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Developing and testing community-based tuberculosis (TB) screening intervention to increase TB referral, case detection and knowledge among sexual minority people in urban Bangladesh: a mixed-method study protocol

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ABSTRACT

Introduction Although Bangladesh is a country of generalised tuberculosis (TB) epidemic, the HIV prevalence is low among general populations, and 3.9% among key populations. Despite the high possibility of HIV–TB co-infection, scientifically tested approaches for increasing TB case detection among sexual minority people are yet to be developed and implemented in Bangladesh. Such approaches could foster service delivery linkages between communities and the government health system. Findings of this experimental research are likely to provide new insights for programme managers and policy planners for adopting similar approaches in order to enhance TB referral, thus ultimately increasing TB case detections and reducing the likelihood of TB-related mortalities and morbidities, irrespective of HIV status.

Methods and analysis This operational research will follow a quasi-experimental design, applying both qualitative and quantitative methods, in two drop-in centres in three phases. Phase 1 will encompass baseline data collection and development of a community-based TB screening approach. In phase 2, the newly developed intervention will be implemented, followed by end-line data collection in phase 3. Qualitative data collection will be continued throughout the first and second phases. The baseline and end-line data will be compared both in the intervention and comparison areas to measure the impact of the intervention.

Ethics and dissemination Ethical approval was obtained from the Institutional Review Board of International Centre for Diarrhoeal Disease Research, Bangladesh. The findings will be disseminated through diverse scientific forums including peer-reviewed journals, presentation at conferences and among the policy-makers for policy implication. The study started in January 2019 and will continue until June 2020.

INTRODUCTION

HIV epidemic poses as an impediment to the effective control of tuberculosis (TB) worldwide.1 Clinical evidence has alluded to a causal relationship between TB and HIV.2 HIV is considered as the most predominant risk factor for the development of Mycobacterium tuberculosis infection to TB disease.2 3 Likewise, TB infection accelerates the progression of HIV infection, reduces survival and even predisposes the infected individual to premature death.4 On the global scale, an estimated 10.0 million
(range 9.0–11.1 million) people were newly diagnosed with TB in 2018, which also constituted to 8.6% of the People Living with HIV (PLHIV). An estimated 1.2 million (range, 1.1–1.3 million) deaths due to TB disease occurs across the globe in 2018, in addition to 251,000 (range, 223,000–281,000) TB deaths happened among PLHIV. This is because PLHIV are 29 (26–31) times more vulnerable to TB than HIV negative individual. In this context, TB is considered as the leading preventable cause of death especially among PLHIV.

Bangladesh is a country with generalised TB epidemic, though it has a low HIV prevalence of less than 0.01% among general populations. However, HIV surveillance findings demonstrated that 3.9% of the key populations (KPs) at risk of HIV contracted HIV infection. Between 1999 to 2002, a study conducted by International Centre for Diarrhoeal Disease Research, Bangladesh (icddr,b) among 958 patients to determine HIV prevalence among patients with TB in Dhaka demonstrated an estimated 0.1% TB/HIV coinfection. Nonetheless, recent data reported by Government of Bangladesh (GoB) from 2009 to 2014 showed that merely 1% of the registered TB patients in the country were documented to know their HIV status. In 2017, the prevalence of HIV among the diagnosed TB patients (n=3917) was 0.36% whereas approximately 16% of the PLHIV (n=559) were tested positive for TB, thus illustrating bidirectional upward trends in TB/HIV coinfection rates over the years. Moreover, despite of being a country of high TB burden, TB case detection rate (the yearly number of newly notified cases per 100,000 population) in Bangladesh is still unsatisfactory as at a 75% case detection rate as of 2018.

In Bangladesh, TB control interventions are being implemented in healthcare facilities throughout the country, from the central to peripheral levels. HIV prevention programmes are primarily concentrated in 23 priority districts based on concentration of HIV positive cases and KPs. Thus, there is inadequate linkage between TB and HIV prevention programmes, that engendering low identification of TB cases among KPs. To foster a functional collaboration between TB and HIV programmes, GoB designed a roadmap of collaboration between the National Tuberculosis Control Program (NTP), AIDS/STD Program, key stakeholders (BRAC, icddr,b, Save the Children, etc) and implementing partners (Damien Foundation, CARE,B, etc) since 2009 with the aim to institute programmatic linkages at each level of healthcare, although with unsatisfactory outcomes.

