Non-consensual disclosure of infectious drug-resistant tuberculosis status in the occupational context: Health workers stuck between a rock and a hard place

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Background. The burden of drug-resistant tuberculosis (DR-TB) remains high in South Africa (SA), despite ongoing control efforts. DR-TB patients who are infectious and continue to work pose a substantial risk of spreading the disease at the workplace. When such patients refuse or interrupt treatment, and are also unwilling to disclose their status at work, they pose an infectious risk in the workplace, creating a conflict between public health good and individual human rights. Should health workers breach confidentiality and disclose patients’ DR-TB status, or take no action while patients pose transmission risks in occupational settings? Non-consensual disclosure of the infectious DR-TB status of such patients is an intervention intended to address this occupational health risk.

Objectives. To provide professional and ethical guidance for health workers in dealing with non-consensual disclosure of the DR-TB status of patients who are unwilling to disclose, continue to go to work and remain infectious.

Methods. A comprehensive literature search was conducted to draw on evidence relating to public health best practices and human rights normative standards. We used the findings, along with guidance from DR-TB programme managers, clinicians, occupational health physicians and human rights lawyers to develop an occupational health protocol for disclosure.

Results. This protocol establishes the normative, ethical and legal framework for dealing with the complexities of non-consensual disclosure, and provides a set of practical standard operative procedures for health workers in the SA setting. It also provides a legal safeguard to protect health workers from potential litigation that may result from such duty.

Conclusion. Non-consensual disclosure to address the spread of DR-TB in occupational settings should be based on professional, ethical and legally sound guidance for health workers.

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Drug-resistant tuberculosis (DR-TB) is of enormous public health concern in South Africa (SA), as a leading cause of morbidity, disability and premature death in the country. With less than 1% of the world’s population, SA accounts for 15% of the global DR-TB burden. The national burden remains high despite ongoing control efforts. This is partly a result of high treatment interruption and loss to follow-up rates, with 30% of patients who start treatment lost to follow-up, and 67% retention at 18 months. It has been recognised that DR-TB patients’ reasons for interrupting treatment may be personal, social or structural. Personal reasons include patients’ perceptions of treatment, hospital admission or regular clinic attendance as intrusions into their work schedule, and they may, therefore, interrupt treatment for the reason of returning to work.

DR-TB poses a serious risk of transmission in the workplace, especially in high-risk settings such as healthcare and mining. Airborne diseases such as TB can be transmitted to contacts who are often unaware of such risk and may be unable to take precautions to protect themselves. Studies have shown that substantial TB transmission occurs outside of household settings, such as in public transport and shared workspaces.

The question as to whether a health worker has a duty to divulge confidential information to protect the public health interest without the patient’s consent continues to be a subject of debate globally. As a result, there is no clear consensus on non-consensual disclosure, with very little evidence on enabling policies and practices contained in World Health Organization (WHO) recommendations, treatment and ethics guidelines or judicial rulings from notable court cases. The WHO’s guidance on the ethics of TB prevention and control recognises that the health of contacts of people with infectious TB is seriously threatened, especially if they are infected with a drug-resistant strain. It therefore recommends that health workers should balance duties to their patients with an obligation to protect the lives of others at risk, and that non-consensual disclosure should be considered when all reasonable efforts to engage the patient’s co-operation have failed.

In the USA, the duty to disclose was first legally defined in the 1976 landmark Tarasoff ruling by the Supreme Court of California, in which it was held that ‘the privilege [of confidentiality] ends where the public peril begins’. Under this ruling, physicians have the duty to warn or protect third parties from patients with severe medical conditions.
In Canada, the confidentiality of patient information is protected by statute except in certain circumstances, when a physician’s duty to the public outweighs the principle of confidentiality. In such cases, Canadian law permits the physician to provide information about a patient without his or her consent to protect society.\(^\text{[12]}\) In a notable court case (Smith v Jones), the Canadian Supreme Court established three criteria that must be present before non-consensual disclosure can be deemed appropriate: (i) there must be a clear risk to an identifiable third party (person or group of people); (ii) there must be a risk of serious health or bodily harm or death to the third party; and (iii) the danger must be imminent.\(^\text{[13]}\)

