 mg Tab			
510	Linezolid (Lzd)			
511	Capreomycin 1 gm Vial (Only for IP)			
512	PAS 4 gm sachet			
513	Omeprazole 20 mg Tab			
514	Domperidone 10 mg Tab			
515	Pyridoxine 25 mg Tab			



516	Multivitamin										
517	Alprazolam 0.5 mg										
518											
519											
520											
521											
522											
523	Others										
524	Syringe, 5 cc										
525	Distilled Water, 3 cc										
526											
527											
528											
529											
530											
531											
532											
533											
534											
535											

Section 7.0 Quantity and Cost of Laboratory Reagents/Supplies/Equipment

Number of Laboratory reagents/ supplies received by the facility last year (October 2015-September 2016- Laboratory Request Form)

Quarter 4 (July 2016- September 2016)

SL#	Reagent/Lab supplies' name	Quantity measures per patient	Quantity received from National TB Control Program (NTP) (units)	Quantity purchased from market (units)	Total received in this quarter	Market price of reagents per unit
		A	В	C	D	E
601	Basic fuchsin	1 gm/sm+				
602	Phenol crystals	5 gm/sm+				
603	Methanol (or denatured ethanol)	10 ml/sm+				
604	Methylene Blue	0.1 gm/sm+				
605	Sulphuric Acid conc.	33 ml/sm+				
606	Burning Spirit	50 ml/sm+				
607	Slides	36 pcs/sm+				
608	Sputum Containers	36 pcs/sm+				•
609	Bamboo Sticks	36 pcs/sm+				
610	Immersion Oil	2 ml/sm+				
611	Xylene	25 ml/sm+				
612	Toilet Paper Rolls	3 rolls/clinic				•
613	Filter Paper Pieces	20 pcs/ clinic				
614	Culture Media (Solid)					
615	Culture Media (Liquid)					



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616													•	
617													•	
618													•	
619													•	
620													•	
621													•	
622													•	

Quarter 3 (April 2016- June 2016)

SL#	Reagent/Lab supplies' name	f	Quai rom Cont (N'	Na trol TP)	tior	al I gra	ΓB ım				sed nits)	 tal 1 this	 	
601	Basic fuchsin				Ī									П
602	Phenol crystals													
603	Methanol (or denatured ethanol)													
604	Methylene Blue													
605	Sulphuric Acid conc.													
606	Burning Spirit													
607	Slides													
608	Sputum Containers													
609	Bamboo Sticks													
610	Immersion Oil							Ħ						
611	Xylene													
612	Toilet Paper Rolls													



613	Filter Paper Pieces									
614										
615										
616										
617										
618										
619										
620										
621										
622										

Quarter 2 (January 2016- March 2016)

SL#	Reagent/Lab supplies' name	f	rom Cont	Na rol	y rection Pro (ur	al T gra	rB m	Quan rom	-	_		 tal 1 this		
601	Basic fuchsin									J			IX	
602	Phenol crystals													
603	Methanol (or denatured ethanol)													
604	Methylene Blue													
605	Sulphuric Acid conc.													
606	Burning Spirit													
607	Slides													



608	Sputum Containers									
609	Bamboo Sticks									
610	Immersion Oil									
611	Xylene									
612	Toilet Paper Rolls									
613	Filter Paper Pieces									
614										
615										
616										
617										
618										
619										
620										
621										
622										

Quarter 1 (October 2016- December 2016)

SL#	Reagent/Lab supplies' name	Qua fron Cor (N	n Na	atio l Pr	nal	TB am			ased nits)	Total received in this quarter
601	Basic fuchsin							141		14
602	Phenol crystals									



		П	1	1	1	1	I I	1	1	1	1
603	Methanol (or denatured ethanol)										
604	Methylene Blue										
605	Sulphuric Acid conc.										
606	Burning Spirit										
607	Slides										
608	Sputum Containers										
609	Bamboo Sticks										
610	Immersion Oil										
611	Xylene										
612	Toilet Paper Rolls										
613	Filter Paper Pieces										
614											
615											
616											
617											
618											
619											
620											
621											
622											

Section 8.0 Cost of Other Supplies

Number of Supplies received by the facility last year (October 2015-September 2016)

Quarter 4 (July 2016- September 2016)

SL#	Supplies' name	Quantity measures	f	rom Con	ı Na trol	y re tior Pro (u	ıal ' ogra	TB am			ased nits)	tal ı this			M	arke	ce of unit	plies
		A				В				C			D				E	
701	TB Register																•	
702	Treatment Card																•	
703	Pen																•	
704	Paper																•	
705	Box																•	
706	Soap																•	
707	Towel																•	
708	Boxes																•	
709	Таре																•	
710	Raincoat																•	
711	Torch light																•	
712	Umbrella																•	
713	Drug Baskets																•	
714	Kit Bag																•	
715	Poster																•	
716	Sticker																	
717	Leaflet																	
718	Flip chart																	



S	
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719	Flash Chart												•	
720	Brochure												•	
721														
722														
723														
724														
725														
726														
727														
728														
729														
730														

SL#	Supplies' name	fro Co	anti m N ntro NTP	atio	nal ' ogra	ГВ am	_	uan om 1	-	_	sed nits)	 	rece qua	
				F						G			Н	
701	TB Register													
702	Treatment Card													
703	Pen													
704	Paper													
705	Box													
706	Soap													
707	Towel													