Worldwide, KPs, including sexual minority people (i.e., men who have sex with men (MSM), male sex workers (MSW) and transgender women (locally known as hijra)), people who inject drug, female sex workers and PLHIV, are considered vulnerable to TB. These populations are particularly susceptible to TB infection due to their hidden and hard-to-reach nature along with their unwillingness to visit lack of access to healthcare facilities like other vulnerable populations. In particular, sexual minority people are more vulnerable not only because of their propensity to contract TB/HIV coinfections but also their highly stigmatised nature. Case notification rates also remain low among these populations due to the implementation of passive case detection of TB control strategy which often exclude marginalised populations, thus leaving them undiagnosed and untreated for TB. Experiences from some countries have shown that successful implementation of active case finding strategies, resulted in increased detection of TB case, especially for hidden and/or hard-to-reach populations. For example, a recent study conducted in tribal India revealed that active case finding interventions contributed to a 52% increase in testing rates and 84% increase in TB case notifications. Similarly, a recent quasi-experimental study that was conducted in Cambodia showed that the TB case notification rates were significantly higher (<0.05) among the intervention group (which adopted active case finding) compared with the control group (which employed passive case finding). Likewise, a recent cohort study conducted in China documented that early case detection rates significantly increased among the higher risk groups, such as PLHIV.

Several studies have suggested that TB screening should be encompassed in an integrated HIV intervention, while also addressing the other healthcare needs of PLHIV. For example, a community outreach model was adopted in rural Mozambique, where TB, HIV, malaria and nutritional services are packaged in a single service delivery point, which helped improve acceptability and satisfaction. However, the acceptability of TB screening programmes among MSM has not been a substantial point of focus in research or programme implementation. In fact, many NTP programmes in low/middle-income countries are oblivious to the role that MSM may contribute in TB epidemics. It was evidenced that even MSM, rooted in higher income countries, are at increased risk for TB. The acceptability of TB screening was found to range from 43% to 91% in many of the studied countries, although the underlying reasons were not specified. TB screening service acceptability among transgender women is also an understudied area. The reach and recruitment rates of transgender cases were hard to quantify since many of the studies used word of mouth or peer referrals through snowball sampling.

In Bangladesh, 77% of the transgender women (locally known as hijra) articulated their willingness to provide sputum samples. According to Mitchell et al, it was inconvenient to reach transgender participants through traditional contact techniques. Rather, the Center for Disease Control recounted an experience where they attempted to recruit transgender women through exploring their residences, social networks and other social spheres as well as common social spheres. Transgender women in Bangladesh, indicated that they felt more comfortable visiting a community-based healthcare facility, particularly if they did not disclose their sexual
orientation and if the facilities were catered to sexual minority populations.17

icddr,b is currently performing verbal TB screening (‘It is a standardised verbal questionnaire for investigating presumptive TB cases on the presence of the following symptoms and signs: cough for more than 2 weeks, haemoptysis, chest pain, breathlessness, evening rise of temperature/night sweating, anorexia, and weight loss/weakness’) for sexual minority people through drop-in centres; STI, sexually transmitted infections.

Despite high possibility of TB/HIV co-infection, a scientifically tested approach for increasing TB case detection among sexual minority people is yet to be developed and implemented in Bangladesh. Therefore, a community-based TB screening approach will be tested among sexual minority people. Such an approach is expected to encourage sustainable service delivery linkages between community systems and the Government health systems. Findings of this study are likely to provide new insights for programme managers (PMs) and policy planners for developing and implementing a community-based TB screening approach to enhance TB referral which ultimately increase TB case detection among sexual minority people.

**RESEARCH OBJECTIVES**

**Primary objective**

To develop and assess a community-based TB screening approach to enhance TB referral for increasing case detection among sexual minority people in Dhaka city, Bangladesh.

**Secondary objectives**

1. To understand barriers and scopes of the existing TB screening approach for sexual minority people.
2. To measure knowledge of TB among the participants.
3. To develop and implement a community-based TB screening approach.
4. To assess the impact of community-based TB screening approach on TB screening, presumptive TB case referral, TB case detection and knowledge related to TB.

**METHODS AND ANALYSIS**

**Research design**

Under the operational research framework, by applying mixed methods (ie, qualitative and quantitative) approach, the study will follow a quasi-experimental design with enlisted sexual minority people in selected DICs.33 Both preintervention and postintervention measurements will be facilitated among non-randomly selected control groups.34 The participants of intervention site will be compared with the participants of the comparison site. To measure changes between two time points, baseline and end-line in the outcome of interest and between the two groups, an analysis will be conducted.

**Research plan**

The study will ensue through the following phases adopted from the operational research framework recommended by WHO and the Global Fund (GF).35—figure 1 shows the activities of this study:

**Phase 1: formative phase**

1. Qualitative data collection to explore implementation barriers and scopes of existing TB screening approach.
2. Survey on sociodemographic characteristics, knowledge and risk behaviour related to TB among the sexual minority people of the both intervention and comparison catchment areas.
4. Baseline data collection on the number of TB screening, referral of presumptive TB cases to directly observed treatment, short-course (DOTS, also known as TB-DOTS) centres and TB case detection within existing DIC STI clinic-based approach of TB screening at both intervention and comparison sites.

**Phase 2: intervention phase**
1. Implementation of community-based TB screening intervention.
2. Qualitative data collection to identify implementation facilitators and challenges of community-based TB screening approach.

**Phase 3: comparison phase**
1. End-line data collection in both intervention and comparison sites on the number of TB screening, referral of presumptive TB cases to DOTS centres, and TB case detection. The comparison will be done between end-line and the baseline data in both intervention and comparison sites.
2. Post test to measure knowledge change as a result of behaviour change intervention (BCI) among sexual minority people in the intervention DIC area and comparison DIC area at the end of the intervention period, of whom the latter will not get BCI. Comparisons will subsequently be drawn within and between these two areas.

**Development of community-based TB screening model**

A community-based TB screening model will be developed based on the first phase findings. At the beginning of the second phase, researchers will develop a training module for the peer educators (PEs) based on the needs assessment, map TB DOTS centres and devise DOTS referral strategies. Sensitisation sessions will be conducted with the DOTS service providers and community leaders. Researchers will train PEs of the intervention area on the TB screening process and then be assigned to serve a separate list of sexual minority people. A work plan will be formulated to screen all sexual minority people at least once during the intervention period. After taking their verbal consents, they will be screened at spots/residences/dera (residence of hijra leader) using a simplified structured verbal TB screening form, while maintaining confidentiality. If the case is negative, post screening information and counselling will be provided and they will be encouraged to seek support if suspected with TB symptoms. If screened positive, they will undergo counselling and be referred to the nearest TB DOTS centre for testing, either through self- or accompanied referral. After the test results are ready, the PE or the sexual minority people themselves will collect the report, and if anyone is tested positive she/he will be linked to nearby TB DOTS centre for treatment and followed up for adherence. PEs will also conduct specialised educational sessions on TB at outreach sites where they will distribute visual materials and discuss TB risks and routes of transmission.

**Study area**

![Study area](image)

**Figure 2** Developing and testing community-based tuberculosis (TB) screening intervention for sexual minority population (men who have sex with men, male sex workers, transgender in Dhaka, Bangladesh Study location/area, January 2019–June 2020). DIC, drop-in centre.

**Study location**

This experimental study will be conducted in two DICs (ie, Jatrabari and Darus Salam DIC) in Dhaka city (figure 2). These DICs are operated by a non-government organisation namely Bandhu Social Welfare Society, under the GF project of icddr,b. The intervention site, Jatrabari DIC, is located in the southern part of Dhaka city. Darus Salam DIC, the comparison site, is in the western part of Dhaka city (Mirpur). These DICs were selected purposively due to similar profiles in terms of duration of service delivery, the concentration of HIV positive cases and number of sexual minority people served.

**Study population**

The sexual minority people enlisted for HIV prevention services in these two DICs will be selected for this study. Operational definitions for the MSM, MSW and hijra are provided in table 1 which are currently used in the provision of HIV prevention services to sexual minority people.

<table>
<thead>
<tr>
<th>Population</th>
<th>Operational definitions</th>
</tr>
</thead>
<tbody>
<tr>
<td>MSM</td>
<td>Men who have had sex with men (with consent) within the last 1 year irrespective of whether or not they have sex with women or have a social or personal gay or bisexual identity, but do not sell sex</td>
</tr>
<tr>
<td>MSW</td>
<td>Man who sell sex to other men in exchange of money or gifts in the last 3 months</td>
</tr>
<tr>
<td>Hijra</td>
<td>Those who identify themselves as part of a traditional hijra subculture and who maintain the guru–chela hijra hierarchy</td>
</tr>
</tbody>
</table>

**Table 1** Definitions of the men who have sex with men (MSM), male sex workers (MSW) and hijra

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For qualitative interview, the following research participants will be recruited purposively in this research:
1. Sexual minority people (MSM, MSW and hijra).
2. DIC staff (ie, DIC managers, medical assistants, outreach supervisors and PEs).
3. Staff of adjacent DOTS centres (ie, physicians, nurses, technicians, etc).
4. Experienced PMs and researchers working with KPs.
5. PMs of NTP.
6. Representative from principal and sub-recipients of the GF project.

Inclusion criteria
1. All sexual minority people aged 18 years or more, and enlisted in the service list of the intervention site (for intervention).
2. According to the sample size, a subset of sexual minority people of both intervention and comparison areas (for baseline and end-line survey).
3. Health service providers and other staff members of the intervention and comparison DICs (for qualitative interview only).
4. Senior management staff members of the PR and SRs, staff of adjacent DOTS centre, experienced PMs of NTP and researchers working with KPs (for qualitative interview only).
5. Those who provide verbal and/or written consent to participate in this study.

Exclusion criteria
Participants who refuse to participate in this research, and sexual minority people who are not enlisted in the intervention DIC. In addition, minors, below 18 years, will not be included in the study.

Study period
The study commenced in January 2019 and is anticipated to continue until June 2020. The Gantt chart is given in table 2:

Sample size and sampling
Quantitative component
To conduct the baseline survey, in both the intervention and comparison sites, the calculation of sample size was done using the following standard formula:\textsuperscript{37}

\[ n = \frac{Z_{1-\alpha}^2p_1(1-p_1)+Z_{1-\beta}^2p_2(1-p_2)}{(p_2-p_1)^2} \times D^2 \]

In the above formula:
- \( D \) = design effect = 2 (due to cluster sampling method).
- \( p_1 \) = assumed proportion of the outcome variable at the time of baseline survey.
- \( p_2 \) = the target proportion at the end line, so that \((p_2 - p_1)\) is the magnitude of change that we want to be able to detect.
- \( Z_{1-\alpha} \) = the Z-score corresponding to desired level of significance = 1.645.\textsuperscript{37}
- \( Z_{1-\beta} \) = the Z-score corresponding to desired level of power = 0.83.\textsuperscript{37}

A literature review was carried out in order to find out estimates of TB knowledge in terms of cause, route of transmission and prevention. The results depicted that the knowledge of cause, routes of transmission and prevention of TB ranged from 2% to 44%, 22% to 88% and 14% to 68%, respectively (table 3).

The sample size was calculated in order to detect 20% changes (1-way change detectable) in the knowledge of TB between two time points with the design effect of 2.0, 95% CI and 80% power. Separate sample sizes were calculated for each of the categories and were adjusted for 5% drop-outs during the interview (table 4). Finally, the largest sample size (n=160) will be selected to conduct the baseline survey each in the intervention and comparison areas. It is worth mentioning that all the findings were from the general population and no literature was found among the sexual minority people. Thereafter, using programmatic experience among sexual minority people in Bangladesh, suitable values were assumed to calculate sample sizes (table 4).

The baseline survey will be conducted using time location sampling.\textsuperscript{37} The trained interviewers will interview sexual minority people at all congregating areas (spots) at a particular time window, usually between 17:00 and 22:00. After collecting the list of spots and the sexual minority people covered throughout each spot, the total sample size will be proportionately distributed among each spot. In each spot, the respondents will be randomly chosen for interview to collect data on sociodemographic characteristics, knowledge of and risk behaviours pertaining to TB and so on. Identification numbers (IDs) from the mother list will be used to avoid duplications during the survey. Similar sample size and sampling method will be used to conduct end line survey both in intervention and comparison areas.

Qualitative component
Non-probabilistic purposeful maximum diversity sampling will be used in the qualitative component. Based on our previous experience of working with sexual minority people, we plan to facilitate in-depth interviews (IDIs) with 15–18 sexual minority people, 10–15 key informants interviews (KIIIs) with service providers. In addition, two focus group discussions (FGDs) will be conducted with sexual minority people and service providers. The sample size may vary depending on the points of data saturation and redundancy.\textsuperscript{38}

Data collection method
Qualitative data collection
Several qualitative data collection approaches will be employed, such as IDIs with the sexual minority people, KIIIs with the service providers and FGDs with sexual minority people and service providers.
### Table 2  Gantt chart

<table>
<thead>
<tr>
<th>Activities</th>
<th>2019 Year</th>
<th></th>
<th></th>
<th></th>
<th>2020 Year</th>
</tr>
</thead>
</table>
|                                                 | January   | February             | March                | April
|                                                 | January   | February             | March                | April
| 1. Training of staff                           | January   | February             | March                | April
| 2. Piloting                                     | January   | February             | March                | April
| Phase-1                                         | January   | February             | March                | April
| 1. Baseline data collection:                    | January   | February             | March                | April
| 2. Quantitative survey                          | January   | February             | March                | April
| 3. Qualitative                                  | January   | February             | March                | April
| 2. Program data                                 | January   | February             | March                | April
| 2. Development of intervention model            | January   | February             | March                | April
| Phase-2:                                        | January   | February             | March                | April
| 1. Implementation of intervention               | January   | February             | March                | April
| 2. Quantitative data collection                 | January   | February             | March                | April
| Phase-3:                                        | January   | February             | March                | April
| 1. End-line data collection:                    | January   | February             | March                | April
| 2. Programme data                               | January   | February             | March                | April
| Data entry, cleaning, analysis and tabulations  | January   | February             | March                | April
| (quantitative)                                  | January   | February             | March                | April
| Data transcription and analysis                 | January   | February             | March                | April
| (Qualitative)                                   | January   | February             | March                | April
| Preliminary report writing                      | January   | February             | March                | April
| Final report and dissemination                  | January   | February             | March                | April

- **January**: January, February, March
- **April**: April, May, June
- **July**: July, August
- **September**: September, October
- **November**: November, December
- **December**: December

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Note: The Gantt chart visually represents the timeline of activities for the project, with different phases marked and their respective months indicated.
Survey on sociodemographic characteristics, knowledge of and risk behaviour related to TB
Survey on sociodemographic characteristics, knowledge of and risk behaviour related to TB will be conducted among selected numbers of sexual minority people of both intervention and comparison areas during baseline and end-line surveys at an approximate interval of 6 months.

Baseline and end-line programmatic quantitative data collection
At the start and end of the intervention phase of the study, data will be collected from programme report of the GF project of icddr,b on the total number of beneficiaries screened for TB, total number of referred cases for presumptive TB and number of TB case detections within the extant STI clinic-based TB screening approach as well as the new intervention approach at both the intervention and comparison sites.

Data collection tool
A structured TB screening format will be used for verbal TB screening. The survey will be conducted via a semi-structured questionnaire in standard Bengali. This questionnaire will encompass the sociodemographic profile, knowledge of and risk behaviours related to TB. Moreover, open-ended guidelines will be used for qualitative interviews and FGDs. All the guidelines and questionnaires will be field tested. Based on field testing, these data collection tools may need to be refined, where appropriate, in order to facilitate the data collection

Table 3 Literature review of the knowledge of tuberculosis (TB)

<table>
<thead>
<tr>
<th>Indicators</th>
<th>Estimates of the indicators</th>
<th>Target group</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Knowledge of TB</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Cause of TB: percentage of beneficiaries correctly identify that TB is caused by bacteria</td>
<td>43.9% 2% 37.7%</td>
<td>General population</td>
<td>Wandwalo and Mørkve</td>
</tr>
<tr>
<td>2. Route of transmission of TB: percentage of beneficiaries mentioned that TB is transmitted by air (by talk/by cough/by sneezing/by spitting TB germ)</td>
<td>53.7% 22%–34% 55.5% 56% 88.0%</td>
<td>General population</td>
<td>Wandwalo and Mørkve</td>
</tr>
<tr>
<td>3. Methods of prevention: percentage of beneficiaries mentioned that TB can be prevented by taking BCG vaccination/covering mouth and nose when patient with TB is coughing or sneezing or spitting or talking</td>
<td>14% 68.2% 29.1%–65.7%</td>
<td>General population</td>
<td>Anjum et al</td>
</tr>
</tbody>
</table>

Table 4 Proposed sample size to conduct survey with sexual minorities (men who have sex with men, male sex workers, transgender) in Bangladesh (January 2019–June 2020)

<table>
<thead>
<tr>
<th>Indicators used in the calculation of sample size</th>
<th>Suitable values assumed for the sexual minority people</th>
<th>One-way change detectable</th>
<th>Design effect</th>
<th>Required Sample Size*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Knowledge of TB:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Cause of TB: Percentage of beneficiaries correctly identify that TB is caused by bacteria</td>
<td>40%</td>
<td>20%</td>
<td>2.0</td>
<td>159</td>
</tr>
<tr>
<td>2. Route of transmission of TB: percentage of beneficiaries mentioned that TB is transmitted by air (by talking/by coughing/sneezing/spitting TB germs)</td>
<td>50%</td>
<td>20%</td>
<td>2.0</td>
<td>152</td>
</tr>
<tr>
<td>3. Methods of prevention: percentage of beneficiaries mentioned that TB can be prevented by taking BCG vaccination/covering mouth and nose when patient with TB is coughing/sneezing/spitting/talking</td>
<td>50%</td>
<td>20%</td>
<td>2.0</td>
<td>152</td>
</tr>
</tbody>
</table>

*5% was added in order to adjust drop-out during survey. TB, tuberculosis.
process. All qualitative and quantitative interviews will be conducted in standard Bengali.

Outcome variables

Data to measure outcome variables will be extracted from the ongoing HIV prevention programme from both intervention and comparison areas. These data will be expressed in terms of percentage points to assist statistical analysis. The outcome measures are defined as follows:

1. Number of TB screenings: number of beneficiaries screened for TB at the community, which will be expressed as percentage points in relation to the total number of beneficiaries enlisted in the mother list.
2. Number of presumptive TB case referrals: number of beneficiaries referred to DOTS centres for TB testing, which will be expressed as percentage points in relation to the total number of beneficiaries screened at the community.
3. Number of TB case detections: number of beneficiaries who will be tested positive for TB, which will be expressed as a percentage points in relation to the total number of presumptive cases tested.

Data on knowledge related to TB will be collected from quantitative surveys before intervention (baseline) and after intervention (end line) both in the intervention and comparison areas. This outcome variable will measure the knowledge of TB, as defined below.39

1. Cause of TB: correctly able to identify that TB is caused by bacteria.
2. Route of transmission of TB: TB is transmitted by air (by talking/coughing/sneezing/spitting TB germs).
3. Methods of prevention: TB can be prevented by taking BCG vaccination/covering mouth and nose when patient with TB is coughing/sneezing/spitting/talking.

Data analysis

Quantitative data

Statistical analysis will be conducted using Stata V.13.0 (Stata Corp). Epi Info (V.6.03; Centres for Disease Control and Prevention, Atlanta, Georgia, USA) will be used for data entry. The populations and measurements will be presented through descriptive statistics. Differences in the outcome variables between intervention and comparison groups at the baseline and at end line will be determined by the independent t-test for parametric continuous data, χ² or Fisher’s exact test for categorical data, and Wilcoxon rank-sum test for non-parametric continuous data. Differences in the outcome variables within intervention and comparison areas between two time points will be determined using the paired t-test for parametric continuous data whereas, Wilcoxon signed rank test for non-parametric continuous data and χ² or Fisher’s exact test for categorical data. P values of <0.05 will be considered as a significant level in all comparisons. In addition, 95% CI will be reported for all quantitative and categorical variables in the study.

Qualitative data

Data will be collected using digital recorders and systematically stored on the computer on a daily basis for a prolonged timeframe, adhering to the requirement of Institutional Review Board (IRB) and data policy of icddr,b. In addition, data collection and data analysis processes will occur in synergy, as qualitative data collection and analysis is an ongoing and reflexive process, and this process will allow for the identification of saturation points.40 41 All qualitative data will be analysed manually by adopting line-by-line content, contextual and thematic analysis strategies; and coding and recoding data into themes and subthemes. The results will be summarised and presented keeping in mind the local context, and some of the findings will be presented verbatim in order to substantiate or reflect noteworthy ideas and perspectives.

Patient and public involvement

As mentioned in the ‘Study population’ section, sexual minority people, enlisted under the intervention and comparison DICs, will be included to explore the research questions and measure outcomes on TB screening, TB presumptive case referral, TB case detection and TB knowledge. While this research did not directly engage them in the design or conceptualisation of the study, researchers planned to involve them throughout the data collection and analysis period. The study participants will be first engaged before the data collection period, when they will be preliminarily oriented about the study objectives and procedure. They will also be involved in facilitating access to and recruiting additional participants, including hidden and hard-to-reach sexual minority people. To design a viable and context-specific community-based TB screening model and optimise selection of actors (ie, community health workers, existing PE, new peer volunteer as TB screener, etc) for engagement in TB screening, insights will be sought from sexual minority people and service providers via formative research. Member checking sessions will be also conducted to verify correct findings and interpretative analyses. Sexual minority people will be the key recipients of the intervention. Guidance for Reporting Involvement of Patients and Public (GRIPP) 2 reporting checklist for patient and public involvement is available in online supplemental table 1.

Study limitations

One of the major limitations is unlike randomised controlled trial, the quasi-experimental study design is unable to establish a strong causal association between intervention and outcome due to the lack of randomisation.42 However, to overcome this limitation, a similar comparison group will be assigned, which will not receive the intervention. Moreover, qualitative methods will be integrated with quasi-experimental methods to support or refute the findings of the quantitative evaluation and to assess the effectiveness of the intervention. Thus, the causal inference will be strengthened. Another potential
limitation might be the chance of insufficient control of some confounding factors, which may influence or contaminate the outcome.43 This limitation will be mitigated through employing preintervention and postintervention measurement, and statistical analyses that adjust (‘control’) the confounder(s). In addition, any positive or negative findings that may influence the expected outcome will be documented and reported to describe the context broadly within which generally causal relationships are constructed and deconstructed. Since the study will adopt the non-probabilistic purposive sampling approach because of the stigmatised and hidden nature of the sexual minority people, this may incur selection bias.43 Therefore, maximum variation sampling will be attempted to minimise this bias.

ETHICS AND DISSEMINATION

Participation in the research process will be voluntary. To ensure maximum congeniality of the informants, verbal consent will be attained from sexual minority people during IDIs and FGDs whereas written consent will be obtained from representative of PRs and SRs, service providers from DOTs centres, experienced PMs, and researchers during KIIs and FGDs. Verbal and/or written consent will also be taken from sexual minority people prior to the quantitative survey, as well as the pretest and post-test phases. Experiences of working with sexual minority people suggest that many sexual minority people withhold information about their identities and practices that characterise them as a KP (eg, involvement in sex trade, sexual orientation) in written form in apprehension of identity disclosure. Structural barriers such as criminalisation of same-sex behaviour and socioreligious stigma impede the uptake of written consent from sexual minority people.

Research participants will be briefed about the objectives of the study, the data collection process and topics of discussion. The researchers ascertained the anonymity of the participants’ responses. Moreover, no identifying information will appear in the data collection tools. If any research participant declines the recording of his responses, researchers will take written notes.

During intervention, if any participant is tested positive for TB, then she/he will be linked to the adjacent DOTs centre, to ensure treatment and follow-up of the treatment adherence.

Ethical clearance was obtained from icddr,b’s IRB which follows international ethical standards to ensure confidentiality, anonymity and informed consent.

Study findings will initially be disseminated at the organisation and then subsequently, among policymakers and relevant stakeholders to inform the policy for implication. In addition, findings will be disseminated to a diverse audience via different scientific forums such as peer-reviewed journals, and presentations at national and international conferences. Further dissemination initiatives include translational and implementation strategies such as the impact of findings in the national policy.

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