Some medical regulatory bodies have also taken clear-cut positions on the issue. The UK General Medical Council in 2004 held that disclosing personal information about a patient without consent may be justified in the public interest if failure to do so may endanger public health safety or expose others to a serious risk of harm.\(^\text{[14]}\) Similarly, the American Psychiatric Association’s guidelines provide that non-consensual disclosure to protect third parties at risk of infection through the behaviour of an HIV-positive patient is ethically permissible if the psychiatrist has exhausted efforts to work with the patient to terminate such behaviour that places others at risk.\(^\text{[15]}\)

The complex question as to whether the constitutional right to freedom and security of the person and/or privacy may be limited where public health interest is paramount has been the subject of a previous case law in SA (Minister of Health for Western Cape v Goliath and Others).\(^\text{[16]}\) In that case, the respondents had all been diagnosed with extensively drug-resistant TB (XDR-TB). Because they were contagious and had failed to comply with the voluntary treatment regimen prescribed for them, the Minister of Health applied for an order compelling the respondents to be detained in a specialist TB hospital to receive treatment. The court considered various factors, including: the respondents’ rights in terms of section 12 of the Constitution,\(^\text{[17]}\) including their rights to freedom of movement; that the respondents were capable of spreading the disease, but had failed to adhere to the voluntary treatment; and the toxicity and associated side-effects of the drugs necessary to treat XDR-TB. Other considerations included the Minister of Health’s duty to prevent and control the spread of communicable diseases in terms of section 7(1)(d) of the National Health Act No. 61 of 2003 (NHA).\(^\text{[18]}\) Following its evaluation of these considerations, the court ruled that the detention and treatment of the respondents, although a breach of their section 12 rights, were both necessary and mandated by section 7(1)(d) of the NHA to protect public health interest. While the judgment is not directly related to the context of non-consensual disclosure of DR-TB status, and did not adequately traverse the requirements outlined in section 36 for limiting human rights, it illustrates the applicable Constitutional principle that public health interest may, in specific circumstances, take preference over an individual’s rights to freedom and security of the person, privacy and bodily integrity.\(^\text{[19]}\)

At the same time, under section 14 of the NHA, healthcare practitioners have the duty to respect patients’ right to autonomy, informed consent and confidentiality, including the right to refuse treatment. However, section 8.2.4.1 of booklet 10 of the guidelines similarly prescribes that, in situations where healthcare practitioners have considered all available means of obtaining consent, but are satisfied that it is not practicable to do so, personal information may be disclosed where the public benefit of disclosure outweighs the patient’s interest in keeping the information confidential (Table 1).

In the context of occupational health and safety, scheduled disclosure of the DR-TB status of workers who are refusing, interrupting or failing DR-TB treatment raises many ethical and human rights concerns. It creates a dilemma between the rights of the individual and the duties of the health system to control infectious diseases for the public good. The relevant regulatory framework incorporating human rights, public health acts and occupational health statutes are summarised in Table 1.

TB is an airborne disease that can be transmitted to contacts who may be unaware of such risk, and unable to take precautions to protect themselves.\(^\text{[20]}\) Effective tracking of treatment defaulters to ensure return to treatment can help reduce the DR-TB burden by 30%, using second-line regimens.\(^\text{[21]}\) There is therefore a need to explore strategies of optimising treatment adherence and reducing infectiousness and transmission.

Disclosure of the DR-TB status of patients refusing or interrupting treatment to their employers or managers without their consent is an intervention that could support infection control measures in workplace settings, protect workers and the public from DR-TB and reduce the stigma associated with this condition. However, such disclosure can bring with it moral, ethical and legal concerns, particularly where patients are reluctant and/or refuse to disclose. There is therefore a need for a standard operating procedure consistent with normative and legal frameworks that balance the public health purpose of such workplace disclosure with its human rights implications. Furthermore, this protocol is necessitated in view of the high burden of HIV/AIDS in SA and the country's workforce, making a substantial proportion of the workforce who are HIV-infected vulnerable to opportunistic TB infection. The protocol should be consistent with the WHO’s recommendation that duty to third parties may justify disclosure of patients’ TB status without their consent, and that public health authorities and TB programmes should develop clear policies, standards and procedures governing such disclosure.\(^\text{[22]}\)

The obligation to protect potential victims of DR-TB transmission in high-risk settings such as workplaces has become an issue that must be dealt with in routine clinical practice and public health systems at large.\(^\text{[23]}\)

The present protocol was therefore developed to provide professional and ethical guidance for health workers in dealing with non-consensual disclosure of the DR-TB status of patients.

**Methodology**

A comprehensive search of the literature was conducted to draw on evidence relating to public health and human rights best practices and normative standards. Inputs were sought from consultative forums with clinicians, occupational health physicians, provincial DR-TB programme managers, municipal environmental health practitioners and human rights lawyers. Additional inputs were also received from provincial DR-TB review committee members.
Table 1. Summary of relevant human rights, public health acts and occupational health statutes (adapted from Adams et al.\cite{19})

| Act/law | Relevant section | Summary of provision | Enforcing agency |
|---------|------------------|----------------------|------------------|
| International Covenant on Economic, Social and Cultural Rights \cite{20} | Right to health (article 12 and general comment 14) | By being party to the covenant, the South African (SA) government assumes obligations and duty to respect, protect and fulfil the right of persons to the enjoyment of the highest attainable standard of physical and mental health. This obligation includes protection from the risk of preventable disease transmission. | Office of the United Nations High Commissioner for Human Rights |
| International Covenant on Civil and Political Rights\cite{21} | Right to privacy (article 17) | No one shall be subjected to arbitrary or unlawful interference with his privacy, family, home or correspondence, nor to unlawful attacks on his honour and reputation. | Office of the United Nations High Commissioner for Human Rights |
| SA Constitution\cite{16} | Right to dignity (s10); right to privacy (s14); right to freedom of trade, occupation and profession (s22); right to an environment that is not harmful to health or wellbeing (s24); conditions under which limitation of rights can be justified (s36) | The state, institutions and individuals have the collective responsibility to respect, protect and fulfil the rights in the Bill of Rights, taking into account the nature of the right and the nature of any duty imposed by the right; rights may be limited under certain condition, and such limitation is justifiable if its purpose is to protect others’ rights, or in the public interest and in the absence of less restrictive means to achieve the purpose. | Department of Justice |
| National Health Act No. 61 of 2003 (NHA) \cite{17} | Right to privacy and confidentiality (s14); health workers’ obligations to notify medical conditions that are deemed notifiable by law (s90) | Healthcare professionals have the duty to protect patients’ rights to confidentiality, and must not disclose any information relating to the patient’s health status without their informed consent, unless a court order or any law (such as s90, which provides for notification) requires that disclosure, or non-disclosure, of the information represents a serious threat to public health. | Department of Health |
| Occupational Health and Safety Act No. 85 of 1993 \cite{22} | Health and safety duties of employers to their employees (s8-10); health and safety duties of employees to their employers and co-workers (s14) | Both employers and employees have obligations to ensure the protection of occupational health and safety. Every employer is required to provide and maintain, as far as is reasonably practicable, a working environment that is safe and without risk to the health and safety of employees, while also requiring employees to take reasonable care for the health and safety of themselves and of other persons who may be affected by their acts or omissions; employers must also do risk assessment and implement medical monitoring of workers who need monitoring. | Department of Labour |
| Basic Conditions of Employment Act No. 75 of 1997 \cite{23} | Sick leave (s22); proof of incapacity (s23) | During every sick leave cycle, an employee is entitled to an amount of paid sick leave. For this, the employee must produce a medical certificate stating that the employee was unable to work for the duration of the absence on account of sickness or injury. | Department of Labour |
| The Labour Relations Act No. 66 of 1995 (as amended in 2014) \cite{24} | Protection from unfair dismissal from work (s191) | An employee who feels unfairly dismissed by the employer may elect to refer the dispute either to arbitration or to the labour court. | Department of Labour |
| Compensation of Occupational Injuries and Diseases Act No. 130 of 1993 \cite{25} | Compensation for occupational diseases (s65) | Provides for medical cover and compensation of occupational injuries or diseases arising from workplace exposures. It also allows for wage replacement if disabled by disease and unable to work, such as for occupationally acquired TB. | Department of Labour |

...continued
**Table 1. (continued) Summary of relevant human rights, public health acts and occupational health statutes (adapted from Adams et al. [19])**

| Act/law                                      | Relevant section                                                                 | Summary of provision                                                                                                                                                                                                 | Enforcing agency                        |
|----------------------------------------------|----------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------|
| Unemployment Insurance Act No. 63 of 2001[20] | Part C (illness benefits) s19 - 23                                              | Provides for the payment of unemployment benefits to employees when they become unemployed and unable to work due to conditions such as illnesses.                                                                          | Department of Labour                     |
| Social Assistance Act No. 13 of 2004[21]     | Eligibility for disability grant (s3)                                            | A person is eligible for a disability grant if he or she is disabled and does not, without good reason, refuse to undergo the necessary medical or other treatment recommended by a medical officer. | SA Social Security Agency                |
| City of Cape Town Environmental Health By-law[22] | Trade and accommodation establishment regulation (s28)                          | The owner of a business or employers must not knowingly cause or permit any person suffering from a communicable disease to be employed in or on the premises unless (s)he is in possession of a medical certificate to the effect that such person is fit to continue his or her employment. | City of Cape Town Environmental Health Services |
| Guidelines for Good Practice in the Health Care Professions[23] | Ethical considerations in seeking informed consent (booklet 9); Protecting patient’s confidentiality (booklet 10) | Healthcare practitioners must respect the patient’s right to autonomy, informed consent and confidentiality, including the right to refuse treatment. However, practitioners also have certain legal obligations as stipulated by the NHA. Section 8.2.4.1 of booklet 10 describes cases in which healthcare practitioners have considered all the available means of obtaining consent, but are satisfied that it is not practicable to do so, personal information may be disclosed in the public interest where the benefits to an individual or to society of the disclosure outweigh the public and the patient’s interest in keeping the information confidential. | Health Professions Council of SA         |

The following pertinent questions and key considerations guided the protocol development. Under what circumstances can non-consensual disclosure be justifiable? How much of a serious public health threat is DR-TB in the workplace? Which of the rights enshrined in the SA Bill of Rights, Constitution and NHA can pose legal dilemmas to non-consensual disclosure? Which of those rights can be limited within reasonable and justifiable considerations? How does the protocol navigate the implementation challenges in the informal and private sectors, where occupational health and safety considerations are often limited or absent? Are there extant provincial occupational and environmental health laws that can lend legal support to the protocol? Are there existing professional statutes, rules and regulations that can provide some measure of normative standards? Are there national and global best practices that can be leveraged? Other questions that arose were: what social security benefits or labour law statutes can be explored to encourage patients’ early return to treatment? What collaborative opportunities can be explored, such as amendment of the Road Traffic Act to ensure that professional drivers are DR-TB free before licensure?

This protocol was structured in line with the End TB strategy of the WHO, and is guided by the following principles:

(i) The non-consensual disclosure of the patient’s health status to a third party interferes with their rights to privacy and confidentiality, both of which are a cornerstone of the health-care worker-patient relationship.

(ii) Third parties such as workplace contacts also have rights, which government has the obligation to protect for the public health good.

(iii) Patients’ right to confidentiality may be limited, but only if such limitation is done either in line with section 36 of the SA Constitution, or Siracusa Principles.[31]

(iv) Recognising that patients are part of their larger communities enables the identification of public health risks and approaches to mitigating such risks.

It lays down normative and legal support, drawn from a broad range of global, national and province-level evidence, for dealing with the ethical complexities of non-consensual disclosure in occupational settings. This protocol proposes an integrated, stepwise incident management and initial risk assessment algorithm adapted from models proposed by Appelbaum[34] and Chaimowitz.[32]

**Results**

**Incident management process**

An incident of DR-TB treatment outcome may comprise one of the following scenarios: (i) treatment interruption; (ii) treatment failure; (iii) on treatment but still infectious; or (iv) refusal to initiate treatment. The key steps in the incident management process are highlighted below and schematically described in flow charts (Figs 1 and 2).
(i) Following the establishment of an incident, the treating physician should notify the health facility’s DR-TB professional nurse, or a similarly designated officer.

(ii) Upon notification, the DR-TB professional nurse, or other designated officer, should, within 3 working days, arrange for a clinic counselling session (in the case of patients who are currently on treatment but remain infectious), or designate social workers to conduct a home visit (in the case of treatment interruption or those refusing to initiate treatment after the DR-TB diagnosis).

(iii) There is a need for careful assessment of each incident, on a case-by-case basis, of current employment status and whether the patient has already disclosed their DR-TB status at the workplace. This is necessary to avoid unwarranted disclosure (e.g. in situations where patients are either currently going to work nor employed, in which case domestic and occupational infection control strategies can be instituted with adequate communication and counselling of patients and household members).

(iv) Any successful contact of patients or their family members/next of kin, either telephonically and/or through a home visit, is a vital opportunity to ascertain their current employment status.

(v) If, after the third home visit, the patient remains untraceable and it is obvious that family members and home contacts do not know his or her whereabouts, the patient should be considered as lost to follow-up.

(vi) However, if at any home visit attempt the patient is untraceable and it is obvious that he or she is deliberately refusing to see or evading the social workers, such home visit attempt should be deemed unsuccessful, as per the incident management algorithm.

(vii) To encourage early return to treatment and care, patients should be made aware of the possibility of accessing social security benefits (a disability grant or unemployment insurance funds, or both), where patients are eligible. These benefits will also lessen the economic implication of loss of income due to sick leave taken on medical grounds, until they achieve at least one negative sputum culture result.

(viii) To encourage disclosure by self in patients still on treatment but who pose a transmission risk, efforts should be made to secure access to social security benefits to allow patients sick leave. Where a patient continues to work after disclosure, there is a need to ensure appropriate infection prevention and control measures at the workplace.

(ix) Documentation of all clinic and home visit counselling and persuasion attempts, with their outcomes, in a tracking log book is recommended.

(x) For patients who are willing to return to treatment and care after a home visit, non-consensual disclosure is usually neither necessary nor appropriate. As far as possible, the treating clinician should grant patients sick leave until they achieve a first negative sputum culture while on resumed treatment.

(xi) The processes of application for social benefits such as a disability grant and/or UIF benefits require medical justification, which should be supported and motivated for by the treating physician.

(xii) For employees in private sector employment, the possibility of accessing temporary incapacity benefits from private insurance

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**Fig. 1. Incident management for patients interrupting treatment and still going to work.**
benefits can be explored. Contracted staff and those without private health insurance benefits should be assisted to access benefits via the Department of Labour’s UIF.

Non-consensual disclosure
Non-consensual disclosure becomes inevitable with persistent unwillingness to return to treatment despite sufficient attempts to get the patient back on treatment. Non-consensual disclosure should be done as respectfully and discreetly as possible by members of the healthcare team, and only information that is vital to protect the public from harm should be disclosed. Disclosure should be made only to those who need to know. Such persons may include employers, human resource managers, occupational health officers or other designated staff responsible for employee health and safety management. Patients should be provided with advance notice of the date and schedule of disclosure. It should also be made in a manner that protects patients from potential violence and discrimination afterwards.

Generally, non-consensual disclosure must be done as an option of last resort, and even when indicated, strict adherence to administratively just procedures must be required.

The justifiable goals of disclosure include the following:
(i) Enable the contact screening of co-workers and other third parties who have had substantial risk of exposure to the patient.
(ii) Institute prophylaxis and preventive therapy, where appropriate.
(iii) Initiate prompt treatment for those who test positive on screening.
(iv) Enable the institution of occupational health and safety measures.

A stepwise approach to implementing non-consensual disclosure is outlined in Table 2.

Situations where disclosure can be deemed inevitable
Disclosure should be considered inevitable in the following situations:
(i) Interrupted treatment, and currently going to work but unwilling to return to care after exhausting all persuasive steps of the incident management algorithm (at least three home visits or clinic persuasive attempts).
(ii) Refusing to initiate treatment, and currently going to work but still unwilling to initiate treatment after exhausting all persuasive steps of the incident management algorithm (at least three home visits or clinic persuasive attempts).
(iii) Failed treatment, currently going to work but unwilling to self-disclose DR-TB status at work after exhausting all persuasive steps of the incident management algorithm.
(iv) On treatment but still infectious, and currently going to work but unwilling to self-disclose DR-TB status at work after exhausting all persuasive steps of the incident management algorithm.

Resources required
Due consideration must be given to the provision of necessary resources and capacity to implement this protocol (Table 3), and a framework for monitoring and evaluation is essential (Table 4).
(i) Human resources: physicians, DR-TB professional nurses, social workers, environmental health officers, TB counsellors, community health workers and health records personnel
(ii) Diagnostics: drug susceptibility testing tracking and results retrieval, National Health Laboratory Services database linkage and prompt response
(iii) Patient-tracking resources, monthly updated patient contact details, monthly occupational-status tracking register and incident management log book
(iv) Channels of collaboration between the Department of Health with other relevant authorities and stakeholders (Table 3).
**Table 2. Stepwise approach to implementing non-consensual disclosure**

| Step | Action |
|------|--------|
| Step 1 | DR-TB professional nurse or any other designated officer in that capacity nominates disclosure team. The team should include a clinician, an EHO and a social worker. |
| Step 2 | Disclosure team contacts workplace, identifies employer, manager or any other focal person, and proposes visit for occupational purpose (patient's identity needs not be revealed at this stage). |
| Step 3 | Make visit as scheduled. Ascertain that patient is currently going to work. Once ascertained, the EHO discusses with workplace contact person the purpose of visit. Assess the workplace for any existing occupational health policy, number of staff, infection control measures and risks of occupational exposure and TB transmission at patient's occupational setting. |
| Step 4 | Assess employer or manager's understanding of the purpose and goals of disclosure. |
| Step 5 | Make disclosure. |
| Step 6 | Provide education to adequately provide relevant information to allay undue fear, apprehension and concerns, while mitigating risks of stigma, discrimination and possible assault on patient. |
| Step 7 | Discuss with employer and explore possibilities of social security and unemployment benefits for patient while booked off work following disclosure. |
| Step 8 | Screening of workplace contacts, if indicated. |
| Step 9 | Monthly visits by the EHO, in exceptional cases where EHO deems transmission risk minimal enough to allow patient to continue to work post disclosure, to monitor the impact of disclosure on patient and to ensure compliance with workplace infection control measures. |

**Table 3. Resources required**

| Agency | Purpose |
|--------|---------|
| South African Social Security Agency | Assist with access to social security opportunities for patients booked off work on medical grounds |
| Municipal and local government health services authorities | Establish contact with workplaces and evaluate risk |
| Department of Health legal department/committee | Advise clinician on legally complex cases |
| Transport authorities | Deal with cases in which taxi drivers pose DR-TB transmission risks to passengers |
| Department of Labour | Ensure compliance with labour legislation, and protection for medically impaired employee |

**Table 4. Monitoring and record-keeping framework**

| Record-keeping | Frequency | Personnel responsible | Tools required |
|----------------|-----------|-----------------------|---------------|
| Documentation of all attempts and outcomes of phone call, home visit and clinic counselling/persuasion activities | Following each activity | DR-TB professional nurse, or any other designated officer in that capacity | Incident management log book |
| Post-disclosure monitoring of impact of disclosure on patients who continue to work for identification of stigma, discrimination and workplace adherence to infection control measure | Monthly | EHO | Post-disclosure monitoring register |
| Tracking and validating current employment status of patients on treatment | Monthly | Social worker | Employment status tracking register |

**Procedure for seeking redress on the grounds of human rights abuses of this protocol**

Consideration of human rights implications is a core element of public health policies and laws. Healthcare providers, as custodians of this protocol, therefore have the responsibility to ensure that they act in accordance with professional ethics and humaneness in the discharge of their duties, while ensuring minimal risk to the fundamental rights and freedoms of the patient. Like any other public health intervention, inadvertent violation of human rights can result from implementing this protocol. Should any undue violation of rights occur from the implementation of this protocol, however, this protocol includes the following procedure for channelling of complaints: it is proposed that the avenue for laying complaints and seeking redress be through existing health committees, or community tribunals, which will independently investigate reported cases and mediate in collaboration with the legal advisory team, including human rights lawyers.

**Discussion**

This protocol provides professional, legal and ethical guidance for health workers in dealing with the dilemma of non-consensual
disclosure to limit the spread of DR-TB in occupational settings. The analysis of global and local bioethical and legal standards highlights the enormous challenge of balancing public health and human rights. We acknowledge that forced public health interventions such as non-consensual disclosure of DR-TB status, for the most part, are not desirable, but may be necessary in circumstances where such interventions are clearly in the public health interest and where alternatives are not feasible or effective. Nevertheless, it is possible to achieve this balance by incorporating human rights principles into policies that limit individual rights in the interests of the public good, and still meet ethical standards.

Foreseeable implementation challenges
(i) Challenges stem from the fact that a substantial proportion of DR-TB patients work in the informal sector, which means they are often unregistered and hard to reach. These include self-employed persons in informal occupational settings, taxi drivers, owners of outlets such as shebeens, baby daycare centres and hair salons.

(ii) Operationalisation can be limited by unavailable or incorrect phone and/or physical address contact details of patients.

(iii) An overburdened health workforce in short-staffed health facilities may also pose a challenge.

Additional recommendations
(i) To aid future tracing of patients, there is a need for proper documentation and tracking (at each clinic visit) of current patients’ contact details, including phone and residential addresses.

(ii) To enable the determination of probable current employment status at any point in time while on treatment, there is a need for patients’ employment status to be monitored monthly during clinic visits, and validated through a contactable next-of-kin.

(iii) Application of municipal by-laws and involvement of specialised health services for environmental health assessment of transmission risk at informal settings such as unregistered creches, hair salons and daycare facilities is needed.

(iv) There is a need for intensified counselling and continuous education of patients on the transmission risks of DR-TB.

(v) The National Transport Act should be amended to make the issue of patients’ sex partners without consent? Stud Family Plan 2007;38(4):297-306.

Conclusion
Addressing the serious problem of DR-TB in the occupational setting and non-disclosure by infectious TB patients may create an ethical dilemma of dual loyalty for health workers. While the limitation of an individual’s rights may be a necessity to protect the public interest, including the prevention and control of communicable diseases, it is important that such a limitation is not undertaken in an arbitrary manner, or without regulation. It must be done in terms of a policy that is consistent with national and international human rights law governing the limitation of rights in the public interest. The present protocol aims to ameliorate that challenge by providing guidance that is based on a consideration of professional, ethical and legally sound principles that support non-consensual disclosure, and balance the public health goals of the intervention with its human rights implications.

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