708	Boxes									
709	Tape									
710	Raincoat									
711	Torch light									
712	Umbrella									
713	Drug Baskets									
714	Kit Bag									
715	Poster									
716	Sticker									
717	Leaflet									
718	Flip chart									
719	Flash Chart									
720	Brochure									
721										
722										
723										
724										
725										
726										
727										
728										
729										
730										



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SL#	r 2 (January 2016- Ma Supplies' name	f	Quant rom N Contr	Natio ol Pr	eceived nal TB ogram nits)			urcha et (u	To		received in s quarter			
			1 1	I		1	J	1	_	-	K			
701	TB Register												<u> </u>	
702	Treatment Card													
703	Pen													
704	Paper													
705	Box													
706	Soap													
707	Towel													
708	Boxes													
709	Tape													
710	Raincoat													
711	Torch light													
712	Umbrella													
713	Drug Baskets													
714	Kit Bag													
715	Poster													
716	Sticker													
717	Leaflet													
718	Flip chart													
719	Flash Chart													
720	Brochure													



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721									
722									
723									
724									
725									
726									
727									
728									
729									
730									

Quarter 4 (October 2015- December 2015)

SL#	Supplies' name	f	Quai rom Cont (N'	Na trol	tion	al I gra	ГВ ım	Quantity purchased from market (units)						Total received in this quarter					
					L						M			N					
701	TB Register																		
702	Treatment Card																		
703	Pen																		
704	Paper																		
705	Box																		
706	Soap																		
707	Towel																		
708	Boxes																		
709	Tape																		

710	Raincoat										
711	Torch light										
712	Umbrella										
713	Drug Baskets										
714	Kit Bag										
715	Poster										
716	Sticker										
717	Leaflet										
718	Flip chart										
719	Flash Chart										
720	Brochure										
721											
722											
723											
724											
725											
726											
727											
728											
729											
730											

Section 8.0 Other Capital Items

Number of Items used by the facility

SL#	Capital Items	Quantity (Unit)	Co	ece Nat ontr	iona ol P	tity from al TB rogram units)	p n	urc	hase	ntity ed from (units)	7	Γota	al is use		ing	Life In Y		M	Market price of item per unit			n per unit	
		A			В	1			C	! ;			D)]	E		,		F	1	1
801	Microscope																						
802	GeneXpert																						
803	Weight Scale																						
804	Signboard																						
805	Computer																						
806	Printer																						
807	Smart Phone																						
808	App development																						
809	Motor Cycle																						
810	Bicycle																						
811	Car																						
812	Ambulance																						
813	LPA																						
814	LJ																						
815	MGIT																						
816																							
817																							
818																							
819																							



Section 9. Other Costs

Section 9.1 Cost of Supervision (facility supervisory visits conducted in last year (October 2015-September 2016)

SL#	Title of person conducting supervisory visit	Number of visits in last year (Oct 2015- Sep 2016)	Duration of visit (total for all visits in days)	Per Diem (total)	Travel expenses (total)	Other expenses	What % of costs paid by facility?
	A	В	C	D	E	F	Н
901							
902							
903							
904							

Section 9.2 Travel cost for Drug /Attending Meeting/Training

SL#	Title of person conducting travel	Number of travel in last year (Oct 2015-Sep 2016)	Purpose of the travel (Use Code from below)^	Duration of visit (total for all visits in days)	Per Diem (total)	Travel expenses (total)	Other expenses	What % of costs paid by facility?
	A	В	С	D	E	F	H	I
905								
906								
907								
908								

[^]Code: 1. Receiving drug 2. Attending Meetings 3. Training 4. Others (Please specify)



Section 9.3 Cost of Other Activities

No	Questions and Filters	Coding categories	Record Response	Skip
909	Was Health education on TB organized by the DOTS center?	Yes		If 2 →911
910	If Health education on TB was organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
911	Was DOTS committee meeting organized by the DOTS center?	Yes1 No2		If 2 →913
912	If DOTS committee meeting was organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
913	Were Contact Tracing activities organized by the DOTS center?	Yes		If 2 →915
914	If Contact Tracing activities were organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
915	Were Refresher training activities organized by the DOTS center?	Yes		If 2 →917
916	If Refresher training activities were organized, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
917	Were Other meetings organized by the DOTS center?	Yes		If 2 →919
918	If Other meetings were organized, How much money was paid for this activity in last year (October 2015- September 2016)?	IF THIS ACTIVITY WAS NOT ORGANIZED AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
919	Were Incentive payments were provided to the providers by the DOTS center?	Yes		If 2 →921



920	If incentive payments were given to the providers, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS PAYMENT WAS NOT MADE AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
921	Were Incentive payments were provided to the patients by the DOTS center?	Yes		If 2 →923
922	If incentive payments were given to the patients, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS PAYMENT WAS NOT MADE AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	
923	Were Incentive payments were provided to the community members by the DOTS center?	Yes		If 2 →END
924	If incentive payments were given to the community members, How much money was paid for this activity in last year (October 2015-September 2016)?	IF THIS PAYMENT WAS NOT MADE AND/OR NO MONEY WAS PAID, RECORD "00000"	Taka	

Thank you for your cooperation! Is there anything you would like to ask or say?

Comments by Interviewer:

Date, Signature by Interviewer: