

*** Supervision and monitoring of the intervention clinics**

Regular weekly visits were made to every intervention clinic. The research nurse examined every tuberculosis patient clinic card for details of sputum monitoring, supporter attendances (these were copied onto clinic cards from the patients' cards), and clinic attendances with clinical details documented at each visit. Drug supplies were inspected, and patient management issues and problems were discussed with staff. The author visited each clinic every three months for the same purpose.

*** Follow-up of non-attending patients**

Clinic nurses were trained to identify any patient who missed a monthly clinic appointment within one or two days. Strategies for follow-up that involved clinic staff action were encouraged. Although emphasis was laid on the clinic staff taking responsibility for the follow-up, their lack of transport and their preoccupation with the daily load of sick patients at the clinic were limiting factors. It became clear that the back-up of researchers, was needed. Where necessary the research nurse went to patients' homes and to their supporters to discuss the problems.

*** Interest and cooperation of clinic nurses**

Most clinic nurses were interested in the new decentralised tuberculosis service, and keen to help their patients. They completed the cards carefully, and some made special efforts in tracing non-attenders. They were receptive to advice and support, and clearly pleased to have regular supportive and instructional visits, and to have colleagues with whom they could discuss their work. The full day training sessions held at a nearby hotel, away from the work situation, were especially popular.

*** Cooperation of hospital and community tuberculosis staff**

The key to the good treatment outcomes throughout the district, not confined to "treatment" clinics, was the cooperation and enthusiasm that developed among the senior nurses in the hospital tuberculosis ward and among the community tuberculosis team consisting of a nurse and two health educators. They all attended the training sessions, and set up regular meetings (monthly and more often as necessary) with the research team to discuss the control programme and issues arising. The ward sister fulfilled an important role in informing the researchers about discharged patients. Help was provided by the researchers in describing and reorganising the roles of the community tuberculosis staff, so that they performed more effectively and efficiently.

*** Support of laboratory staff**

The laboratory staff had not previously been informed or consulted about tuberculosis work. Once they were invited to training sessions, and quality control systems were set up through the research programme (with their cooperation), the turn-around time for specimens improved. Measuring quality of laboratory work did not form part of the study, but a quality control system set up by the South African Institute of Medical Research as a result of the research, had results which were said to be excellent.

*** Availability of drugs**

Anti-tuberculosis drugs were erratic in supply before the research project. Combinations drugs were unused and unknown. The pharmacist, with the help of the newly established provincial drug supply system, eventually responded to suggestions of the researchers and cooperated to ensure that all recommended drugs were available. Issues of supplies for up

to five months were given to the research nurse to take to "treatment" clinics to be stored for patients who attended there.

*** Interest and willingness of community supporters**

The social system in this district whereby many community members took responsibility for one another was an asset to the tuberculosis supporter system. With appropriate initiation from research and service health workers, a variety of lay-people became supporters, and successfully trod the path to cure or completion of treatment with many patients.

20.2.2 What were the constraints in the process?

There were a number of factors that constrained the process of decentralising tuberculosis services to district clinics. These could be classified as factors in the physical and social environment of this area, health services deficiencies, and research deficiencies and limitations.

*** Physical and social environmental factors**

Major constraints for the provision of all health care in this district are the long distances from where people live to the hospital and to clinics along bad, dangerous roads, and where public transport is expensive.

The researchers became aware of the social and cultural beliefs that were barriers to early health service attendance, both because the disease was stigmatised through local beliefs that the cause was unacceptable behaviour, and because traditional healers had to be consulted. Until different health care providers cooperate, patients will be torn, and will

present to clinics and hospitals with late stage tuberculosis.

* **District management deficiencies**

In the absence of a district health system, management was actually that of the hospital. The district suffered from a number of management deficiencies, many consequent on the previous system of "homelands", two of which, Lebowa and Gazankulu, had formed part of the previous area of the district. Any initiatives that communities or health workers might have shown had been stifled by this system, so that demotivated staff and absent resources were the norm. Exceptions existed of course, at management and implementation levels.

The newly established provincial government could not be expected to facilitate immediate changes in attitudes or behaviours. Health service management was hospital-based, and there was poor understanding and limited activity around community services, including clinic management. Resources and support for clinics were minimal. As with many government services, senior staff spent long hours travelling to and attending meetings called at short notice at head office, with resultant disruption of routines, tiredness and frustration.

The research initiative met opposition from such managers. This was not overt, but manifest in lack of active support for the new decentralised system, failure to ensure accountability of clinic staff for their actions in tuberculosis patient management and failure to be seen supporting the work, resulting in confused lines of authority for clinic staff. Organising training sessions (at no cost to the hospital) was difficult as managers would not commit themselves to allowing staff to attend, and failed to help with forward planning. Much time and energy was expended in diplomatic negotiations, and in trying

to understand the underlying issues.

The old Lebowa clinics, under Mapulaneng hospital, posed particular problems. Drug and vaccine supplies were irregular, empty shelves were the norm, staff felt and were incompetent as they had to send patients away. The causes were multiple, including poor ordering, theft at various levels en route to clinics, and "lack of money". Water supplies for clinics were inadequate, indeed frequently non-existent. Accommodation facilities for clinic staff were below standard.

*** Perceptions of researchers by district health workers**

Although it was not stated, the behaviours of managers and clinic staff, and experiences of academics working in the related University project, led the researchers to believe that they were perceived by some as temporary sojourners in the district working for academic recognition. Noone was impolite enough to state this, but it must have been part of the apparent demotivation that constrained full involvement. One had sympathy for health workers in difficult and constantly challenging situations, whose standards were being questioned, and who were being asked to do more in quantity and quality, without apparent personal satisfaction. This applied equally to senior district staff.

*** Researchers as service providers**

The researchers assumed the role of tuberculosis service providers for "treatment" clinics. This, although discussed and agreed upon before the project started, inevitably caused some problems. Staff of the "treatment" clinics became accountable to the researchers, without the visible support of the hospital community nursing managers. The researchers had tried, without success, to organise management presence at an introductory meeting

at each of the clinics involved.

Another issue was that tuberculosis services became "verticalised", with a specific team discussing only that disease and its management. This could have lost credibility for the comprehensive approach, and may have alienated the researchers from clinic staff who had a number of responsibilities apart from management tuberculosis.

*** Absence of districts and a functional district health system**

Productivity depends on clear work structures and lines of accountability. During the period of the study, the district system did not exist. In fact the three proposed districts of the Bushbuckridge sub-region were still non-functional in the first quarter of 1998! The hoped for changes in boundary lines of responsibility, described since 1994, did not occur, causing confusion and frustration of health staff, and contradicting the training on the district system that the researchers and others carried out. The delays were partly due to the need to ensure democratic decision-making by communities unused to involvement in local government. As a result of the non-district system, there was further demotivation of the clinic nurses, for whom this would have the greatest impact.

The (old) Mhala area clinics, many kilometers distant from Tintswalo hospital, remained the responsibility of that hospital's community health staff. They had neither staff capacity, time, nor vehicles to service these far areas, while assuming an increased load of work with responsibility for areas closer to the hospital (previously under Mapulaneng hospital). They had gained work load, but lost none. In fact it was surprising that the hospital-based community team managed to improve tuberculosis management, as they did. Referrals of patients who lived outside the proposed district was very difficult, as authorities responsible for other areas were far away, there was no means of

communication (senior staff were not even allowed to telephone other authorities from the hospital ward to enact referrals), and these other authorities were not clearly defined.

*** The limited role of clinics in tuberculosis patient management**

During the period of the study, any patient who attended a clinic, and in whom tuberculosis was suspected, was asked to go to Tintswalo hospital for diagnosis, and initiation of treatment, usually as an in-patient. Only on discharge was the patient referred back to the clinic for continuation of the course of treatment, to be referred back to the hospital again for final assessment and discharge from treatment. The clinic staff saw the patients only three, four or sometimes five times at monthly visits, with roles limited to clinical assessment, checking of supporter attendances (and copying the supporter's ticks from the patient card to the clinic card), and making the next appointment. Clinic staff had no contact with supporters. Without the research team, who did random visits to patients and supporters, non-attendances of patients at their daily supporter visits would not have been identified until the next monthly clinic visit. Patients who failed to attend for their monthly appointments would be identified within a few days of the failure by clinic staff, but in the absence of time and transport to trace those patients at home, clinic staff would be powerless to act, and would need to contact the district tuberculosis team for them to seek out such patients. Contact was dependent upon reliable communication systems from the clinics, the ability of the district team to access transport, and upon sufficient district staff assigned to tuberculosis work. Frequently all three of these factors were absent. The role of clinic staff was thus limited, and frustrating for professionals trained in a comprehensive patient approach.

*** Failure to monitor sputum bacteriology during and at the end of treatment**

Sputum tests during and at the end of treatment (2 and 5-6 months for new patients) were part of the research (and the national) policy, but were not done, despite the availability of the research nurse to transport the specimens. This was a combined deficiency of the researchers who failed to insist, and to remind clinic nurses about the timely collection of specimens, and of the clinic nurses who did not accept responsibility for doing this. There was no help from management who should have insisted on the implementation of this policy.

*** No organised laboratory network**

Tuberculosis services, as with many other clinic-based services, are dependent on a rapid and reliable laboratory link. The absence of an organised system of delivery of sputum specimens to the district laboratory, and the return of results within 48 hours, has been noted as a major constraint in many parts of the world.²⁵ The absent district system, long distances, bad roads, the tuberculosis control programme with emphasis on bacteriological diagnosis only just emerging, and inadequate organisational and management skills meant that there was no network.

For "treatment" clinics, the researchers themselves provided the transport for specimens to the laboratory and results back, but during the last six months of the study, made strenuous attempts to get the service to organise the link. Suggestions were to utilise senior community nurses on supervisory visits, the hospital pharmacy team delivering drugs to clinics, ambulances transporting patients, or even local taxis. A system was never set up, since it was beyond the capacity of the researchers to establish this. The Tintswalo hospital laboratory was not part of the SAIMR network, so the improvements

taking place in other provinces, notably Gauteng and Mpumalanga (personal communications Drs. L Floyd and E Balt respectively), were not a feature of this area.

*** Inadequate organisation and management of the community supporter system**

The organisation of the supporter system was well managed at an individual level by the research nurse or the hospital community team. They selected an person acceptable to the patient, informed that person what was expected, and made occasional visits to monitor the process of daily attendance. However, the process was unsystematic, and without community consultation. The considerable resources within the community could be better accessed and utilized if volunteers within villages were formally recruited and trained, so that they formed a team, ready for the task of supporting tuberculosis patients as needed. They could play a part in general health promotion, including the Extended Programme of Immunisation, nutrition, home-based management of diarrhoeal disease, family spacing, tuberculosis and other disease case-finding, and environmental control.

The system whereby patients attended clinics to fetch their tablets, and took these to their supporters, was illogical, and must have seemed so to patients. It would have been better for drugs to be taken to supporters, or for them to collect these at the clinic. These aspects needed further discussion and trials to formulate a district supporter plan.

A further important issue was the lack of any incentive for supporters. This would not necessarily be monetary, but could take the form of certificates presented at an annual function, transport allowances, or badges to denote membership of supporter groups. The strength of the community "care group" movement that contributed significantly to the control of trachoma in northern South Africa, was the authority and status given to members.⁴⁰⁵

*** Poor provincial government commitment**

District management and commitment, even had these been present and adequate, would have been very difficult without provincial government commitment to a national tuberculosis policy, and appropriate allocation of resources, including provincial management and training staff. Only towards the end of the project in mid-1996, were provincial tuberculosis co-ordinator/manager and trainer posts filled.

*** Non-involvement of hospital medical staff**

Although tuberculosis case management, community control and the research project were presented and discussed at at least three formal doctors' meetings at Tintswalo hospital, doctors did not award the disease priority status, and remained distant from the community-based system. Patients were often started on treatment without microscopy confirmation of diagnoses, sometimes with incorrect drugs, and occasionally discharged without notification or referral to the tuberculosis ward for notification and registration. Both clinical diagnosis and death certification (as discussed in chapter 17) sometimes incorrectly included tuberculosis when it was unproven or unlikely. It may well have been missed occasionally when it was present.

20.2.3 How can the improvements be sustained and further improvement achieved?

An operational model for the analysis of tuberculosis control programmes has been described.⁴⁰⁶ The model states 8 steps: motivation (of patients to present to health facilities), selection (of suspect cases for smear microscopy), examination (of smears in the laboratory), sensitivity (of smears), prescription (of correct treatment), treatment

(patient obtains correct treatment), regularity (adherence) and effectiveness (probability of cure).

This model for good tuberculosis control and management requires efficient and effective actions of health workers at a number of levels. District, provincial and national management commitment and support means training of staff, and the provision of the infrastructure required for diagnosis and treatment, as well as those resources required for efficient follow-up of non-adherent patients, and for routine supervision of clinic staff.

The roles of district health workers need clear definition. An effective tuberculosis programme presupposes the existence of a functioning district health system, the acceptance of the priority status of tuberculosis at all levels of the health service, good district service management, which includes well managed and well motivated health personnel. The suggested roles of different levels of health staff are described hereunder.

The district clinic nurse

A clinic nurse has many duties in the provision of primary care to the people served by a district clinic. Regarding tuberculosis, which is best integrated within these duties, the nurse must be able to:

- + decide which patients have suspected tuberculosis
- + confirm the diagnosis, using the laboratory network
- + treat patients with confirmed disease
- + organise clinic drug supplies
- + refer patients with clinical problems or those who require hospitalisation
- + contact the appropriate person (district co-ordinator or hospital clinicians) for

clinic-based management problems

- + record all processes and outcomes on the relevant tuberculosis programme forms
- + follow-up patients who have adherence problems during treatment, or call on the district tuberculosis coordinator to help with this

The clinic should be organised according to the needs and comfort of the patients and the staff. It is well described that patients respond well to a relationship with the care provider. It is thus best for the same nurse to see the same tuberculosis patients for the duration of their treatment, as far as can be organised. It may be useful to have a tuberculosis patient day once a week, or alternatively to structure visits any day of the week.

For a district with a population of up to 250 000 people, with rates of tuberculosis of 200-300 per 100 000 population (double that measured over the last 5 years in the Tintswalo district, and allowing for the rise with the HIV epidemic), there would be 500-750 patients per year, or 250-375 at any one time. The Tintswalo study showed that 28% of patients lived close to the hospital, leaving 72% who would require treatment at one of the 12 district clinics. Assuming equal distribution of patients attending the clinics, between 15 and 20 could be on treatment at each. At clinics in more densely populated villages, this number could be 30. If each attended monthly on one specific day, there would be about five to eight patients per clinic day. This number would seem manageable, even in the context of clinics delivering comprehensive services. If the rise in incidence is more than double that in the early 1990s, and if case-finding rates improve, numbers will be higher, which is a good reason to ensure an efficient system now.

The district communicable disease coordinator

This district-based health worker should have responsibility for all communicable diseases in a district, including malaria, meningitis, typhoid fever, diarrhoeal disease, haemorrhagic fevers, tuberculosis and any others. The job includes a knowledge of the district population, its distribution, health problems, and the health services. The role of this person is to:

- + train health staff about tuberculosis and its management
- + train tuberculosis patient supporters and organise appropriate incentives
- + visit all clinics in a district on a regular basis to teach and to supervise clinic staff
- + respond to calls for help in tracing non-attending patients
- + monitor laboratory and drug supply systems, and report problems to district management

Once the tuberculosis control programme was functioning well, with all clinic staff trained, the routine visits of one or sometimes two clinics per day could be completed in a few hours, so that all district clinics would be visited once in two weeks. This would leave time for training, and for tracing non-attenders at their homes.

Given the responsibilities for other communicable diseases, and outbreak investigations, the task may be too much for one person. An additional district staff member, at the level of a health educator, or nurse should assist. Both would require transport, always a limiting factor in the past. However, an efficient programme with cure rates exceeding 80%, would make the use of adequate staff and transport resources cost effective, particularly if other, less common communicable diseases and good surveillance and outbreak response systems are simultaneously managed in the district.

District management:

District management has to take responsibility for the provision of the infrastructure, upon which a tuberculosis control programme depends. Clinic staff and the district communicable disease coordinator need to be consulted and to report problems, but district management should plan and organise the systems for:

- * Laboratory network
- * Drug supplies
- * Vehicles
- * Stationery
- * Staff management

An efficient and reliable laboratory network between clinics and district laboratory is essential. In many urban areas, a daily collection service from clinics has been arranged, with results faxed back within 24 hours. (Gauteng province, personal experience) In rural districts such a service may be less easy to implement. For those new patients not sick enough to be referred for admission to hospital, and who are to be diagnosed at clinics, bacteriological results should be available within one or two days, so that treatment can start as soon as possible. This requires transport of specimens for which innovative ways of reaching the laboratory must be devised. They may include using health workers on supervisory visits, delivering drug supplies, or fetching patients in ambulances. It may necessitate specific communication with the district tuberculosis team, or the use of local taxis. Whatever the means, it has to be reliable. There is no reason why fax machines could not be installed and functional in every clinic for the results. The costs of organising a laboratory network would be balanced by savings in the short and long terms.

Another district management responsibility is the regular and timeous delivery of drugs for tuberculosis, as well as essential drugs for primary care use, to every clinic. Tuberculosis drugs were not available on shelves of "treatment" clinics in this study, for reasons described by hospital management as related to rules about dispensing drugs. These were not logical to researchers, and should be overcome without difficulty. The supply for each patient attending "treatment" clinics had to be hand-delivered by the research team.

Vehicles for the district communicable disease coordinator and assistant must be available for routine visits to clinics and supporters, for collecting specimens if required, and for tracing non-attending patients. The problem with sharing vehicles with other services is non-availability in times of need. Again a functioning control programme will cost far less than one which does not promote cure.

The provision and delivery of all stationery required for the tuberculosis control programme is the responsibility of district management. Without attention to this seemingly small detail, the ability to monitor and evaluate the programme is lost, and staff become demotivated and confused.

The descriptions of roles and responsibilities above assumes a comprehensive role for clinic staff to manage tuberculosis, different from their limited role during this study. The increased responsibility for correctly diagnosing and treating tuberculosis, and the ability to evaluate progress of each patient should provide greater incentives to clinic nurses. Given the public health impact of this disease, the responsibility is large, but

should not be overwhelming for professionals required to manage a number of life threatening, as well as routine problems which include dehydrated infants, complicated birth deliveries, ordering of drugs and equipment, and responding to community needs and demands.

Adaptations would have to be made for those patients requiring hospitalisation for clinical or risk of interruption reasons, and for those on re-treatment regimens, for whom daily streptomycin injections are needed, either in hospital or from district clinics.

The logistics and the infrastructure described above require good planning, organisation and budgeting by district management.

Staff motivation and factors that demotivate are complex, although in many rural primary health care environments, problems are not difficult to identify. Herzberg's motivation-hygiene theory, based on a study of 200 professionals, proposes that motivators (achievement, recognition, work, responsibility and advancement) lead to satisfaction at work.⁴⁰⁷ For maximum satisfaction, and thus productivity, from clinic nurses, attention needs to be paid to their "job-enrichment" factors. They certainly have work and responsibility, but little recognition and sense of achievement because of the remoteness of their posts, and the lack of management presence to acknowledge the challenges and difficulties of their jobs. Herzberg's "hygiene" factors, which he found to be most frequently quoted causes of dissatisfaction, are those related to the context of work. Named problems were with company policy and administration (management), technical supervision, salary and relationships with supervisor. Researchers studying nurses working at Agincourt Health Centre, one of the health facilities in Bushbuckridge, under Tintswalo

hospital management, found that the absence of close supervision and effective management impacted negatively on their morale, exacerbating feelings of isolation.⁴⁰⁸

The work context of clinic nurses does not always meet the needs of Maslow's hierarchy theory.⁴⁰⁹ Some basic physiological needs may be covered, but acceptable accommodation, and adequate water and sanitation are often absent. Many clinic buildings in the Tintswalo district were in a poor state, certainly not conducive to high standards of productivity. Safety and security are often not assured, nor, at the next level of need, is a need to belong to a group catered for. The isolation and loneliness of professional life in far distant clinics, away from relatives and friends, without means of transport and often without any form of communication, can be profound. The higher needs of Maslow include the motivational factors of Herzberg already mentioned - status recognition and self-fulfilment.

The World Health Organisation have stated that "if health personnel are to provide a caring service, it is necessary for the public health services to show that it too cares".⁴¹⁰ Much is lacking in both the physical and social environment, and in the professional management of nurses assigned to work in rural clinics of Bushbuckridge, and in many other areas, both rural and urban.

20.2.4 How can the community and patient perspectives be incorporated into planning and implementing tuberculosis services?

This study identified some consistent beliefs about the origin of tuberculosis among local communities. Health workers confined to facilities, especially if they do not come from the community they serve, are in danger of missing important information that impacts on the management of diseases. They all have a responsibility to consult community groups

and leaders, and traditional healers, and to work with and through local beliefs and customs in order to provide an acceptable effective service. Traditional healers should be educated about the medical treatment of tuberculosis, and persuaded to refer their clients to clinics or hospitals for concurrent medical management. Continuing attempts to destigmatise the disease through community-based education would facilitate early patient presentation and treatment adherence.

CHAPTER 21

RECOMMENDATIONS FOR GOOD, SUSTAINED TUBERCULOSIS CONTROL

The findings of this study prompt a number of recommendations. In keeping with the principles of integrating all primary care activities, some are general health service recommendations, others relate specifically to tuberculosis. Many would be generalisable to other districts in South Africa, and perhaps to other developing countries.

GENERAL HEALTH SERVICE RECOMMENDATIONS

Commitment to tuberculosis control as a priority for health services

A functioning district health system

Incentives and support for health workers

Task definitions and protocols for management of common health problems

Communication systems

Supervisory visits

Management of material resources - drugs, equipment, water and buildings

The need for good management is a recurring theme in this thesis.

- * A prerequisite in the context of tuberculosis is commitment by management to the priority status of this disease, recognition that the incidence will increase, that the disease is preventable by cost-effective means, and that tuberculosis care is best integrated within a district primary care network.

- * The creation of a functional district health system is fundamental to all health care

delivery.

Staff and material resources need good management.

- * Staff management starts with recruitment of appropriately trained and experienced health personnel. Much has been debated around issues of incentives for doctors who work in remote areas, but little attention has been given to salary supplementation, opportunities for promotion, extra leave and other forms of recognition for the onerous duties that nurses perform in rural primary care settings. Staff rotations should be considered, balancing the desire to give clinic nurses a break from work in remote areas with the need for some permanency and continuity of care. It certainly seems disadvantageous to leave a nurse at the same clinic for years on end.
- * Tasks to be performed should be clearly defined, with protocols, training and retraining on the management of common important health problems (including tuberculosis). This aspect is important too for staff at other levels in the district, including district communicable disease coordinators and district management. Protocols on how to manage common health problems and their complications must be available, together with resources for implementation. Essential components are equipment, telephones, fax machines, and ambulances as well as available consultation services with technical guidance.
- * There must be open communication channels between all clinics, the district communicable disease coordinator, the district headquarters and the hospital, to use routinely for common health problems and their complications.
- * Regular supervisory and monitoring visits by district managers, notably the communicable disease coordinator and the community health nursing manager, are

essential to motivate staff, to address their problems, and to check that work is being performed according to set standards.

- * Clinics cannot function without drugs, vaccines, equipment, constant water supplies and buildings in reasonable repair. Functioning systems for procurement and distribution of drugs and equipment have to be in place, with delegated responsibilities for their maintenance.

SPECIFIC TUBERCULOSIS SERVICE RECOMMENDATIONS

Train all health workers in tuberculosis management

Decentralise tuberculosis care to district clinics

Organise laboratory network for diagnosis and monitoring

Organise community supporters

Provide district support for community outreach functions

Keep standardised records at clinic level, to be collated at district level

Transfer patients efficiently from one district to another

Within the context provided by good general management, there are specific recommendations for tuberculosis management emanating from this study.

- * All health workers, including nurses, doctors, managers, laboratory workers and pharmacists, must be trained in the management of tuberculosis. Education at tertiary and other levels for all health workers must include management of patients with tuberculosis, as well as the principles of the national and international control programmes. Continuing education with on-site training is an essential component to maintain standards and interest. Manuals must be produced, and

courses coordinated and standardised at the national tuberculosis directorate, with extra resources supplied as necessary to provinces unable to supply the needs.

- * Tuberculosis can be and should be managed at district clinics by nurses trained in how to diagnosis and treat the disease. There is no reason why they should not have responsibility, with supervision and appropriate advice for problems, for managing suspects, making a definite diagnosis and initiating and continuing treatment. Simplification of treatment regimens currently recommended by the South African tuberculosis control programme has been suggested by many experts.
- * Systems of transport and communication between all clinics and the district laboratory is essential for a functioning tuberculosis programme. Innovative ways and systems must be developed to supply drugs and other requisites, and to collect specimens, all of which will benefit the delivery of comprehensive health care.
- * All patients should have supporters. Community members make excellent supporters, but need training incentives, monitoring and evaluation.
- * Backup support for recruiting community supporters, and for following up non-adherent patients will always be needed. To provide transport at each clinic for clinic staff to do this themselves is probably not cost-effective. At district level it would be, if the staff responsible for tuberculosis was also responsible for other communicable diseases, surveillance systems and outbreak responses.
- * Completion of the various cards, register and regular reports should also be done by clinic nurses, who will find the reports easier and more acceptable when some modifications are made, and with continuing training. The district communicable disease coordinator should monitor the correct completion of all forms, and collate those of all clinics into a district report.

- * Transfer of patients who are leaving the district should be facilitated by encouraging those with intended movements to discuss where they are going. Transfers to other services must be formalised with appropriate documentation and direct communication to the receiving health services. Work should proceed on implementing a national computerised register of all patients with tuberculosis, which will facilitate transfers as well as record details of current and previous treatment of all patients.

REFERENCES

1. Cave AJE, The evidence for the incidence of tuberculosis in ancient Egypt. *Br J Tuberc* 1939;33:142-152.
2. Formicola V, Milanese Q, Scarsini C. Evidence of spinal tuberculosis at the beginning of the fourth millenium BC from Arene Candide Cave (Liguria, Italy) *Am J Phy Anthropol* 1987;72:1-6.
3. Stirland A, Waldron T. The earliest cases of tuberculosis in Britain. *J Arch Sc.* 1990; 17: 221-230. In: Ryan Frank Tuberculosis: The Greatest Story never told. Great Britain: Swift, 1992:5-6.
4. Dubos R, Dubos J. The White Plague: Tuberculosis, Man and Society. Second edition. Rutgers University Press, New Brunswick, New Jersey.1992.
5. Grigg ERN The Arcana of Tuberculosis. *Am Rev of Tuberc Pulm Dis* 1958;78:151-172.
6. Bignall JR. A century of treating tuberculosis. *Tubercle* 1982;63:19-22.
7. de Kruif P. Koch: the Death Fighter In: Microbe Hunters Pocket ed. New York:Pocket Books, Inc, 1926.
8. Snider Gordon J. Tuberculosis Then and Now: A Personal Perspective on the Last 50 Years. *Ann Intern Med* 1997; 126(3):237-243.
9. Comstock George W. Tuberculosis: Is the Past Once Again Prologue? *Am J Public Health* 1994;84(11):1729-1731.
10. Beyers AD, Donald PR, van Helden PD, Ehlers MRW, Steyn L, Mizrahi V. Tuberculosis researh - the way forward. *S Afr Med J* 1996;86(1):30-33.
11. Horton R. Towards the elimination of tuberculosis. *Lancet* 1995;346:790.
12. Dunlap NE, Briles DE. Immunology of Tuberculosis. *Med Clin North Am* 1993;77(6):1235-1251.
13. Bloom B, Murray CJL. Tuberculosis: Commentary on a Reemergent Killer. *Science* 1992;257:1055-1064.
14. Stehbens WE. On the "cause" of Tuberculosis. *Pathology* 1987;19:115-119.
15. Rieder HL, Cauthen GM, Comstock GW, Snider DE Jr. Epidemiology of Tuberculosis in the United States. *Epidemiol Rev* 1989;11:79-96.
16. Grzybowski S, Barnett GD, Styblo K. Contacts of cases of active pulmonary tuberculosis. *Bull Int Union Tuberc* 1975;50:90-106.

17. Beyers N, Gie RP, Scaaf HS, Van Zyl S, Talent JM, Nel ED, Donald PR. A prospective evaluation of children under the age of 5 years living in the same household as adults with recently diagnosed pulmonary tuberculosis. *Int J Tuberc Lung Dis* 1997;1(1):38-43.
18. Kenyon TA, Valway SE, Ihle WW, Onorato IM, Castro KG. Transmission of Multidrug-resistant *Mycobacterium Tuberculosis* during a long airplane flight. *N Engl J Med* 1996;334(15):933-938.
19. Styblo K, Meier J, Sutherland I. The transmission of tubercle bacilli: its trend in a human population. *Tuberculosis Surveillance Report no.1 Bull Int Union Tuberc* 1969;42:1-104.
20. Sudre P, ten Dam G, Kochi A. Tuberculosis: a global overview of the situation today. *Bull World Health Organ* 1992;70(2):149-159.
21. Sutherland I. Research into the control of tuberculosis and leprosy in the community. *Br Med Bull* 1988;44(3):665-678.
22. Armstrong JA, D'Arcy Hart P. Response of cultured macrophages to *Mycobacterium tuberculosis*, with observations on fusion of lysosomes with phagosomes. *J Exp Med* 1971;134:713-740.
23. Ehlers MRW. *Mycobacterium tuberculosis* : infection or disease? Key events in pathogenesis and the immune response. *Specialist Medicine* 1994;11:48-55.
24. Cole ST. Why sequence the genome of *Mycobacterium tuberculosis*? *Tubercle Lung Dis* 1996;77(6):486-490.
25. Styblo K. Epidemiology of Tuberculosis. Royal Netherlands Tuberculosis Association. The Hague, The Netherlands 1984. Reissued in: *Selected Papers* 1991.
26. Small PM, Moss A. Molecular epidemiology and the new tuberculosis. *Infectious Agents and Disease*, Raven Press 1993;2(3):132-138.
27. Small PM, Hopewell PC, Singh SP, Paz A, Parsonnet J, Ruston DC et al. The Epidemiology of Tuberculosis in San Francisco: a population-based study using conventional and molecular methods. *N Engl J Med* 1994;330:1703-1709.
28. Murray CJL, Styblo K, Rouillon A. Tuberculosis in developing countries: burden, intervention and cost. *Bull Int Union Tuberc Lung Dis* 1990;65(1):6-24.
29. Comstock GW, Livesay VT, Woolpert SF. The prognosis of a positive tuberculosis reaction in childhood and adolescence. *Am J Epidemiol.* 1974;99:131-138.
30. Selwyn PA, Hartel D, Lewis VA et al. A prospective study of the risk of tuberculosis among intravenous drug users with human immunodeficiency virus

infection. *N Eng J Med* 1989;320:545-550

31. De Cock KM, Soro B, Coulibaly IM, Lucas SB. Tuberculosis and HIV infection in Sub-Saharan Africa. *JAMA* 1992;268:1581-1587
32. van den Broek J, Borgdorff MW, Pakker NG, Chum HJ, Klokke AH, Senkoro KP, Newell JN. HIV-1 Infection as a Risk factor for the Development of Tuberculosis: a Case Control Study in Tanzania. *Int J Epidemiol* 1993;22(6):1159-1165.
33. CDC. Tuberculosis-United States, 1985-and the possible impact of human lymphotropic virus type III/lymphadenopathy-associated virus infection. *Morb Mort Wkly Rep* 1986;35:76-86.
34. Brudney K, Dobkin J. Resurgent Tuberculosis in New York City. *Am Rev Respir Dis* 1991;144:745-749.
35. Reichman LB. The U-shaped Curve of Concern (editorial). *Am Rev Resp Dis* 1991;144:741-742.
36. Richards SB, St Louis MB, Nieberg P, Coulibaly IM, Coulibaly D, Abouya I, Gayle HD, De Cock KM. Impact of the HIV epidemic on trends in tuberculosis in Abidjan, Cote d'Ivoire. *Tubercle Lung Dis* 1995;76(1):11-16.
37. Daley CL, Small PM, Schecter GF, Schoolnik G, McAdam R, Jacobs W, et al. An outbreak of tuberculosis with accelerated progression among persons infected with the human immunodeficiency virus. *N Eng J Med* 1992;326:231-235
38. Nunn P, Mungai M, Nyamwaya J, Gicheha C, Brindle RJ, Dunn DT, Githui W, Were JO, McAdam KPWJ. The effect of Human Immunodeficiency Virus type-1 on the infectiousness of tuberculosis. *Tubercle Lung Dis* 1994;75:25-32.
39. Raviglione MC, Snider DE Jr, Kochi A. Global Epidemiology of Tuberculosis: Morbidity and Mortality of a Worldwide Epidemic. *JAMA* 1995;273:220-226.
40. Styblo K. The global aspects of tuberculosis and HIV infection *Bull Int Union Tubercle Lung Dis* 1990;65(1):28-33.
41. Grzybowski S. Tuberculosis in the Third World. *Thorax* 1991;46:689-691.
42. Tuberculosis, HIV on collision course in Asia. World Health Organisation, Office of Information, press release, August 10, 1994. (WHO/63)
43. Narain JP, Raviglione MC, Kochi A. HIV-associated tuberculosis in developing countries: epidemiology and strategies for prevention. *Tubercle Lung Dis* 1992;73:311-321.
44. Schulzer M, Fitzgerald JM, Enarson DA, Grzybowski S. An estimate of the future size of the tuberculosis problem in sub-Saharan Africa resulting from HIV-

infection. *Tuber Lung Dis* 1992;73:52-58.

45. Dolin PJ, Ravigliione MC, Kochi A. Global tuberculosis incidence and mortality during 1990-2000. *Bull World Health Organ* 1994;72:213-220.
46. UNAIDS Fact Sheet, December 1996
47. Cantwell MF, Binkin NJ. Tuberculosis in sub-Saharan Africa: a regional assessment of the impact of the human immunodeficiency virus and National Tuberculosis Control Program quality. *Tuber Lung Dis* 1996;77:220-225.
48. Spence DPS, Hotchkiss J, Williams CSD, Davies PDO. Tuberculosis and poverty. *Br Med J* 1993;307:759-761.
49. Snider DE Jr. The relationship between tuberculosis and silicosis (Editorial) *Am Rev Respir Dis* 1978;118:455-460.
50. Cowie RL. The Epidemiology of Tuberculosis in Gold Miners with Silicosis. *Am J Respir Crit Care Med* 1994;150:1460-1462.
51. Yu G, Hseih C, Peng J. Risk factors associated with the prevalence of pulmonary tuberculosis among sanitary workers in Shanghai. *Tubercle* 1988;69:105-112.
52. Buskin SE, Gale JL, Weiss NS, Nolan CM. Tuberculosis risk factors in Adults in King County, Washington 1988 through 1990. *Am J Public Health* 1994;84:1750-1756.
53. Alcaide J, Altet MN, Plans P, Parron I, Folguera LI, Salto E et al Salleras LI. Cigarette smoking as a risk factor for tuberculosis in young adults: a case contro' study. *Tuber Lung Dis* 1996;77(2):112-116.
54. Stead WW, Lofgren JP. Does the risk of tuberculosis increase in old age? *J Infect Dis* 1983;147:951-955
55. Rocha M, Pereira S, Barros H, Seabra J. Does pulmonary tuberculosis change with ageing? *Int J Tuberc Lung Dis* 1997;1(2):147-151.
56. Grzybowski S, Enarson DA. The fate of cases of pulmonary tuberculosis under various treatment programmes. *Bull Int Union Tuberc* 1978;53:70-75.
57. Rieder HL. Epidemiology of tuberculosis in Europe. *Eur Respir J* 1995;8 Suppl.20:620s-632s.
58. Styblo K. The relationship between the risk of infection and the risk of developing infectious tuberculosis. *Bull Int Union Tuberc* 1985;60(3-4):117-119.
59. Grzybowski S. Natural history of tuberculosis. *Bull Int Union Tuberc Lung Dis* 1991;66:193-194.

60. Styblo K. Tuberculosis and its control: lessons to be learned from past experience, and implications for leprosy control programmes. The Kellersberger Memorial Lecture 1982. *Ethiop Med J* 1983;21:101-122.
61. Grzybowski S, Styblo K, Dorken E. Tuberculosis in Eskimos. *Tubercle* 1976; 57(4) Suppl;S1-S58.
62. Nunn PP, Elliot AM, McAdam KPWJ. Impact of human immunodeficiency virus on tuberculosis in developing countries. *Thorax* 1994;49:511-518.
63. World Health Organisation. The HIV/AIDS and Tuberculosis Epidemics: Implications for control. WHO, Geneva, 1994. WHO/TB/CARG(4)/94.4
64. Rouillon A, Perdrizat S, Parrot R. Transmission of Tubercle Bacilli: the effects of chemotherapy. *Tubercle* 1976;57:275-299.
65. Toman K. Sensitivity, specificity and predictive value of diagnostic tests. *Bull Int Union Tuberc* 1981;56(1):18-27.
66. Pust RE. The risk factor approach to sputum diagnosis. *World Health Forum* 1982;3(1):78-80
67. Rouillon A. The Mutual Assistance Programme of the IUATLD. Development, contribution and significance. *Bull Int Tuberc Lung Dis* 1991;66:159-172
68. Enarson DA. The International Union Against Tuberculosis and Lung Disease model National Tuberculosis Programmes. *Tuber Lung Dis* 1995;76:95-99
69. Grzybowski S. The value of different diagnostic and preventive programmes. *S Afr Med J* 1982; 17 Nov Special issue;6-8.
70. Toman K. How reliable is smear microscopy? In: Tuberculosis case finding and chemotherapy. Geneva: WHO, 1979:15.
71. Gebre N, Karlsson U, Jonsson G, Macaden R, Wolde A, Assefa A et al. Improved microscopical diagnosis of pulmonary tuberculosis in developing countries. *Trans R Soc Trop Med Hyg* 1995;89:191-193.
72. Schluger NW, Rom WN. Current Approaches to the Diagnosis of Active Pulmonary Tuberculosis. *Am J Crit Care Med* 1994;149:264-267
73. Elliott AM, Luo N, Tembo G, Halwiindi B, Steenbergen G, Machiels L, et al. Impact of HIV in tuberculosis in Zambia: a cross sectional study. *Br Med J* 1990;301:412-415.
74. Barnes PF, Bloch AB, Davidson PT, Snider DE, Jr. Tuberculosis in patients with human immunodeficiency virus infection. *N Eng J Med* 1991;324:1644-1650.

75. Toman K. What are the advantages and disadvantages of fluorescent microscopy? In: Tuberculosis case finding and chemotherapy. Geneva:WHO,1979:25.
76. Toman K. What is the additional case yield from repeated sputum examinations by microscopy and culture? In: Tuberculosis case finding and chemotherapy. Geneva: WHO,1979:40-42.
77. Urbanczik R. Present position of Microscopy and of Culture in Diagnostic Mycobacteriology. Zbl Bakt Hyg 1985;A260:81-87.
78. Harries AD, Kamenya A, Namarika D, Msolomba IW, Salaniponi FM, Nyangulu DS, Nunn P. Delays in diagnosis and treatment of smear positive tuberculosis and the incidence of tuberculosis in hospital nurses in Blantyre, Malawi. Trans R Soc Trop Med Hyg;1997;91:15-17.
79. Mitchison DA. Organisation of Tuberculosis Laboratory Services in Developing Countries. Bull Int Union Tuberc 1982;57(42):140-145.
80. Aluoch JA, Edwards EA, Stott H, Fox W, Sutherland I. A fourth study of case finding methods for pulmonary tuberculosis in Kenya. Trans R Soc Trop Med Hyg 1982;76:679-691.
81. Toman K. What is the probability of obtaining a negative culture from a sputum specimen found positive by smear microscopy? Tuberculosis case finding and chemotherapy. Geneva: WHO,1979:38-39.
82. Gordin FM, Slutkin G, Schechter G, Goodman PC, Hopewell PC. Presumptive diagnosis and treatment of pulmonary tuberculosis based on radiographic findings. Am Rev Respir Dis 1989;139:1090-1093.
83. Toman K. What is the clinical and epidemiological significance of consistent negativity by smear microscopy in patients positive by culture only? In: Tuberculosis case finding and chemotherapy. Geneva: WHO,1979:50-55.
84. Shaw JB, Wynn-Williams N. Infectivity of pulmonary tuberculosis in relation to sputum status. Am Rev Tuberc 1954;69:724-732.
85. Fox W. Short-course chemotherapy for pulmonary tuberculosis and some problems of its programme application with particular reference to India. Lung India 1984;II(2):161-174.
86. Tuberculosis research and development. Report of a WHO meeting Geneva October 1990. Tuberculosis Unit, Division of Communicable Diseases, WHO;1990.
87. Woodring JH, Vandiviere HM, Fried AM, Dillon ML, Williams TD, Melvin IG. Update: the radiographic features of Pulmonary tuberculosis. Am J Roentgenol 1986;146:497-506.

88. Toman K. How reliable is chest radiography? In: Tuberculosis case finding and chemotherapy. Geneva: WHO, 1979:28-37.
89. Toman K. Mass radiography in tuberculosis control. WHO Chron 1976; 30: 51-57.
90. WHO Technical Report Series, No. 552. Geneva 1974:15-16.
91. De Cock KM Impact of Interaction with HIV. In: Porter JDH, McAdam KPWJ, editors. Tuberculosis Back to the future. Chichester: John Wiley and sons Ltd, 1994:41.
92. Chaulet P. Diagnosis of tuberculosis in children. Children in the tropics: Childhood tuberculosis is still with us. Paris: International Children's Centre, 1992.
93. Godfrey-Fausset P. Of molecules and men: the detection of tuberculosis, past, present and future. In: Porter JDH, McAdam KPWJ, editors. Tuberculosis Back to the future. Chichester: John Wiley and sons Ltd, 1994:79-95.
94. Long R, Scalchini M, Manfreda J, Jean-Baptiste M, Hershfield E. The Impact on the Usefulness of Sputum Smears for the Diagnosis of Tuberculosis. Am J Public Health 1991;81:1326-1328.
95. Elliott AM, Namaambo K, Allen BW, Luo N, Hayes RJ, Pobee JOM, McAdam KPWJ. Negative sputum smear results in HIV-positive patients with pulmonary tuberculosis in Lusaka, Zambia. Tuber Lung Dis 1993;74:191-194.
96. Long R, Scalchini M, Manfreda J, Jean-Baptiste M, Hershfield E. The Impact of HIV on the Usefulness of Sputum Smears for the Diagnosis of Tuberculosis. Am J Public Health 1991;81:1326-1328.
97. Aluoch JA, Oyoo D, Swai OB et al. A study of the use of maternity and child welfare clinics in case finding for pulmonary tuberculosis in Kenya. Tubercle 1987;68:93-103.
98. Banerji D, Anderson S. Sociological Study of Awareness of Symptoms among persons with Pulmonary tuberculosis. Bull World Health Organ 1963;29:665-683.
99. Elink Schuurman MW, Srisaenpang S, Pinitsoontorn S, Bijleveld I, Vaeteewoothacharn K, Methapat C. The rapid village survey in tuberculosis control. Tuber Lung Dis 1996;77:549-554.
100. Rieder HL. Case finding. In: Reichman LB, Hershfield ES, editors. Tuberculosis - a comprehensive international approach. New York: Marcel Dekker, Inc, 1993:167-182.
101. Pamra SP. Problems of tuberculosis in developing countries. Clin Chest Med 1980;1(2):265-271.

102. McAdam KPWJ. Back to the future: 'the ten commitments.' In Porter JDH, McAdam KPWJ eds: *Tuberculosis: back to the future*. Chichester: John Wiley, 1994;267-276.
103. Fine PEM. The BCG story, lessons from the past and implications for the future. *Rev Infect Dis* 1989;Suppl2:353-359.
104. Ferebee SH. Controlled chemoprophylaxis trials in tuberculosis. *Adv Tuberc Res* 1970;17:28-106.
105. Comstock GW, Woolpert SF. Preventive therapy. In: Kabuki GP, Wayne LG, editors. *The Mycobacteria. A source-book*. New York and Basel, Marcel Dekker, 1984:1071-1082.
106. IUATLD. Tuberculosis in children - guidelines for diagnosis, prevention and treatment. *Bull Int Union Tuberc Lung Dis* 1991;66:61-67.
107. American Thoracic Society/Centers for Disease Control. Treatment of tuberculosis and tuberculosis infection in adults and children. *Am Rev Respir Dis* 1986;134:355-363.
108. Centers for Disease Control. The use of preventive therapy for tuberculosis infection in the United States. Recommendations of the Advisory Committee for the Elimination of Tuberculosis. *Morb Mortal Wkly Rep* 1990;39(RR-6):9-12.
109. Comstock G. Prevention of tuberculosis. *Bull Int Union Tuberc Lung Dis* 1990/1991;66(Suppl):9-11.
110. Styblo K. Preventive therapy for tuberculosis control in developing countries. *Bull Int Union Tuberc Lung Dis* 1990/1991;66(Suppl):27-28.
111. Mwinga AG. Preventive therapy for tuberculosis. Discussion. In: Porter JDP, McAdam KPWJ, editors. *Tuberculosis: Back to the future*. Chichester: John Wiley and Sons Ltd, 1994:166-169.
112. Pape JW, Jean SS, Ho JL, Hafner A, Johnson WD. Effect of isoniazid on incidence of active tuberculosis and progression of HIV infection. *Lancet* 1993;342:268-272.
113. Hawken MP, Meme HK, Elliott LC, Chakaya JM, Morris JS, Githui WA et al. Isoniazid preventive therapy for tuberculosis in HIV-1-infected adults: results of a randomised controlled trial. *AIDS* 1997;11:875-882.
114. Centers for Disease Control. Purified protein derivative (PPD) - tuberculin anergy and HIV-infection: guidelines for anergy testing and management of anergic persons at risk of tuberculosis. *Morb Mortal Wkly Rep* 1991;40(supplRR-5):27-33.

115. Editorial. Africa's tuberculosis burden and chemoprophylaxis. *Lancet* 1991;335:1249-1250.
116. Grzybowski S. Preventive treatment for tuberculosis control in developing countries. *Bull Int Union Tuberc Lung Dis* 1990/1991;66 Suppl:25.
117. TB Programme. DOTS stops TB at the source: WHO report on the tuberculosis epidemic. 1995 (WHO/TB/95.183).
118. Harries AD, Maher D. TB/HIV: A Clinical Manual. World Health Organisation Global Tuberculosis Programme, 1996.
119. Porter JDH, McAdam KPWJ. Aspects of AIDS in Africa: 1. Tuberculosis in Africa in the AIDS era - the role of chemoprophylaxis. *Trans R Soc Trop Med Hyg* 1992;86:467-469.
120. Joint statement by the International Union against Tuberculosis and Lung Disease and the Global Programme on AIDS and the Tuberculosis Programme of the World Health Organisation. Tuberculosis preventive therapy in HIV-infected individuals. *Tuber Lung Dis* 1994;75:96-98.
121. Monaco A. Antituberculous chemoprophylaxis in silicotics. *Bull Int Union Tuberc* 1964;35:51-54.
122. Hong Kong Chest Service/Tuberculosis Research Centre, Madras/British Medical Research Council. A Double-Blind Placebo-controlled Clinical Trial of Three Antituberculosis Chemoprophylaxis Regimes in Patients with Silicosis in Hong Kong. *Am Rev Respir Dis* 1992;145:36-41.
123. Cowie RL. Short course chemoprophylaxis with rifampicin, isoniazid and pyrazinamide for tuberculosis evaluated in gold miners with chronic silicosis: a double-blind placebo controlled trial. *Tuber Lung Dis* 1996;77:239-243.
124. Mitchison DA. Chemotherapy of tuberculosis: a bacteriologist's viewpoint. *Br Med J* 1965;1:1331-1338.
125. Medical Research Council Investigation. Streptomycin treatment of pulmonary tuberculosis. *Br Med J* 1948;2:769-782.
126. Fox W. The modern management and therapy of pulmonary tuberculosis. *Proceedings of the Royal Society of Medicine* 1977;70:4-15.
127. Crofton J. Drug treatment of tuberculosis. 1 Standard chemotherapy *Br Med J*. 1960;2:370-373.

128. Tuberculosis Chemotherapy Centre, Madras: A concurrent comparison of intermittent (twice weekly) isoniazid plus streptomycin and daily isoniazid plus PAS in the domiciliary treatment of pulmonary tuberculosis. *Bull World Health Organ* 1964;31:247-54.
129. Tuberculosis Chemotherapy Centre, Madras. A Controlled Comparison of a Twice-weekly and Three Once-Weekly Regimens in the Initial Treatment of Pulmonary Tuberculosis. *Bull World Health Organ* 1970;43:143-206.
130. British Medical Research Council. Co-operative controlled trial of a standard regimen of streptomycin, PAS, and isoniazid and three alternative regimens of chemotherapy in Britain. *Tubercle* 1973;54:99-129.
131. Fox W. Whither short-course chemotherapy? *Br J Dis Chest*. 1981;75:331-357.
132. Styblo K, Chum HJ. Treatment results of smear positive tuberculosis in the Tanzania National Tuberculosis and Leprosy Programme: standard and short-course chemotherapy. Proceedings of the XXVIth World Conference on Tuberculosis and Respiratory Disease. Singapore 4-7 November 1986. Tokyo, Professional Postgraduate Services, 1987:122-126.
133. Grosset J. Bacteriologic basis of short-course chemotherapy for tuberculosis. *Clin Chest Med* 1980;1:231-241.
134. Mitchison DA. The action of anti-tuberculosis drugs in short-course chemotherapy. *Tubercle* 1985;66:219-225.
135. Hopewell PC. 'The cure':organisation and administration of therapy for tuberculosis. In: Porter JDH, McAdam KPWJ, editors. *Tuberculosis: Back to the future*. Chichester: John Wiley and Sons Ltd, 1994:99-121.
136. WHO. Treatment of tuberculosis: Guidelines for National Programmes. WHO, Geneva 1997.WHO/TB/97.220
137. Enarson DA, Rieder HL, Arnadottir T, Trebusq A. Tuberculosis Guide for low income countries. International Union Against Tuberculosis and Lung Disease. 4th ed. Frankfurt, Germany:pmi Verslagsgruppe GmbH. 1996.
138. Doyle JA, Lambertini A, Dolmann A. Intermittent chemotherapy with three drugs in the treatment of pulmonary tuberculosis. *Tubercle* 1970;51:397-402.
139. Hong Kong Chest Service/British Medical Research Council. Controlled Trial of 2, 4, and 6 Months of Pyrazinamide in 6-Month, Three-Times-Weekly Regimens for Smear-positive Pulmonary Tuberculosis, Including an Assessment of a Combined Preparation of Isoniazid, Rifampicin, and Pyrazinamide. *Am Rev Respir Dis* 1991;143:700-706.

140. Cohn DL, Catlin BJ, Peterson KL, Judson FN, Sbarbaro JA. A 62-Dose, 6-Month Therapy for Pulmonary and Extrapulmonary Tuberculosis. *Ann Intern Med* 1990;112:407-415.
141. Dutt AK, Moers D, Stead WW. Short-Course Chemotherapy for Tuberculosis with Mainly Twice-Weekly Isoniazid and Rifampicin: Community Physicians' Seven-Year Experience with Mainly Outpatients. *Am J Med* 1984;77:233-242.
142. Hong Kong Chest Service/British Medical Research Council. Acceptability, compliance, and adverse reactions when isoniazid, rifampicin, and pyrazinamide are given as a combined formulation or separately during three-times-weekly anti-tuberculosis chemotherapy. *Am Rev Respir Dis* 1989;140:1618-1622.
143. Macnab MF, Bohmer PD, Seager JR. Evaluation of the 3-drug combination, Rifater, versus 4-drug therapy in the ambulatory treatment of tuberculosis in Cape Town. *S Afr Med J* 1994;84:325-328.
144. Cowie RL, Brink BA. Short course chemotherapy for pulmonary tuberculosis with a rifampicin-isoniazid-pyrazinamide combination tablet. *S Afr Med J* 1990;77:390-391.
145. Singapore Tuberculosis Service/British Medical Research Council. Assessment of a daily Combined Preparation of Isoniazid, Rifampicin and Pyrazinamide in a Controlled Trial of Three 6-month Regimens for Smear Positive Pulmonary Tuberculosis. *Am Rev Respir Dis* 1991;143:707-712.
146. Fox W. Drug combinations and the bioavailability of rifampicin. *Tubercle* 1990;71:241-245.
147. Joint statement of the IUATLD and the Tuberculosis Programme of the WHO. The promise and reality of fixed combinations with rifampicin. *Tuber Lung Dis* 1994;75: 180-181.
148. Rieder HL. Sputum smear conversion during directly observed treatment for tuberculosis. *Tuber Lung Dis* 1996;77:124-129.
149. Mitchison DA. Assessment of new sterilizing drugs for treating pulmonary tuberculosis by culture at 2 months [correspondence]. *Am Rev Respir Dis* 1993;147:1062-1063.
150. Rieder HL, Enarson DA. Rebuttal: time to call a halt to emotions in the assessment of thioacetazone. *Tuber Lung Dis* 1996;77:109-111.
151. Murray C, Styblo K, Rouillon A. Tuberculosis. In: Jamison DT, Mosley WH, Measham AR, Bobadilla JL, editors. *Disease Control: Priorities in Developing Countries*. New York: Oxford University Press, 1993:233-259.

152. Nunn PP, Brindle R, Carpenter L, Odhiambo J, Wasunna K, Newnham R et al. Cohort study of HIV infection in tuberculosis patients, Nairobi, Kenya. Analysis of early (6-months) mortality. *Am Rev Respir Dis* 1992;146:849-854.
153. Pertinens JH, Colebunders RL, Karahunga C, Willame J-C, Jeugmans J, Kagoto M, et al. Increased Mortality and Tuberculosis Treatment Failure rate among Human Immunodeficiency Virus (HIV) Seropositive Compared with HIV Seronegative Patients with Pulmonary Tuberculosis Treated with "Standard" Chemotherapy in Kinshasa, Zaire. *Am Rev Respir Dis* 1991;144:750-755.
154. Wilkinson D, Moore DAJ. HIV-related tuberculosis in South Africa- clinical features and outcome. *S Afr Med J* 1996;86:64-67.
155. Haken M, Nunn P, Gathua S, Brindle R, Godfrey-Faussett P, Githui W et al. Increased recurrence of tuberculosis in HIV-1 infected patients in Kenya. *Lancet* 1993;342:332-337.
156. Sepkowitz KA, Raffalli J, Riley L, Kiehn TE, Armstrong D. Tuberculosis in the AIDS era. *Clin Microbiol Rev* 1995;8:180-199.
157. Centers for Disease Control. Diagnosis and management of mycobacterial infection and disease in persons with human immunodeficiency virus infection. Centers for Disease Control, US Department of Health and Human Services. *Ann Intern Med* 1987;106:254-256.
158. Holt E, Cantave M, Johnson M, Clermont HC, Atkinson J, Chaisson RE. Efficacy of supervised, intermittent, short course therapy of tuberculosis in Haiti. *Abst WS-B09-4:52* In: Program and abstracts of the 9th International Conference on AIDS 1993.
159. Small PM, Schecter GF, Goodmark PC, Sande MA, Chaisson RE, Hopewell PC. Treatment of tuberculosis in patients with advanced immunodeficiency virus infection. *N Engl J Med* 1991;324:289-294.
160. Elliott AM. An approach to the management of tuberculosis in HIV endemic areas. *Trop Doct* 1992;22:147-150.
161. Elliott AM, Foster SD. Thiacetazone: time to call a halt? *Tuber Lung Dis* 1996;77:27-29.
162. Ipuge YAI, Rieder HL, Enarson DA. Adverse cutaneous reactions to thiacetazone for tuberculosis treatment in Tanzania. *Lancet* 1995;346:657-660.
163. Nunn P, Porter J, Winstanley P. Thiacetazone - avoid like poison or use with care? *Trans R Soc Tropical Med Hyg* 1993;87:578-582.
164. Escreet BC, Langton ME, Cowie RL. Short-course chemotherapy for silicotuberculosis. *S Afr Med J* 1984;66:327-330.

165. Cowie RL, Langton ME, Becklake MR. Pulmonary Tuberculosis in South African Coal Miners. *Am Rev Resp Dis* 1989;139:1086-1089.
166. Suo J, Lee CN, Lee JJ, Yang SP. Short-course Chemotherapy of Pulmonary Tuberculosis in Pneumoconiotic patients. *Am Rev Respir Dis* 1987;136:808-810.
167. Cowie RL. Silicotuberculosis: long-term outcome after short-course chemotherapy. *Tuber Lung Dis* 1995;76:39-42.
168. Grange JM. Drug resistance and tuberculosis elimination. *Bull Int Union Tuberc Lung Dis* 1990;65(2-3):57-59.
169. Rieder HL. Drug-resistant tuberculosis: issues in epidemiology and challenges for public health. (Editorial) *Tuber Lung Dis* 1994;75:321-323.
170. Crofton J. Multidrug resistance: danger for the Third World. In Porter JDH, McAdam KPWJ. Editors: *Tuberculosis: Back to the future*. Chichester: John Wiley and Sons Ltd, 1994:231-233.
171. Nunn P, Felten M. Surveillance of resistance to anti-tuberculosis drugs in developing countries. *Tuber Lung Dis* 1994;75:163-167.
172. Cohn D, Bustreo F, Raviglione M. Drug resistance in tuberculosis: review of the worldwide situation and WHO surveillance. *Clin Infect Dis* 1996 (in press), quoted in: Crofton J, Chaulet P, Maher D. *Guidelines on the management of drug resistant tuberculosis*. World Health Organisation, Geneva. 1996 WHO/TB/96.210.
173. Bustreo F, Migliori GB, Nardini S, Raviglione MC. Antituberculosis drug resistance: is it worth measuring?. The WHO/IUATLD working group on antituberculosis drug resistance surveillance. *Monaldi Arch Chest Dis* 1996;51(4):299-302.
174. Centers for Disease Control. Nosocomial transmission of multidrug resistance among HIV infected persons; Florida and New York, 1988-1991. *JAMA* 1991;266:1483-1485.
175. Beck-Sague C, Dooley SW, Hutton MD, Otten J, Breedon A, Crawford JT, et al. Hospital outbreak of multidrug resistant *Mycobacterium tuberculosis* infections: factors in transmission to staff and HIV-infected patients. *JAMA* 1992;268:1280-1286.
176. Edlin BT, Tokars JJ, Grieco MH, Crawford JT, Williams J, Sordillo EM et al. An outbreak of multidrug resistant tuberculosis among hospitalised patients with the acquired immunodeficiency syndrome. *N Engl J Med* 1992;326:1514-1521.

177. Pearson ML, Jereb JA, Frieden TR, Crawford JT, Davis BJ, Dooley SW, et al. Nosocomial transmission of multidrug-resistant *Mycobacterium tuberculosis*: a risk to patients and health workers. *Ann Intern Med* 1992;117:191-196.
178. Nolan CM. Multidrug-resistant tuberculosis in the USA: the end of the beginning. *Tubercle* 1996;77:293-294.
179. Frieden TR, Fujiwara PI, Washko RM, Hamburg MA. Tuberculosis in New York City - Turning the tide. *N Engl J Med* 1995;333:229-233.
180. Neville K, Bromberg A, Bromberg R, Bonk S, Hanna BA, Rom WN. The Third Epidemic - Multidrug Resistant Tuberculosis. *Chest* 1994;105:45-48.
181. Enarson DA. Will HIV hasten appearance of resistance? Proceedings of the conference on Global Lung Health and the 1996 Annual Meeting of the International Union Against Tuberculosis and Lung Disease; 1996 Oct2-5; Paris. *Tubercle* 1996;77(suppl2):14.
182. Crofton J. The prevention and management of drug-resistant tuberculosis. *Bull Int Union Tuberc Lung Dis* 1987;62:6-11.
183. Chaulet P. Drug resistant tuberculosis: can it be cured? Proceedings of the conference on Global Lung Health and the 1996 Annual Meeting of the International Union Against Tuberculosis and Lung Disease; 1996 Oct2-5; Paris. *Tubercle* 1996;77(suppl2):9.
184. Crofton J, Chaulet P, Maher D. Guidelines on the management of drug resistant tuberculosis. World Health Organisation, Geneva. 1996 WHO/TB/96.210.
185. Kochi A. The global tuberculosis situation and the new control strategy of the World Health Organisation. *Tubercle* 1991;72:1-6.
186. WHO Tuberculosis Programme: Framework for effective tuberculosis control. WHO, Geneva, 1994. WHO/TB/94.179.
187. Fox W. The problem of self-administration of drugs: with particular reference to pulmonary tuberculosis. *Tubercle* 1958;39:269-74.
188. Fox W. Self-administration of medicaments: a review of published work and a study of the problem. *Bull Int Union Tuberc* 1962;32:307-331.
189. Sbarbaro JA. Compliance: Inducements and Enforcements. *Chest* 1979 Suppl;76:750-756.
190. Bayer R, Wilkinson D. Directly observed therapy for tuberculosis: history of an idea. *Lancet* 1995;345:1545-1548.

191. McDonald RJ, Memon AM, Reichman LB. Successful Supervised Ambulatory Management of Tuberculosis Treatment failures. *Ann Intern Med* 1982;96:297-302.
192. Weis SE, Slocum PC, Blais FX, King B, Nunn M, Matney GB, et al. The effect of directly observed therapy on the rates of drug resistance and relapses in tuberculosis. *N Engl J Med* 1994;330:1179-1184.
193. Idukitta GO, Bosman MCJ. The Tuberculosis Manyatta Project for Kenyan nomads. *Bull Int Union Tuberc Lung Dis* 1989;64:44-47.
194. Neher A, Breyer G, Shrestha B, Feldmann K. Directly observed intermittent short-course chemotherapy in the Kathmandu valley. *Tuber Lung Dis* 1996;77:302-307.
195. Chowdhury AMR, Chowdhury S, Islam MN, Islam A, Vaughan P. Control of tuberculosis by community health workers in Bangladesh. *Lancet* 1997;350:169-172.
196. Dick J, Schoeman JH, Mohammed A, Lombard C. Tuberculosis in the community: 1. Evaluation of a volunteer health worker programme to enhance adherence to anti-tuberculosis treatment. *Tuber Lung Dis* 1996;77:274-279.
197. Iseman MD, Cohn DL, Sbarbaro JA. Directly Observed Treatment of Tuberculosis: We Can't Afford Not to Try it. *N Engl J Med* 1993;328:576-578.
198. Floyd K, Wilkinson D, Gilks CF. Community-based, Directly Observed Therapy for tuberculosis: An economic analysis. Tygerberg, South Africa: Medical Research Council, 1997.
199. Nardell EA. Beyond Four Drugs: Public Health Policy and the Treatment of the Individual Patient with Tuberculosis [editorial]. *Am Rev Respir Dis* 1993;148:2-5.
200. Wilkinson D. Compliance with tuberculosis treatment in a rural district. [correspondence] *Tuber Lung Dis* 1995;76:180.
201. Wilkinson D, Davies GR. Coping with Africa's increasing tuberculosis burden: are community supervisors an essential component of the DOT strategy? *Trop Med Int Health* 1997;2(7):700-704.
202. DiFerdinando GT, Klein SJ, Novick LF. [correspondence]. *N Engl J Med* 1993;329:135-136.
203. Annas GJ. Control of Tuberculosis - the law and public's health. *N Engl J Med* 1993;328:585-588.
204. Maher D, Hausler HP, Raviglione MC, Kaleeba N, Aisu T, Fourie B, Nunn P. Tuberculosis care in community care organisations in Sub-Saharan Africa: practice and potential. *Int J Tuberc Lung Dis* 1997;1(3):276-283.

205. Iseman MD. Directly-observed therapy, patient education and combined drug formulations: complementary, not alternative, strategies in tuberculosis control [editorial]. *Tuber Lung Dis* 1996;77:101.
206. World Health Organisation. Breakthrough in TB control announced by the WHO. Press Release WHO/23, 19th March 1997.
207. Volmink J, Garner P. DOTS - are we over-optimistic? [correspondence]. *S Afr Med J* 1997;87:905-6.
208. Advisory Council for the Elimination of Tuberculosis. Initial therapy for tuberculosis in the era of multidrug resistance; recommendations of the Advisory Council for the Elimination of Tuberculosis. *Morb Mortal Wkly Rep* 1993;42:rr77.
209. Nagpaul DR, Vishwanath MK, Dwarakanath G. A Socio-epidemiological Study of Out-patients Attending a City Tuberculosis Clinic in India to Judge the Place of Specialised Centres in a Tuberculosis Control Programme. *Bull World Health Organ* 1970;43:17-34.
210. Paramasivam R, Rajasekaran R, Selvaraj D. Evolution of admission policy in a tuberculosis hospital - a suggested model. *Lung India* 1992;X(3):110-114.
211. Chaulet P. Compliance with chemotherapy for tuberculosis. Responsibilities of the Health Ministry and of physicians. *Bull Int Union Tuberc Lung Dis* 1990/1991;66(Suppl):33-35.
212. Fox W. Compliance of patients and physicians: experience and lessons from tuberculosis-II. *Br Med J* 1983;287:101-105
213. Hong YP, Kwon DW, Kim SJ, Chang SC, Kang MK, Lee EP et al Survey of knowledge, attitudes and practices for tuberculosis among general practitioners. *Tuber Lung Dis* 1995;76:431-435
214. Marsh D, Hashim R, Hassany F, Hussain N, Iqbal Z, Irfanullah A et al Front-line management of pulmonary tuberculosis: an analysis of tuberculosis and treatment practices in urban Sindh, Pakistan *Tuber Lung Dis* 1996;77:86-92.
215. Allan WGL, Girling DJ, Fayers PM, Fox W. The symptoms of newly diagnosed pulmonary tuberculosis and patients' attitudes to the disease and to its treatment in Hong Kong. *Tubercle* 1979;60:211-223.
216. Rothe TB, Karrer W. Short-course therapy of pulmonary tuberculosis: doctors' compliance. *Tuber Lung Dis* 1996;77:93-97.
217. Menzies D, Adhikari N, Tannenbaum T. Patient characteristics associated with failure of tuberculosis prevention. *Tuber Lung Dis* 1996;77:308-314.

218. Ormerod LP, Prescott RJ. Inter-relations between relapses, drug regimens and compliance with treatment in tuberculosis. *Respir Med* 1991;85:239-242.
219. Sbarbaro JA. A challenge - to our practices and to our principles. [editorial] *Tuber Lung Dis* 1996;77:2-3.
220. Raviglione MC, Dye C, Schmidt S, Kochi A. Assessment of worldwide tuberculosis control. *Lancet* 1997;350:624-629.
221. Graf P. Tuberculosis control in high-prevalence countries. In: Davies PDO, editor. *Clinical Tuberculosis*. London: Chapman & Hall, 1994:325-339.
222. Chum HJ. The Tanzania National Tuberculosis/Leprosy Programme in the face of HIV infection. *Bull Int Union Tuberc Lung Dis* 1990/1991;66 Suppl:53-55
223. China Tuberculosis Control Collaboration. Results of directly observed short-course chemotherapy in 112 842 Chinese patients with smear-positive tuberculosis. *Lancet* 1996;347:358-362.
224. Salomao MA, Parkkali. Case-finding and treatment results of pulmonary tuberculosis in Mozambique, 1985-1988. *Bull Int Union Tuberc Lung Dis* 1990/1991;66 Suppl:47-49.
225. Chan SL, Wong PC, Tam CM. 4-, 5- and 6-month regimens containing isoniazid, rifampicin, pyrazinamide and streptomycin for treatment of pulmonary tuberculosis under programme conditions in Hong Kong. *Tuber Lung Dis* 1994;75:245-250.
226. Nyangulu DS, Nkhoma WN, Salaniponi. Factors contributing to a successful Tuberculosis Control Programme in Malawi. *Bull Int Union Tuberc Lung Dis* 1990/1991;66 Suppl:45-46.
227. World Health Organisation Report, Weil DC. Tuberculosis Control Programmes in Brazil. Tuberculosis Unit WHO, 1990
228. Kumaresan JA, Maganu ET. Case holding in patients with tuberculosis in Botswana. *Br Med J*;305:340-341.
229. Sukrakanchana-Trikham P, Puechal X, Rigal J, Rieder HL. 10-year assessment of treatment outcome among Cambodian refugees with smear-positive tuberculosis in Khao-I-Dang, Thailand. *Tuber Lung Dis* 1992;73:384-387.
230. Gninafon M. The Anti-tuberculosis Programme of Benin. *Bull Int Union Tuberc Lung Dis* 1990/1991;66 Suppl:57-58.
231. Arguello L. Results of the tuberculosis programme in Nicaragua in 1984-89. *Bull Int Union Tuberc Lung Dis* 1990/1991;66 Suppl:51-52.

232. Caselle MT, Galvagno S. Tuberculosis in a Rural Hospital in Northern Kenya. *E Afr Med J* 1992;69:464-467.
233. Mushtaque A, Chowdhury R. Controlling a forgotten disease: the case of tuberculosis in a primary health care setting in rural Bangladesh. Proceeding of the Workshop on case studies for evaluation of Tuberculosis Control programmes in various primary care settings; 1990 May 25-28; Sturbridge, Massachusetts, USA: WHO, 1990.
234. Datta M, Radhamani MP, Selvaraj R, Paramasivan CN, Gopalan BN, Sudeendra CR, et al. Critical assessment of smear-positive pulmonary tuberculosis patients after chemotherapy under the district tuberculosis programme. *Tuber Lung Dis* 1993;74:180-186.
235. Hopewell PC, Sanchez-Hernandez M, Baron R, Ganter B. Operational Evaluation of Treatment for Tuberculosis: results of a "standard" 12-month Regimen in Peru. *Am Rev Respir Dis* 1984;129:439-443.
236. Manalo F, Tan F, Sbararo JA, Iseman MD. Community-based Short-course Treatment of Pulmonary Tuberculosis in a developing Nation. *Am Rev Respir Dis* 1990;142:1301-1305.
237. Williams JC, Nsinga M, Mulumba K. Fate of smear-positive pulmonary tuberculosis patients after 12 months on an entirely ambulatory primary treatment in Kinshasa (Zaire). *Bull Int Union Tuberc Lung Dis* 1986;61:43-45.
238. Grzybowski S. Drugs are not enough [editorial]. *Tuber Lung Dis* 1993;74:145-146.
239. Gangadharam PRJ. Chemotherapy of tuberculosis under programme conditions. *Tuber Lung Dis* 1994;75:241-244.
240. Chakraborty AK [correspondence]. *Tuber Lung Dis* 1994;75:316-320.
241. De Cock KM, Wilkinson D. Tuberculosis control in resource-poor countries: alternative approached in the era of HIV. *Lancet* 1995;346:675-677.
242. Wilkinson D, Bechan S, Connolly C, Standing E, Short G. Should we take a history of prior treatment, and check sputum status at 2-3 months when treatment patients for tuberculosis? *Int J Tuberc Lung Dis* 1998;2:52-55.
243. Trebucq A, Rieder HL. Two excellent management tools for national tuberculosis programme: history of prior treatment and sputum status at two months. *Int J Tuberc Lung Dis* 1998;2(3):184-186.
244. Helberg H Tuberculosis programme: fragmentation or integration? [Editorial] *Tuber Lung Dis* 1995;76:1-3

245. Fox W. Compliance of patients and physicians: experience and lessons from tuberculosis-I. *Br Med J* 1983; 287:33-35
246. Rouillon A. Problems in organising effective ambulatory treatment of tuberculosis patients. *Bull Int Union Tuberc Lung Disease* 1972;47:68-83
247. Rubel AJ, Garro LC. Social and Cultural Factors in the Successful Control of Tuberculosis. *Public Health Rep* 1992;107(6):626-636.
248. Grange JM, Festenstein F. The Human dimension of tuberculosis control. *Tuber Lung Dis* 1993;74:219-222.
249. Sumartojo E. When Tuberculosis Treatment Fails: a Social Behavioral Account of Patient Adherence. *Am Rev Respir Dis* 1993;147:1311-1320.
250. Haynes RB. Introduction. In: Haynes RB, Taylor DW, Sackett DL, editors. *Compliance in Health Care*. Baltimore: Johns Hopkins University Press, 1979;1-7.
251. Menzies R, Rocher I, Vissandjee B. Factors associated with compliance in treatment of tuberculosis. *Tuber Lung Dis* 1993;74:32-37
252. Wardman AG, Knox AJ, Muers MF, Page RL. Profiles of Non-Compliance with Antituberculosis Therapy. *Br J Dis Chest* 1988;82:285-289.
253. Pozsik CJ. Compliance with tuberculosis therapy. *Med clin North Am* 1993;77(6):1289-1301.
254. Becker MH, Maiman LA. Sociobehavioral Determinants of Compliance with Health and Medical Care Recommendations. *Med Care* 1975;13(1):10-24.
255. Williams B. Patient Satisfaction: a valid concept. *Soc Sci Med* 1994;38(4):509-516.
256. Jenkins CD. Group Differences in Perception: A Study of Community Beliefs and Feelings about Tuberculosis. *Am J Sociol* 1966;71:417-429.
257. Johansson E, Diwan VK, Huong ND, Ahlberg BM. Staff and patient attitudes to tuberculosis and compliance with treatment: an exploratory study in a district in Vietnam. *Tuber Lung Dis* 1996;77:178-183.
258. Dick J, Van der Walt H, Hoogendoorn L, Tobias B. Development of a health education booklet to enhance adherence to tuberculosis treatment. *Tuber Lung Dis* 1996;77:173-177
259. Cassell J. A comprehensive health program among South African Zulus. In: *Health, culture, and community*. editor Paul B. Russel Sage Foundation, New York, 1995:15-41

260. Metcalf CA, Bradshaw D, Stindt WW. Knowledge and beliefs about tuberculosis among non-working women in Ravensmead, Cape Town. *S Afr Med J* 1990;77:408-411.
261. Liefoghe R, Michiels N, Habib S, Moran MB, De Muynck A. Perception and Social Consequence of Tuberculosis: A focus group study of tuberculosis patients in Sialkot, Pakistan. *Soc Sci Med* 1995;41:1685-1692.
262. Cornwall J. Tuberculosis: a clinical problem of international importance. *Lancet* 1997;349:660-661.
263. Westaway MS. Knowledge and attitudes about tuberculosis of black hospitalised TB patients. *Tubercle* 1990;71:55-59.
264. Nichter M. Illness Semantics and International Health: the weak lungs/TB complex in the Phillipines. *Soc Sci Med* 1994;38(5):649-663.
265. Emmel ND, O'Keefe P. Participatory analysis for redefining health delivery in a Bombay slum. *J Public Health Med* 1996;18(3):301-307
266. Carey JW, Oxtoby MJ, Nguyen LP, Huynh V, Morgan M, Jeffery M. Tuberculosis Beliefs Among Recent Vietnamese Refugees in New York State. *Public Health Rep* 1997;112:66-72.
267. Molantoa KEM. Traditional attitudes towards tuberculosis. *S Afr Med J* 1982; Special Issue 17th November:29-31.
268. Westaway MS, Wolmarans L. Cognitive and affective reactions of black urban South Africans towards tuberculosis. *Tuber Lung Dis* 1994;75:447-453.
269. Ndeti K. Sociocultural aspects of tuberculosis defaultation: a case study. *Soc Sci Med* 1972;6:397-412.
270. Barnhoorn F, Adriaanse H. In search of factors responsible for noncompliance among tuberculosis patients in Warda district, India. *Soc Sci Med* 1992;34(2):291-306.
271. Farmer P. Social scientists and the new tuberculosis. *Soc Sci Med* 1997;44(3):347-358.
272. Thomson EM, Myrdal S. Tuberculosis-the patients' perspective. *S Afr Med J* 1986;70:263-264.
273. Westaway MS, Wessie GM. Tuberculosis diagnosis and treatment of young South African children: experiences and perceptions of care-givers. *Tuber Lung Dis* 1994;75:70-74.

274. Buchanan N, Shuenyane E, Mashigo S, Mtangai P, Unterhalter B. Factors Influencing Drug Compliance in Ambulatory Black Urban Patients. *S Afr Med J* 1979;55:368-373.
275. Morisky DE, Malotte CK, Choi P, Davidson P, Rigler S, Sugland B et al. A Patients Education Program to Improve Adherence Rates with Antituberculosis drug regimens. *Health Educ Q* 1990;17(3):253-267.
276. Cuneo WD, Snider DE Jr. Enhancing Patient Compliance with Tuberculosis Therapy. *Clin Chest Med* 1989;10(3):375-379.
277. Liefvooghe R, Baliddawa JB, Kipruto EM, Vermeire C, De Munynck AO. From their own perspective. A Kenyan community's perception of tuberculosis. *Trop Med Int Health* 1997;2(8):809-821.
278. Ngubane H. Aspects of Clinical Practice and Traditional Organisation of Indigenous Healers in South Africa. *Soc Sci Med* 1981;15B:3.
279. McDonald SJ, Ma M. Identifying Compliance Incentives for Screening and Treatment of Tuberculosis. *J Community Health Nurs* 1987;4(3):131-143.
280. Sbarbaro JA. Strategies to improve compliance with therapy. *Am J Med* 1985;79 Suppl 6A:34-37.
281. Edginton ME. Tuberculosis: the need for bio-social care. *S Afr Med J* [editorial] 1997;87(8):1041.
282. Ellis JHP, Beyers N, Bester D, Gie RP, Donald PR. Sociological anthropological factors related to the community management of tuberculosis in the Western Cape communities of Ravensmead and Uitsig. *S Afr Med J* 1997;87:1047-1051.
283. Kimerling ME, Petri L. Tracing as part of tuberculosis control in a rural Cambodian District during 1992. *Tuber Lung Dis* 1995;76:156-159.
284. Wardman AG, Knox AJ, Muers MF, Page RL. Profiles of Non-Compliance with Antituberculosis Therapy. *Br J Dis Chest* 1988;82:285-289.
285. Dick J, Schoeman HH. Tuberculosis in the Community: 2. The perceptions of members of a tuberculosis health team towards a voluntary health worker programme. *Tuber Lung Dis* 1996;77:380-383.
286. Mellins RB, Zimmerman B, Clark NM. Patient Compliance: Are We Wasting Our Time and Don't Know It? [editorial] *Am Rev Respir Dis* 1992;146:1376-1377.
287. Dick J, Clarke M, Tibbs J, Schoeman H. Combating tuberculosis - lessons learned from a rural community project in the Klein Drakenstein area of the Western Cape. *S Afr Med J* 1997;87:1042-1047.

288. Werhane MJ, Snukst-Torbeck G, Schraufnagel DE. The Tuberculosis Clinic. *Chest* 1989;96(4):815-818.
289. Rideout M, Menzies R. Factors affecting compliance with preventive treatment for tuberculosis at Mistassini Lake, Quebec, Canada. *Clin Invest Med* 1994;17(1):31-36.
290. Buri PS, Vathesatogkit P, Charoenpan P, Kiatboonsri S, Buranaratchada S. A Clinic Model for a Better Tuberculosis Treatment Outcome and Factors Influencing Compliance. *J Med Assoc Thai* 1985;68(7):356-360.
291. Smukst-Torbeck G, Werhane MJ, Schraufnagel DE. Treatment of tuberculosis in a nurse-managed clinic. *Heart Lung* 1987;16(1):30-33.
292. Ramachandran P, Prabhakar R. Defaults, defaulter action and retrieval of patients during studies on tuberculosis meningitis in children. *Tuber Lung Dis* 1992;73:170-173.
293. Jin BW, Kim SC, Shimao T. The impact of intensified supervisory activities on tuberculosis treatment. *Tuber Lung Dis* 1993;74:267-272.
294. van der Werf TS, Dade GK, van der Mark TW. Patient compliance with tuberculosis treatment in Ghana: factors influencing adherence to therapy in a rural service programme. *Tubercle* 1990;71:247-252.
295. Teklu B. Reasons for failure in treatment of pulmonary tuberculosis in Ethiopians. *Tubercle* 1984;65:17-21.
296. Miles SH, Maat RB. A Successful Supervised Outpatient Short-Course Tuberculosis Treatment Program in an Open Refugee Camp on the Thai-Cambodian Border. *Am Rev Respir Dis* 1984;130:827-830.
297. Haynes RB. Determinants of Compliance: The Disease and the Mechanics of Treatment In: Haynes RB, Taylor DW, Sackett DL, editors. *Compliance in Health Care*. Baltimore: Johns Hopkins University Press, 1979;49-62.
298. Curry FJ. Neighborhood clinics for more effective outpatient treatment of tuberculosis. *N Engl J Med* 1968;279:1262-1267.
299. Hill JP, Ramachandran. A simple scheme to improve compliance in patients taking tuberculosis medication. *Trop Doct* 1992;22:161-163.
300. Giuffrida A, Torgerson DJ. Should we pay the patient? Review of financial incentives to enhance patient compliance. *Br Med J* 1997;315:703-707.
301. Collins TBF The history of southern Africa's first tuberculosis epidemic. *S Afr Med J* 1982;62:780-788.

302. Packard RM. White Plague, Black Labour. Pietermaritzburg: University of Natal Press, 1989.
303. Metcalf C. A history of tuberculosis. in: Coovadia HM, Benatar SR. A century of Tuberculosis. South African perspectives. Oxford: Oxford University Press 1991.
304. Dubovsky H. Rural Tuberculosis in the Republic of South Africa. Public Health 1987;101:85-91.
305. Benatar SR. Failure of the tuberculosis control in South Africa - the need for a unitary national health system. [editorial] S Afr Med J 1986;70:247-248.
306. Report of the Tuberculosis Research Committee 1932; Tuberculosis in South African natives, with special reference to the disease among the mine labourers on the Witwatersrand. Johannesburg: South African Institute for Medical Research.
307. Stevenson-Hamilton J. The Low-veld: its wildlife and its people. London, Toronto, Melbourne and Sydney. Cassell and Company, Ltd, 1929.
308. Schneider J. Tuberculin testing and Mass Miniature X-Ray Survey of the Northern and Eastern Transvaal. S Afr Med J. 1954;28:689-692.
309. Department of Health. Tuberculosis update. Epidemiological Comments 1995;22(1).
310. Department of Health. Tuberculosis Control Programme 1992. Epidemiological Comments 1994;21(3).
311. TB in South Africa The People's Plague. The Department of Health 1997.
312. Weyer K, Fourie PB. Assessment of the tuberculosis epidemic in South Africa: Historical perspective and critical evaluation of current information. 1996. Report for the Tuberculosis Control Programme Review Task Group. National Tuberculosis Research Programme, Medical Research Council, Pretoria.
313. Department of Health. Tuberculosis Control Programme 1991. Epidemiological Comments 1993;20(1):2-11
314. Weyer K, Kleeberg HH. Primary and acquired drug resistance in adult black patients with tuberculosis in South Africa: results from a continuous national drug resistance programme involvement. Tubercle Lung Dis 1992;73:106-112.
315. Strebel PM, Seager JR. The epidemiology of tuberculosis in South Africa. In: Coovadia HN, Benatar SR, editors. A Century of Tuberculosis: South African Perspectives. Oxford: Oxford University Press, 1991;58-0.
316. Botha JL, Bradshaw D. African vital statistics - a black hole? S Afr Med J 1985; 67: 977-981.

317. Weyer K. Tuberculosis in rural populations in Southern Africa: the association between sputum origin, sputum quality and bacteriological prevalence. 1990 Doctoral thesis, University of Pretoria.
318. Fourie PB, Knoetze K. Tuberculosis prevalence and risk of infection in Southern Africa. *S Afr J Sci* 1986;82:287.
319. Berman S, Kibel MA, Fourie P, Strebel PM. Childhood tuberculosis and tuberculosis meningitis: high incidence rates in the Western Cape. *Tubercle Lung Dis* 1992;73:349-355.
320. Styblo K. Recent advances in epidemiological research in tuberculosis. *Adv Tuberc Res* 1980;20:1-63.
321. Styblo K. Surveillance of tuberculosis. *Int J Epidemiol* 1976;5:63-68.
322. Department of Health. Further hypotheses on the tuberculosis epidemic in South Africa. *Epidemiological Comments* 1993;20(7):106-118.
323. Department of Health. Policy Statement: Tuberculosis Control in the Republic of South Africa. Pretoria: Department of Health, 1979.
324. Tibbit LR. The economics of short-course therapy for tuberculosis in the Cape Divisional Council area. *S Afr Med J* 1982; Special Issue 17th November:31-33.
325. Glatthaar E. Tuberculosis: Basic Perspectives. Pretoria: Medunsa 1982.
326. Tuberculosis Control Programme: Strategy and Policy Manual 1992. Department of National Health and Population Development.
327. Lee T, Buch E. Tuberculosis Control in South Africa. In Coovardia HM, Benatar SR, editors. *A Century of Tuberculosis: South African Perspectives*. Cape Town: Oxford University Press, 1991:287-302.
328. Glatthaar E. Tuberculosis control in South Africa: where have we gone wrong? *S Afr Med J*. 1982; Special Issue 17th November:36-41.
329. Thomson EM, Myrdal S. Regional variations in tuberculosis policy in the Cape and Ciskei. *S Afr Med J*. 1986;70:253-257.
330. Yach D, Hoffman M, van Herzeele A. Western Cape local authority compliance with tuberculosis policy, 1984. *S Afr Med J* 1988;73:33-35.
331. Koornhof HJ, Coetzee GJ. A National policy for the laboratory diagnosis of pulmonary tuberculosis [editorial]. *South Afr J Epidemiol Infect* 1995;10:94.
332. Benatar SR. Tuberculosis in the 1980s, with particular reference to South Africa. *S Afr Med J* 1982;62:359-364.

333. Taylor SP, Benatar SR. The Tuberculosis Control Programme - a time to re-evaluate? [editorial] *S Afr Med J* 1989;76:639-640.
334. La Grange MAC, Glatthaar E. The tuberculosis epidemic - why we are slipping up. *S Afr Med J* 1991;80:329-331.
335. Westaway MS, Fourie PB. Tuberculosis control. *Contin Med Educ* 1989;7:305-311.
336. Yach D. Tuberculosis in the Western Cape Health Region of South Africa. *Soc Sci Med*; 1988;27:683-689.
337. Styblo K. Report on an evaluation of the South African Tuberculosis Control Programme. March 1994. South African Department of Health. Pretoria.
338. Graf P, Chaulet P, Eriki P. Report on a Mission to South Africa, July 1995. World Health Organisation.
339. Lee T, Price M. Indicators and research methods for rapid assessment of a tuberculosis control programme: case study of a rural area in South Africa. *Tuber Lung Dis* 1995;76:441-449.
340. Arnadottir The. Assessment of tuberculosis: is 'rapid assessment' needed? *Tuber Lung Dis* 1995;76:375-376.
341. Burney P, Shahyar S. Tuberculosis in the Transkei. In: Wilson F, Westcott G, editors. *Economics of Health in South Africa. Volume II: Hunger, Work and Health*. Johannesburg: Raven Press, 1980.
342. Whitehouse AB. The Modern Management of Tuberculosis in a Rural Area. *S Afr Med J* 1980;58:695-696.
343. Saunders LD, Irwig LM, Wilson TD, Kahn A, Groenewald H. Tuberculosis management in Soweto. *S Afr Med J* 1984;66:330-333.
344. Griffiths ML, Makgothi MM, Nordesjö G. Tuberculosis management in a rural community - factors in failure. *S Afr Med J* 1981;59:14-16.
345. Eason C, Johnson K, Mashabane R. Can good tuberculosis care be provided in the face of poverty? Paper presented to the Second Carnegie Inquiry into Poverty and Development in Southern Africa. Cape Town: SALDRU, University of Cape Town. 1984.
346. Fredlund VG. Six-month intermittent chemotherapy for tuberculosis in the Mseleni Health Ward of KwaZulu. *S Afr Med J* 1990;77:405-407.
347. Conradie HH. Compliance to TB outpatient treatment in the Hewu district of Ciskei. Is hospitalisation necessary? *SA Fam Prac* 1986;7:356-361.

348. Yeats JR. Attendance compliance for short-course tuberculosis chemotherapy at clinics in Estcourt and surroundings. *S Afr Med J* 1986;70:265-266.
349. Collie A, Küstner HGV. The Tuberculosis Control Programme, 1985-1986. *S Afr Med J* 1989;76:675-680.
350. Westaway MJ. Managing tuberculosis at the clinic level. [correspondence]. *S Afr Med J* 1990;77:539-540.
351. Bell J, Yach D. Tuberculosis patient compliance in the western Cape, 1984. *S Afr Med J* 1988;73:31-33.
352. Westaway MS, Conradie PW, Remmers L. Supervised out-patient treatment of tuberculosis: evaluation of a South African rural programme. *Tubercle* 1991;72:140-144.
353. Edginton ME, Price M. A record review of patients notified with TB to an urban local authority. Conference paper at the Annual Conference of the Epidemiological Society of Southern Africa, Johannesburg 1991.
354. Wilkinson D. High-compliance tuberculosis treatment programme in a rural community. *Lancet* 1994;343:647-648.
355. Dick T, Schoeman JH, Mohammed A, Lombard C. Tuberculosis in the community: 1. Evaluation of a volunteer health worker programme to enhance adherence to anti-tuberculosis treatment. *Tuber Lung Dis* 1996;77:274-279.
356. Tuberculosis Control in South Africa: Joint Programme Review, June 1996. WHO, Geneva. WHO/TB/96.208.
357. Millard FJC. Assessment of resources for tuberculosis control on Sekukhuneland, Northern Province. *S Afr J Epidemiol Infect* 1997;12(3):75-78.
358. Wilkinson D, de Cock KM. Tuberculosis control in South Africa - time for a new paradigm? [editorial] *S Afr Med J* 1996;86:33-35.
359. TB talk. National Tuberculosis Control Programme, Department of Health. August 1997 and September/October 1997.
360. Notifiable Medical Conditions. Epidemiological Comments, Department of Health 1996/7;23:24-25.
361. World Health Organisation, United Nations Children's Fund. Primary Health Care. Report of the International Conference on Primary Health Care. Alma Ata, USSR, 6-12 September 1978. Geneva: World Health Organisation, 1978.
362. Newell K. The way ahead for district health systems. *World Health Forum*, 1989;10:80-87.

363. World Health Organisation. The Challenge of Implementation. District Health Systems for Primary Health Care. World Health Organisation, Division of Strengthening Health Services, 1988.
364. Kasongo Project Team. Primary care for less than a dollar a year. World Health Forum. 1984;5:211-215.
365. Amonoo-Lartson R, Ebrahim GJ, Lovel HJ, Ranken JP. District Health Care: Challenges for Planning, Organisation and Evaluation in Developing Countries. Basingstoke and London: MacMillan Education Ltd, 1984.
366. World Health Organisation. Integration of Health Care Delivery. WHO Technical Report Series No. 861, Geneva, 1996.
367. Tarimo E. Towards a health district: Organising and managing district health systems based on primary health care. World Health Organisation, Geneva. 1991.
368. Daniel TM. Selective Primary Health Care: Strategies for Control of Disease in the Developing World.II Tuberculosis. Rev Infect Dis 1982;4(6):1254-1265.
369. Leowski J. Tuberculosis control at the Primary Health Care level. Bull Int Union Tuberc 1986;61:29-31.
370. Chaulet P. Integrating tuberculosis control activities into the primary health care system at district level Bull Intern Union Tuberc Lung Dis 1989;64:33-34.
371. Zidouni N. Involvement of a properly structured network of primary health care in the management of tuberculous patients during a community survey in 45 districts in Algeria. Bull Intern Union Tuberc 1989;64:36-38.
372. Gokce C, Gokce O, Erdogmus Z, Arisoy E, Arisoy S, Koldas O et al Problems in running a tuberculosis dispensary in a developing country: Turkey Tubercle 1991;72:268-276.
373. Huwaha FN. The integration of tuberculosis control into primary health care in Uganda during this era of the human immune deficiency virus epidemic CHASA J Comprehen Health 1995;6(1):46-48.
374. Tuberculosis Control as an Integral Part of Primary Health Care. WHO. Geneva;1988.
375. Zwarenstein M, Barron P, Tollman S, Crisp N, Frankish J, Toms I, Solarsh G. Primary Health Care depends on the district system. [editorial] S Afr Med J 1993;83:558.
376. Department of Health. Health Policy Co-ordinating Unit. A policy for the development of a district health system in South Africa. 1994.

377. Pick WM. Alternative approaches to the delivery of health care in South Africa. *S Afr J Sci* 1996;92:17-23.
378. Gilson L, Balfour T, Goosen V. Towards Well-Functioning Health Districts in South Africa: A Vision and Indicators for Assessing Progress. Technical Report No. TK27. 1997. The Centre for Health Policy, Department of Community Health, University of the Witwatersrand.
379. Harrison D. A Pocket Guide to District Health Care in South Africa. 1997. Health Systems Trust.
380. World Health Organisation. The Health Centre in District Health Systems. Division of Strengthening Health Services, World Health Organisation, Geneva. 1994.
381. Yach D, Tollman SM. Public Health Initiatives in South Africa in the 1940s and 1950s: Lessons for a Post-Apartheid era. *Am J Public Health* 1993;83:1043-1050.
382. Tollman SM, Rispel L. Organisation, Planning and Management. In: South African Health Review. Health Systems Trust and Henry J Kaiser Family Foundation. Durban and California, 1995;75-88.
383. Tollman S, Mkhabela S, Pienaar J. Developing District Health Systems in the rural Transvaal: Issues based on the Tintswalo/Bushbuckridge experience. *S Afr Med J* 1993;83:565-568.
384. Tyrell B, Jergens P. African Heritage. Johannesburg: MacMillan South Africa Pty Ltd, 1983.
385. Pugh AO. The District Health System. Paper presented at the District Health Systems Workshop, 30th September and 1st October 1992.
386. Project for Statistics on living standards and development. South Africa's rich and poor: baseline household statistics. SALDRU, School of Economics, University of Cape Town. August 1994.
387. Coleman I, Dolan C, Russo R. The Village Development Programme Income and Expenditure: an analysis. Unpublished report 1993; Wits Rural Facility, University of the Witwatersrand, Klaserie.
388. Hirschowitz R, Orkin M. A National Household Survey of Health Inequalities in South Africa. Prepared by The Community Agency for Social Enquiry (CASE). The Henry J Kaiser Family Foundation, October 1995.
389. Department of Health. Notifiable medical conditions: number of reported cases of selected conditions, January to December 1992. *Epidemiological Comments*. 1993;20(6):103.

390. Department of Health. The trend of Tuberculosis through the eyes of cohorts . *Epidemiological Comments*. 1992;19(11):184-197.
391. Chum HJ. Ten years of the National Tuberculosis/Leprosy Programme in Tanzania. *Bull Int Union Tuberc Lung Dis* 1989; 64: 34-36.
392. Holmes CB, Hausler H, Nunn P. A review of differences in the epidemiology of tuberculosis. *Int J Tuberc Lung Dis* 1998;2(2):96-104.
393. Department of Health. National antenatal surveillance. *Epidemiological Comments* 1995;22(5):90-96.
394. Wilkinson D. Tuberculosis: Hlabisa Health Ward. Department of Health, *Epidemiological Comments*. 1993;20:73-75.
395. Xie HJ, Enarson DA, Chao CW, Allen FA, Grzybowski S. Deaths in tuberculosis patients in British Columbia, 1980-1984. *Tuber Lung Dis* 1992;73:77-82.
396. Weyer K, Groeneveld PJ, Zwarenstein M, Lombard CJ. Tuberculosis drug resistance in the Western Cape. *S Afr Med J* 1995;85(6):499-504.
397. Jolly R, King M. The organisation of health services. In King M, editor. *Medical Care in Developing Countries*. Lusaka, Addis Ababa, London: Oxford University Press, 1966:2.1-2.15.
398. Okot-Nwang M, Wabire-Mangen F, Kagezi VBA. Increasing prevalence of tuberculosis among Mulaga Hospital admissions, Kampala, Uganda (1985-1989). *Tuber Lung Dis* 1993; 74: 121-125.
399. Aisu T, Nochur SV. Monitoring of Tuberculosis Therapy Compliance. Abstracts for the Annual Meeting of the American Society of Microbiology, 16-20 May, 1993, Atlanta, Georgia.
400. Sumartojo E. Comparison of Measures of Adherence to TB treatment. Unpublished manuscript. Division of Tuberculosis Elimination, National Center for HIV, STD and TB Prevention. Centers for Disease Control and Prevention. Atlanta, Georgia.1996.
401. Moorman J, Edginton ME. Cause of death of patients on treatment for tuberculosis: a study at Tintswalo Hospital. Submitted to *Int J Tuberc Lung Dis*, June 1998.
402. Khan ME, Anker M, Patel BC, Barge S, Sadihwani H, Kohle R. The use of focus groups in social and behavioural research: some methodological issues. *World Health Stat Q* 1991;44:145-149.
403. Kitzinger J Introducing focus groups. *Br Med J* 1995;311:299-302.

404. Centre for Disease Control and Prevention. Improving Tuberculosis Treatment and Control: an Agenda for Behavioral, Social and Health Services Research. Proceedings :Tuberculosis and Behavior. National; Workshop on Research for the 21st Century. Atlanta; CDC. 1995.
405. Sutter EE, Ballard RC. Community participation in the control of trachoma in Gazankulu. *Soc Sci Med* 1983;17(22):1813-1817.
406. Dujardin B, Kegels G, Buve A, Mercenier P. Tuberculosis control: did the programme fail or did we fail the programme? [editorial] *Tropical Med and Int Health* 1997;2(8):715-718.
407. Herzberg F, Mausner, Snyderman BB. *The Motivation to Work*. New York: John Wiley & Sons, 1959.
408. Beattie A, Rispel L, Broomberg J, Price M, Cabral J. The Quality of Primary Care in South Africa: the results of a facility-based assessment. Centre for Health Policy Paper no.44, Department of Community Health, University of the Witwatersrand. September 1995.
409. Maslow AD. A Theory of Human Motivation. *Psychological Review* 1943;July:370-396.
410. Improving the performance of health centres in district health systems. Report of a WHO Study Group. WHO Technical Report Series 869. Geneva 1997.

APPENDIX A: TABLES

A1 TREATMENT OUTCOMES AND SEX - NEW SMEAR POSITIVE PATIENTS (1992-94)

SEX	COMPLETED	INTERRUPTED	DIED	TOTAL
Female	42 (66%)	16 (25%)	6 (9%)	64 (31%)
Male	97 (68%)	36 (25%)	10 (7%)	143 (69%)
Total	139 (67%)	52 (25%)	16 (8%)	207 (100%)

A2 TREATMENT OUTCOMES AND AGE GROUPS - NEW SMEAR POSITIVE CASES (1992-94)

AGE GROUP in yrs	COMPLETED	INTERRUPTED	DIED	TOTAL
0-14	3 (60%)	1 (20%)	1 (20%)	5 (2%)
15-29	40 (66%)	17 (28%)	4 (7%)	61 (29%)
30-59	75 (70%)	25 (23%)	7 (7%)	107 (52%)
60 +	21 (62%)	9 (26%)	4 (12%)	34 (16%)
Total	164 (79%)	65 (31%)	19 (9%)	207 * (100%)

* excludes 3 transferred cases

A3 TREATMENT OUTCOMES AND DISTANCE FROM TINTSWALO HOSPITAL, NEW SMEAR POSITIVE PATIENTS (1992-94)

DISTANCE in kms	COMPLETED	INTERRUPTED	DIED	TOTAL
0-9	49 (65%)	21 (28%)	5 (7%)	75 36%
10-25	62 (76%)	15 (18%)	5 (6%)	82 40%
26-49	19 (61%)	7 (23%)	5 (16%)	31 15%
50 +	8 (44%)	9 (50%)	1 (6%)	18 9%
TOTAL	138 (67%)	52 (25%)	16 (8%)	206 100%

A4 DISTANCE FROM TINTSWALO HOSPITAL AND AGE GROUPS (1994-96)

AGEGROUP (years)	DISTANCE CATEGORIES (kms)				TOTAL
	0 -9	10-25	26-49	50+	
0-14	17 (37%) 12%	9 (20%) 8%	19 (41%) 15%	1 (2%) 4%	46 11%
15-29	27 (30%) 19%	26 (29%) 22%	32 (36%) 25%	4 (4%) 15%	89 21%
30-59	84 (37%) 58%	68 (30%) 58%	61 (27%) 48%	17 (7%) 63%	230 56%
60+	16 (33%) 11%	14 (29%) 12%	14 (29%) 11%	5 (10%) 19%	49 12%
Total	144 (35%)	117 (28%)	126 (30%)	27 (7%)	414

**A5 DISTANCE FROM TINTSWALO HOSPITAL
AND SFX (1994-96)**

DISTANCE (kms)	FEMALE	MALE	TOTAL
0-9	38 (26%) 31%	106 (74%) 36%	144 (100%)
10-25	34 (29%) 28%	83 (71%) 28%	117 (100%)
26-49	45 (36%) 37%	81 (64%) 28%	126 (100%)
50+	5 (19%) 4%	22 (81%) 8%	27 (100%)
TOTAL	122 (29%) 100%	292 (71%) 100%	414 (100%)

**A6 DISTANCE FROM TINTSWALO AND
PATIENT CATEGORY (1994-96)**

DISTANCE (kms)	NEW PATIENTS	RE- TREATMENT PATIENTS	TOTAL
0-9	129 (90%) 37%	15 (10%) 24%	144 (100%)
10-25	92 (79%) 26%	25 (21%) 40%	117 (100%)
26-49	104 (83%) 30%	22 (17%) 35%	126 (100%)
50 +	26 (96%) 7%	1 (4%) 2%	27 (100%)
TOTAL	351 (85%) 100%	63 (15%) 101%	414

A7 AGES AND TREATMENT GROUPS

Age Group	TREATMENT GROUP				TOTAL
	Treatment Clinic	Control Clinic	Hospital	Outside District	
0-14	16 (35%) 12%	12 (26%) 16%	10 (22%) 8%	8 (17%) 9%	46 (11%)
15-29	22 (26%) 17%	19 (21%) 25%	26 (29%) 22%	22 (25%) 25%	89 (21%)
30-59	79 (34%) 60%	36 (16%) 48%	70 (30%) 58%	45 (20%) 52%	230 (56%)
60+	15 (31%) 11%	8 (16%) 11%	14 (29%) 12%	12 (24%) 14%	49 (12%)
TOTAL	132 32%	75 18%	120 29%	87 21%	414 100%

A8 SEX AND TREATMENT GROUPS

Sex	TREATMENT GROUP				TOTAL
	Treatment Clinic	Control Clinic	Hospital	Outside District	
FEMALE	37 (30%) 28%	21 (17%) 28%	34 (28%) 28%	30 (25%) 34%	122 (100%) 29%
MALE	95 (33%) 72%	54 (18%) 72%	86 (29%) 72%	57 (20%) 66%	292 (100%) 71%
TOTAL	132 (32%) 100%	75 (18%) 100%	120 (29%) 100%	87 (21%) 100%	414 (100%) 100%

A9 AGES AND TREATMENT SUPPORTERS

AGEGROUPS (years)	SUPPORTED		NOT SUPPORTED		TOTAL	
	n	%	n	%	n	%
0-14	41	13	4	6	45	12
15-59	238	75	58	81	296	76
60+	37	12	10	14	47	12
Total	316	100	72	101	388 *	100

* excludes patients transferred out, and those who died before supporters could be organised

A10 SEX AND TREATMENT SUPPORTERS

SEX	SUPPORTED		NOT SUPPORTED		TOTAL	
	n	%	n	%	n	%
Female	95	30	20	28	115	30
Male	221	70	52	72	273	70
Total	316	100	72	100	388 *	100

* excludes patients transferred out, and those who died before supporters could be organised

A11 NEW AND RETREATMENT PATIENTS AND SUPPORTERS

	NEW PATIENTS	RETREATMENT PATIENTS	TOTAL
NO SUPPORTER	62 (81%) 23%	15 (19%) 26%	77 23%
SUPPORTED	213 (84%) 77%	42 (16%) 74%	255 77%
	275 (83%)	57 (17%)	332 * 100%

* pulmonary disease patients

A12 TREATMENT GROUPS AND SUPPORTER CATEGORIES

Supported	Treatment Clinic		Control Clinic		Hospital		Outside District		Total	%
	n	%	n	%	n	%	n	%		
SHOP	43	35	28	43	41	45	11	30	123	39
FAMILY	36	29	20	31	33	36	16	43	105	33
CLINIC	24	19	6	9	1	1	6	16	37	12
NEIGHB	12	10	4	6	6	7	4	11	26	8
TEACHER	4	3	2	3	2	2			8	3
WORK	2	2	3	5	1	1			6	2
CHURCH	1	1	1	2	4	4			6	2
CHW	1	1	-	-	-	-			1	
INDUNA	1	1	1	2	1	1			3	1
HOSP	-	-	-	-	2	2			2	1
TOTAL	124	100	65	100	91	99	37	100	317	101

A13 TREATMENT OUTCOMES OF NEW SMEAR POSITIVE PATIENTS FOR 1992-94, CATEGORISED ACCORDING TO INTERVENTION STUDY GROUPS

PLACE FOR TREATMENT	COMPLETED	INTERRUPTED	DIED	TOTAL
INTERVENTION CLINICS IN DISTRICT	45 (74%)	11 (18%)	5 (8%)	61 (100%) (38% of in-district pats.)
CONTROL CLINICS IN DISTRICT	34 (67%)	12 (24%)	5 (10%)	51 (100%) (32% of in-district pats.)
TINTSWALO HOSPITAL DISTRICT PATIENTS	33 (69%)	14 (29%)	1 (2%)	48 (100%) (30% of in-district pats.)
TOTAL IN DISTRICT	<u>112 (70%)</u>	<u>37 (23%)</u>	<u>11 (7%)</u>	<u>160 (100%)</u> (78% of all pats.)
TINTSWALO HOSPITAL OUTSIDE DISTRICT 10-25 kms	11 (85%)	2 (15%)	0 (0%)	13 (100%)
TINTSWALO HOSPITAL OUTSIDE DISTRICT 26-49 kms	7 (47%)	4 (27%)	4 (27%)	15 (100%)
TINTSWALO HOSPITAL OUTSIDE DISTRICT 50+ kms	8 (44%)	9 (50%)	1 (6%)	18 (100%)
TOTAL OUTSIDE DISTRICT	<u>26 (57%)</u>	<u>15 (33%)</u>	<u>5 (11%)</u>	<u>46 (100%)</u> (22% of all pats.)
GRAND TOTAL	138 (67%)	52 (25%)	16 (8%)	206 * (100%)

* excludes 3 transfers and 1 patient with missing data

A14 TREATMENT OUTCOMES AND SEX - ALL PATIENTS (1994-96)

SEX	CURED AND COMPLETED	INTERRUPTED	FAILED	DIED	TOTAL
FEMALE	94 (79%)	15 (13%)	2 (2%)	8 (7%)	119
MALE	208 (73%)	47 (17%)	14 (5%)	15 (5%)	284
TOTAL	302 (75%)	62 (15%)	16 (4%)	23 (6%)	403 *

A15 TREATMENT OUTCOMES AND SEX - NEW SMEAR POSITIVE PATIENTS (1994-96)

SEX	CURED AND COMPLETED	INTERRUPTED	FAILED	DIED	TOTAL
FEMALE	39 (85%)	4 (9%)	2 (4%)	1 (2%)	46 (26%)
MALE	96 (72%)	22 (17%)	7 (5%)	8 (6%)	133 (74%)
TOTAL	135 (75%)	26 (15%)	9 (5%)	9 (5%)	179 * (100%)

* excludes 4 new smear positive patients transferred out

**A16 OUTCOMES AND AGE GROUPS - NEW SMEAR POSITIVE PATIENTS
(1994-96)**

AGE GROUP in years	CURED AND COMPLETED	INTERRUPTED	DIED	FAILED	TOTAL
0-14	3 (75%)	1 (25%)	0	0	4 (2%)
15-29	41 (91%)	4 (9%)	0	1 (2%)	45 (25%)
30-59	79 (70%)	20 (18%)	8 (7%)	6 (5%)	113 (63%)
60+	12 (71%)	2 (12%)	1 (18%)	2 (12%)	17 (10%)
TOTAL	135 (75%)	26 (15%)	9 (10%)	9 (5%)	179 (100%)

A17 TREATMENT OUTCOMES AND HOSPITAL ADMISSIONS (1994-96)

HOSPITAL ADMISSION	COMPLETED & CURED	INTERRUPTED	DIED	FAILED	TOTAL
NO	35 (73%)	10 (21%)	1 (2%)	2 (4%)	48 (13%)
YES	267 (79%)	49 (15%)	7 (2%)	13 (4%)	336 (88%)
TOTAL	302 (79%)	59 (15%)	8 (2%)	15 (4%)	384 * (100%)

* excludes 11 transferred and 19 for whom admission information not available (of these, 15 died, 1 failed, 3 interrupted)

A18 TREATMENT OUTCOMES AND DURATION OF STAY IN HOSPITAL,
ONE WEEK AND MORE (1994-96)

DAYS IN HOSPITAL	COMPLETED & CURED	INTERRUPTED	DIED	FAILED	TOTAL
1-6	17 (71%)	5 (21%)	1 (4%)	1 (4%)	24 (7%)
7+	250 (80%)	44 (14%)	6 (2%)	12 (4%)	312 (93%)
TOTAL	267 (79%)	49 (15%)	7 (2%)	13 (4%)	336 (100%)

A19 TREATMENT OUTCOMES AND DURATION OF STAY IN HOSPITAL,
ONE MONTH AND MORE (1994-96)

DAYS IN HOSPITAL	COMPLETED & CURED	INTERRUPTED	DIED	FAILED	TOTAL
1-27	157 (77%)	34 (17%)	6 (3%)	7 (3%)	204 (63%)
28+	110 (83%)	15 (11%)	2 (2%)	5 (4%)	132 (37%)
TOTAL	267 (79%)	49 (15%)	8 (2%)	12 (4%)	336 (100%)

A20 TREATMENT OUTCOMES AND TREATMENT SUPPORT,
NEW SMEAR POSITIVE CASES (1994-96)

SUPPORT	COMPLETED	INTERRUPTED	DIED	FAILED	TOTAL
YES	120 (84%)	15 (10%)	1 (1%)	7 (5%)	143 (84%)
NO	15 (54%)	11 (39%)	0	2 (7%)	28 (16%)
TOTAL	135 (79%)	26 (15%)	1 (1%)	9 (5%)	171 * (100%)

* excludes 5 transfers and 8 deaths

TABLE 21: COMPARISON OF PATIENTS WHO HAD 3, 2, 1
AND NO INTERVIEWS

VARIABLE		All 3 inter-views n=185		Two inter-views n=103		One inter-view n=17		No inter-view n=22	
AGE (years)	< 15	n	%	n	%	n	%	n	%
		19	10	15	15	1	6	3	14
	15-59	146	79	76	74	12	71	18	82
	60+	20	11	12	12	4	24	1	5
SEX	Male	129	70	77	75	12	71	17	77
	Female	56	30	26	25	5	29	5	23
DISTANCE (kms)	0-9	92	50	52	51	10	59	12	55
	10-25	70	38	37	36	5	29	6	27
	26-49	23	12	14	14	2	12	4	18
TREAT- MENT OUTCOMES	Cure/ Compl.	163	88	84	82	3	18	3	14
	Interr	9	5	10	10	6	35	10	46
	Transf	2	1	2	2	3	18	1	5
	Died	0	0	5	5	5	29	8	36
	Failed	11	6	2	2	0	0	0	0

A22 SEX AND EDUCATIONAL STATUS

SEX	HIGHEST EDUCATION ACHIEVED			
	Informal	Primary	Secondary	Total
FEMALE	37 (43%) (28%)	24 (28%) (29%)	25 (29%) (29%)	86 (100%) (28%)
MALE	96 (44%) (72%)	59 (27%) (71%)	62 (29%) (71%)	217 (100%) (72%)
TOTAL	133 (44%)	83 (100%) (27%)	87 (29%)	303 (100%) (100%)

A23 PERCENTAGE AGREEMENT FOR SAMPLE OF FIRST AND REPEAT INTERVIEWS

VARIABLE	% AGREEMENT
INTERVIEW 1:	
Address	88
Treatment place	100
Facility first attended	75
Attended: clinic	75
trad healer	100
faith healer	100
hospital	100
Treatment: frequency	75
duration	100
Problems anticipated	88
Belief in cause (open question)	38*
Belief: spread	88
inadequate treatment	75
mines	88
poor nutrition	63*
alcohol	100
smoking	100
food quality	75
Previous TB	88
Family TB now/previous	100/88
Type of work	57*
Education	67*
Post-school education	83
Household size	50*
No. under 15 yrs	33*
No. under 5 yrs	17*
No earning	83
Old age pensioners	67*
Disability pensioners	100
No. sick	83
INTERVIEW 2	
Supporter presence / who	100/73
Transport to hospital clinic	55*
Satisfaction with hospital/clinic	100

VARIABLE	% AGREEMENT
INTERVIEW 2 contd.	
Time to reach supporter	27
Post-hospital consult:	
trad healer	73
faith healer	100
another hospital	100
another clinic	91
private dr.	100
INTERVIEW 3	
Attending supporter	67*
Problems attending supporter	100
Cost	100
CLINIC PATIENTS	
Place attended	100
Time to reach	0*
Transport	50*
Cost	0*
Waiting time	0*
Staff care	100
Preference	100
HOSPITAL PATIENTS	
Transport	90
Cost	67*
Waiting time	0*
Staff care	100
Problems	44*
Preference	67*
GENERAL	92
Difficult treatment	93
Side effects	54*
Staff kind	69*
Trad healer role	93

* < 70% agreement

TABLE 24 SUMMARY OF BIVARIATE ANALYSIS OF TREATMENT OUTCOMES AND VARIABLES n=296

VARIABLE	OUTCOME		p	OUTCOME	
	Cured / Compl.	Interrup / Failed/Died		Interrup/ Failed	p
Education: n=296					
Informal	111	21	0.89	14	0.36
Formal	137	27		24	
Informal & primary	177	35	0.82	25	0.48
Secondary	71	13		13	
Below std 9 & 10	32	4	0.37	4	0.68
Std 9 or 10	216	44		34	
Age (years): n=319					
0-5	16	4	0.93	4	0.39
6+	237	62		44	
0-15	33	5	0.22	5	0.61
16+	220	61		43	
0-29	89	11	0.004	11	0.09
30+	164	55		37	
0-59	226	56	0.31	41	0.43
60+	27	10		7	
15-29	56	6	0.009	6	0.6
30+	164	55		37	
15-59	193	51	0.40	36	0.47
60+	27	10		7	
Sex: n=319					
Female	78	13	0.074	6	0.009
Male	175	53		42	
Work: n=319					
Unemployed	90	24	0.90	14	0.39
Employed	163	42		34	
Manual	120	29	0.55	22	0.29
Other employed	43	13		12	
Skilled/semi-skilled	43	13	0.55	12	0.29
Manual	120	29		22	

VARIABLE	OUTCOME		p	OUTCOME	
	Cured / Compl.	Interrupt / Failed/Died		Interrupt / Failed	p
Time to reach supporter: n=319					
0 (same home)	81	9	0.003	9	0.06
Rest (>0 mins)	172	57		57	
Stated problems with treatment access: n=282					
Yes	28	12	0.0006	10	0.005
No	217	25		22	
How patients reached treatment source:					
Those nearest hospital (hospital treatment): n=101					
Walk	27	7	0.35		
Other	58	9			
Those on hospital treatment (Control treatment) n=64					
Walk	1	1	0.20		
Other	56	6			
Those on clinic treatment: n=117					
Walk	57	7			
Other	46	7			
CLINIC PATIENTS: n=74					
Time to reach clinic:					
< 30 mins	50	6			
30+ mins	15	3			
Cost to reach:					
0 - R3	61	8			
> R3	4	1			
0	37	2			
>0 (R1+)	28	7			
How reached:					
Walk	37	2			
Other	28	7			
Waiting time:					
<30 mins	51	14			
30+ mins	8	1			
Problems experienced					
Yes	6	59			
No	2	7			

VARIABLE	OUTCOME		p	OUTCOME	
	Cured / Compl.	Interrup / Failed/Died		Interrup/ Failed	p
HOSPITAL PATIENTS: n=112					
Time to reach:					
<30 mins	58	5	0.32		
30+ mins	43	6			
Cost to reach:					
0 - R3	72	7	0.41		
>R3	29	4			
0	29	4	0.41		
>0 (R1+)	72	7			
How reached: n=113					
Walk	30	4	0.54		
Other	72	7			
Waiting time:					
<30 mins	18	1	0.40		
30+ mins	83	10			
Problems experienced:					
Yes	28	2	0.39		
No	74	9			

UNIVERSITY OF THE WITWATERSRAND, JOHANNESBURGDivision of the Deputy Registrar (Research)COMMITTEE FOR RESEARCH ON HUMAN SUBJECTS (MEDICAL)
Ref: R14/49 (Registry)CLEARANCE CERTIFICATEPROTOCOL NUMBER M940103PROJECTRecord review of TB patients
admitted to Tintswalo HospitalINVESTIGATORS

Dr M E Edginton

DEPARTMENT

Community Health

DATE CONSIDERED

940128

DECISION OF THE COMMITTEE

Approved unconditionally

DATE 940204

CHAIRMAN *P E Cleaton-Jones*
(Professor P E Cleaton-Jones)* Guidelines for written "informed consent" attached where
applicable.

=====

To be completed in duplicate and ONE COPY returned to the Secretary at Room
10001, 10th Floor, Senate House, University.I/We fully understand the conditions under which I am/we are authorized to
carry out the abovementioned research and I/we guarantee to ensure
compliance with these conditions. Should any departure be contemplated
from the research procedure as approved I/we undertake to resubmit the
protocol to the Committee.DATE... *Nov 1993* ...SIGNATURE... *P E Cleaton-Jones* ...

UNIVERSITY OF THE WITWATERSRAND, JOHANNESBURG

B2

Division of the Deputy Registrar (Research)

COMMITTEE FOR RESEARCH ON HUMAN SUBJECTS (MEDICAL)

Ref: R14/49 Edginton

CLEARANCE CERTIFICATE

PROTOCOL NUMBER M 940806

PROJECT

Study to evaluate a strip test to detect the drug INH and its metabolites in urine of patients taking INH orally

INVESTIGATORS

Dr M E Edginton

DEPARTMENT

Community Health, Medical School

DATE CONSIDERED

940826

DECISION OF THE COMMITTEE *

Approved unconditionally

DATE

940830

CHAIRMAN. *Patrick P. Cleaton-Jones* (Professor P E Cleaton-Jones)

* Guidelines for written "informed consent" attached where applicable.

c c Supervisor: Dr M E Edginton
Dept of Community Health, Medical School

Works2\other\hclear\ 940806

DECLARATION OF INVESTIGATOR(S)

To be completed in duplicate and ONE COPY returned to the Secretary at Room 10001, 10th Floor, Senate House, University.

I/we fully understand the conditions under which I am/we are authorized to carry out the abovementioned research and I/we guarantee to ensure compliance with these conditions. Should any departure to be contemplated from the research procedure as approved I/we undertake to resubmit the protocol to the Committee.

DATE..... *1/9/94* SIGNATURE *[Signature]*

UNIVERSITY OF THE WITWATERSRAND, JOHANNESBURGDivision of the Deputy Registrar (Research)COMMITTEE FOR RESEARCH ON HUMAN SUBJECTS (MEDICAL)

Ref: R14/49 (Registry)

CLEARANCE CERTIFICATEPROTOCOL NUMBER M940203

PROJECT Tuberculosis in the Eastern
Transvaal: a study in management

INVESTIGATORS Dr M E Edginton

DEPARTMENT Community Health

DATE CONSIDERED 940225

DECISION OF THE COMMITTEE

Approved with the attached condition

DATE 940302

 CHAIRMAN
 (Professor P E Cleaton-Jones)

* Guidelines for written "informed consent" attached where applicable.

 =====
DECLARATION OF INVESTIGATOR(S)

To be completed in duplicate and ONE COPY returned to the Secretary at Room 10001, 10th Floor, Senate House, University.

I/we fully understand the conditions under which I am/we are authorized to carry out the abovementioned research and I/we guarantee to ensure compliance with these conditions. Should any departure to be contemplated from the research procedure as approved I/we undertake to resubmit the protocol to the Committee.

DATE 8/3/74 SIGNATURE

QUESTIONNAIRE FOR DATA ENTRY INTO EPIINFO

name _____ tb reg no. # / # / #
 age in years # # sex <A> address _____ nearest clinic _____
 date treatment started <dd/mm/yy> treatment regime #
 source of patient <A> if H state which _____ patient category <A>
 diagnostic criteria <A> internat code # # #
 sputum exam. pretreatmnt. 8 weeks 6 months
 result date result date result date
 smear <A> <dd/mm/yy> <A> <dd/mm/yy> <A> <dd/mm/yy>
 culture <A> <dd/mm/yy> <A> <dd/mm/yy> <A> <dd/mm/yy>
 resistance <A> <A> <A>
 date of discharge from hospital <dd/mm/yy>
 post-discharge treatment place _____
 if supervised, who supervises _____, who organised _____
 how did patient reach clinic _____
FOR CLINIC TREATMENT PATIENTS date first seen at clinic <dd/mm/yy>
 subsequent clinic visits no. days supervsd trtmnt during mnth
 scheduled actual
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 any hospital visits <dd/mm/yy> why _____
 <dd/mm/yy> _____
 <dd/mm/yy> _____
 <dd/mm/yy> _____
 final hospital visit for discharge <dd/mm/yy>
FOR HOSPITAL TREATMENT PATIENTS dates of attendance
 scheduled actual days of supervision
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 <dd/mm/yy> <dd/mm/yy> # #
 readmission for tb admission date <dd/mm/yy> discharge date <dd/mm/yy>
 outcome <A> TRF # date <dd/mm/yy> HIV <A>
 interview 1 <dd/mm/yy>
 2 scheduled <dd/mm/yy>
 actual <dd/mm/yy>
 3 scheduled <dd/mm/yy>
 actual <dd/mm/yy>

remarks _____

REPORT OF PATIENT ATTENDANCES AND SPUTA RESULTS

DATE: _____

PATIENT'S NAME: _____

TB NUMBER: 9____/____

CLINIC (name) OR HOSPITAL PATIENT: _____

DATE TREATMENT STARTED: _____

DATE DISCHARGED FROM HOSPITAL: _____

PATIENT TAKEN TO CLINIC BY: _____

DATE FIRST SEEN AT CLINIC: _____

SUPPORTER: _____

SUPPORTER ORGANISED BY: _____

ATTENDANCES:

SCHEDULED	ACTUAL	DOCTOR (D) PILLS (P)	CLINIC (C) HOSPITAL (H)	DAYS OF SUPPORT
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____

SPUTUM TESTING:

INITIAL DATE	RESULT	2 MONTHS DATE	RESULTS
_____	_____	_____	_____
_____	_____	_____	_____

5-6 MONTHS DATE	RESULT
_____	_____
_____	_____

FINAL DISCHARGE: _____

[illegible]

TINTSWALO TUBERCULOSIS PROGRAMME
DATA SHEET FOR INTERRUPTER CASE CONTROL STUDY

INTERUPPTER OR CONTROL

BASIC INFORMATION FROM RECORDS: (MEE to enter from records)

TB NUMBER...../9.. NAME.....

AGE..... GENDER M/F HIV P/N SOURCE..... CATEGORY.....

TB CODE..... SMEAR P/N CULTURE P/N

ADDRESS (name of village).....

FIRST ADMISSION/READMISSION F/R

If READMISSION, for first admission:

TB NO...../9.. DATE..... OUTCOME.....

FAMILY SIZE POSITION OF THIS PATIENT IN FAMILY

Days in hospital TB ward

TB TREATMENT PLACE (clinic name/Tintswalo)

Supporter Category

Supporter attendance per month

Outcome:

Cure S

Compleat.C1(monthly) C2(monthlyx1st 2m) C3(2-3monthly) C4

Interrupter I1 (incompl.1st 2m.) I2(compleat.1st 2 m.only)

I3(never seen postdisc)

RHT from hospital..... RHT

Failed to return after passout FR

Other.....

ACCESS - INFORMATION ON TREATMENT PLACE: (MEE to ask CS/SN/HEDs)

What is the best way to get from patient's home to the treatment place

If best way is by taxi/bus,

How much does it cost to get from patient's home to that place

R.....

How long does it usually take

Are taxis/buses regular Y/N

If best way is walking,

How long does it take to walk (one way home to treatment place)

.....

INFORMATION FROM REPORT OF CONNIE/SELLITA/PHILLIP/SOLLY:

ANY PROBLEMS IDENTIFIED BY STAFF AS:

DISTANCE TO TREATMENT PLACE	Y/N
NO MONEY	Y/N
ATTITUDE/BEHAVIOUR OF STAFF AT TR. PLACE (fr.pt)	Y/N
ALCOHOL	Y/N
PATIENT AWAY AT WORK	Y/N
PATIENT CANNOT BE TRACED	Y/N
Other (say what)	

.....

.....

INFORMATION FROM PATIENT INTERVIEW

First place attended for treatment before T C/TH/FH/AnotherH/GP/Other

Belief re TB (open)

Family TB NOW/PREVIOUS

Work type

Education (highest level)

Household size

Under 15.....Under 5 OAPen Disab.Pen

Household income R.....

Knowledge re treatment G/B

Treatment Place same as records Y/N

Supporter same as records Y/N

Clinic access by Walk/Taxi/Bus/Other.....

Clinic transport cost R

Clinic wait timeminutes

Clinic staff problem Y/N

Other clinic problems Y/N what.....

Hospital access by Walk/Taxi/Bus/Other.....

cost R.....

wait timeminutes

staff problem Y/N

other problems Y/N what.....

General statements (ring only if problem)

Pills difficult to take

Pills make sick

Nurses not good

Traditional healer essential

- A2 Where did you go first when you got sick with this TB?

☐ 39

Tintswalo hospital	Other hospital	Clinic	Traditional healer
Faith healer	Private doctor	Other - say where	

TUBERCULOSIS KNOWLEDGE

- K1 Please tell me about the treatment you must take for your TB:

For how long must you take the treatment?

How often should you take the treatment?

☐ 40

- K2 Where will you go regularly to fetch your TB treatment after you are discharged from hospital?

☐ 41

- K3 Do you think you will have any problems in getting to the place where you will get your treatment?

Yes	No
-----	----

☐ 42

- K4 If yes, please explain _____

☐ 43

- K5 Are there questions you would like to ask about your TB treatment

☐ 44

(Interviewer: if you cannot answer the questions, ask someone qualified to do so)

TUBERCULOSIS CAUSE

- C1 Why do you believe you got TB?

☐ ☐ 45

TINTSWALO DISTRICT TUBERCULOSIS PROGRAMME

Interview for all patients who live in the Tintswalo district before their discharge from hospital

Card No.

1

 1

Name of Patient: _____

TB Registration No. _____

9				
---	--	--	--	--

 2

Date of Admission

				9	
--	--	--	--	---	--

 7

Date of interview

				9	
--	--	--	--	---	--

 13

Date of discharge

				9	
--	--	--	--	---	--

 19

Place of Interview: _____

Address: Name of village/town

--	--

 25

Exact address with landmarks:

--	--

 26

Place for follow up treatment _____

--	--

 26

Interviewer: Check records and/or ask ward sister where this patient is to receive follow-up treatment and write it here.

TUBERCULOSIS ACTION

A1 Please tell me about places you went to when you got sick with TB, before you came to hospital. Have you been to a:

clinic

Yes	No
-----	----

--

 30

say where _____

--

 31

traditional healer

Yes	No
-----	----

--

 32

faith healer

Yes	No
-----	----

--

 33

hospital other than this one

Yes	No
-----	----

--

 34

say which _____

--

 35

private doctor

Yes	No
-----	----

--

 36

other place/person

Yes	No
-----	----

--

 37

say where/who _____

--

 38

C2 Do you think people get tuberculosis from:-

1	Bewitchment Neglect of ancestors	Yes	No	Don't know
2	Disobeying rules when family member dies	Yes	No	Don't know
3	Sexually transmitted disease	Yes	No	Don't know
4	Spread from people with TB Germs	Yes	No	Don't know
5	TB inadequately treated in the past	Yes	No	Don't know
6	Work in mines or dusty conditions	Yes	No	Don't know
7	Poor nutrition or diet	Yes	No	Don't know
8	Alcohol	Yes	No	Don't know
9	Smoking cigarettes	Yes	No	Don't know
10	Food or water contamination	Yes	No	Don't know
11	Other	Yes	No	Don't know

47

48

49

50

51

52

53

54

55

56

57

PREVIOUS TUBERCULOSIS

P1 Have you ever had TB before this time?

☐ 58

Yes	No
-----	----

P2 If yes,
What year was this 19_____ 59

P3 Where did you have treatment

 61

P4 For how long were you treated (in months)_____

 62

FAMILY TB

F1 Is anyone in your family on treatment for TB
now

Yes	No
-----	----

☐ 64F2 Has anyone in your family been on treatment in
the past

Yes	No
-----	----

☐ 65

WORK OUTSIDE THE HOME

W1 When did you last work for money? (write month and year) _____

		9	
--	--	---	--

 66

W2 For how many years have you worked for money in your life?

(Interviewer: write down all the different jobs and add up the total time worked)

Total Time _____

--	--

 70

W3 What is the main kind of work you have done? (Interviewer ask for the job the person has had for the longest) _____

--	--

 72

Card No.

2

 1

TB Registration No.

9				
---	--	--	--	--

 2

EDUCATION

E1 What is the highest school standard that you passed eg Nil, SubA, SubB, Std1 - 10 _____

--	--

 7

E2 Did you have any education or training after you left school

Yes	No
-----	----

--

 9

E3 If yes, what was this _____

--	--

 10

HOUSEHOLD INFORMATION

Tell me about the people in your household. (Tick appropriate squares)

	First name	Relation- ship to you	Year of birth	Age	Earning money	Old Age pension	Disa- bility Pension	Sick now for 2 wks or more
1		yourself						Yes
2								
3								
4								
5								
6								
7								
8								
9								
10								
11								
12								
13								
14								
15								
16								
17								
18								

Subject's age	☐	12
Total No.	☐	14
Under 15 years	☐	16
Under 5 years	☐	17
Earning	☐	18
Old age pension	☐	19
Disability pension	☐	20
Sick (other than patient)	☐	21

How much money comes into your household each month (in Rands) _____

☐	☐	☐	☐	☐	22
--------------------------	--------------------------	--------------------------	--------------------------	--------------------------	----

Thank you for answering my questions.

TINTSWALO DISTRICT TUBERCULOSIS PROGRAMME

PATIENT INTERVIEW AT 8 WEEKS AFTER TREATMENT ONSET

Card No

3

 1

Name of Patient: _____

TB Registration No. _____

9				
---	--	--	--	--

 2Date of interview: _____

				9	
--	--	--	--	---	--

 7Place of interview: _____

--

 13If patient is under 15 years of age, name of mother
or person interviewed on behalf of child: _____

--

 14Are you currently taking treatment for TB? _____

--

 15

Yes	No
-----	----

If no:
Please explain why not _____

--

 16If yes:
How often do you take your pills? _____

--

 17Does someone supervise your treatment i.e. give you
your pills? _____

--

 18

Yes	No
-----	----

If yes:
Who is your supervisor? _____

--

 19

Name: _____

Position or relationship: _____

How long does it take you to go to your supervisor
(in minutes) _____Please explain _____

--	--

 20When you go for a check up and to get more pills do
you go to the _____

Hospital	Clinic
----------	--------

--

 22How do you get to the clinic or hospital? _____

--

 23

Do you have any problems getting to the clinic/hospital to fetch your pills?

Yes	No
-----	----

☐ 24

If yes, please tell me about these problems.

☐ 25

What do you feel the pills do for you?

☐ 26

Why do you feel this?

☐ 27

For how many months must you take the pills? _____
(from start to finish)

☐ ☐ 28

Are you satisfied with the service you are getting at the clinic/hospital?

☐ 30

(Interviewer delete whichever is not applicable)

Yes	No
-----	----

If no, please tell me about the problems.

☐ 31

Have you been to see any of the following since you were discharged from hospital?

Traditional healer

Yes	No
-----	----

☐ 32

Faith healer

Yes	No
-----	----

☐ 33

Another Hospital

Yes	No
-----	----

☐ 34

Another clinic

Yes	No
-----	----

☐ 35

A private doctor

Yes	No
-----	----

☐ 36

Any other healer or service
say which: _____

Yes	No
-----	----

☐ 37

Do you have any questions about your sickness or things you would like to discuss?

☐ 38

(Interviewer if you cannot answer the questions, ask someone who can to see this patient.)

Thank you for answering these questions.

INTERVIEW SCHEDULE FOR TB PATIENTS WHO HAVE COMPLETED THEIR TREATMENT

Interview to take place at Tintswalo hospital at the time of final discharge

Interviewer: Introduce yourself, and explain that the patient has the right to refuse to answer questions, and that answers will be kept confidential.

Name of patient: _____

Card No. 1

TB registration number

2

Date of interview

7

Place of interview _____

13

Patient's address: (name of village or place)

14

Which is your nearest clinic (ask patient this)?

16

SUPPORTER FOR TREATMENT

Have you been going daily to a supporter to get your treatment?

Yes No

16

If yes, who was that supporter?

[shop/ induna/ family member/ clinic nurse]

Say exactly who it was? _____

19

Did you have any problems getting to your supporter every day?

Yes No

21

If yes, please tell me about your problems.

22

Did it cost you money to go to your supporter every day?

Yes No

If yes, how much did it cost? (Rands and cents) _____

23

WHERE HAVE YOU BEEN ATTENDING EVERY MONTH TO SEE A DOCTOR OR NURSE TO FETCH MORE PILLS?

26

IF YOU HAD TREATMENT AT A CLINIC:

Which clinic did you go to? _____

 27How long did it take you to get there? _____
(In minutes) 29

How did you get there? _____

 32

How much money did you have to pay to get there?

(in Rands and Cents) _____

 33How long did you have to wait at the clinic before
you were seen by the nurse (in minutes)? _____ 37Did the clinic staff look after you well?

Yes	No
-----	----

If no, please explain _____

 39Please explain any problems you had in getting your
treatment at the clinic?

 40Would you have preferred to go to hospital for your
treatment rather than to the clinic

Yes	No
-----	----

 41

Whether person said yes or no, ask:-

Please explain why you have said this

 42

IF YOU HAD TREATMENT AT TINTSWALO HOSPITAL:

How long did it take you to get there? (in minutes)_____

 43

How did you get there? _____

 46How much money did you have to pay to get there?
(in Rands and Cents) _____ 47How long did you spend at the hospital
(from the time you got to the ward until
you left to go home)? (in minutes) _____ 51

Did the hospital staff look after you well?

Yes	No
-----	----

 54

If no, please explain _____

 55Please explain any problems you had in getting your
treatment at the hospital?

If yes, explain the problems _____

 56Would you have preferred to go to your nearest
clinic for your treatment rather than to the hospital?

Yes	No
-----	----

 57

Please explain why you have said this (whether yes or no)

 58

GENERAL COMMENTS:

If one of your friends or a neighbour got TB, and he asked you about TB, what would you tell that person about the treatment for TB?

Would you tell him that TB pills are difficult to take?

☐ 59

Yes	No
-----	----

☐ 60

Would you tell him that the pills sometimes make you feel sick?

Yes	No
-----	----

☐ 61

Would you say the nurses at the clinic or hospital were good/kind/helpful?

Yes	No
-----	----

☐ 62

Would you tell him he should go to a traditional healer?

Yes	No
-----	----

☐ 63

Would you tell them anything else about TB or TB treatment?

☐ 64

Do you feel that your TB is now cured?

Yes	No
-----	----

☐ 65

If no, please explain why not _____

☐ 66

Have you got a job earning money since you were treated for TB?

Yes	No
-----	----

☐ 67

Interviewer:

Note yes or no here if this person came from Mozambique, and if possible about how long ago (in years). Do not ask the person directly.

Yes	No
-----	----

☐ 68

How many years ago did he/she come _____?

☐ 69

Thank the patient for answering your questions.

REPORT ON PATIENT NOT FOUND WHEN LOOKED FOR AT HOME

FIRST VISIT/SECOND VISIT/THIRD VISIT (ring the one that applies)
Do not make more than 3 visits to try to find a patient

Name of patient..... TB no/9..

Did you find patient's home? Yes/No

Who was at the home?

If someone at patient's home, did they
know the patient ? Yes/No

If they know patient, where did they
say he/she was ?
.....

Does he ever come home? Yes/No

If yes, when will he next come home?

Is he/she well? Yes/No

Is he/she taking TB treatment? Yes/No

If yes, where does he get the pills?

If no-one was found at patient's home

Did neighbours or people nearby know
the patient? Yes/No

If yes, did they know where he/she is? Yes/No

If yes, where did they say patient is

If patient is said to be somewhere else in the Tintswalo or the
old Mhala district (even if outside our study area), look for the
patient there and interview. If he is said to be beyond the
Tintswalo or old Mhala area, don't look further.

Author: Edginton M.E

Name of thesis: Tuberculosis in the Tintswalo district of the Northern Province

PUBLISHER:

University of the Witwatersrand, Johannesburg

©2015

LEGALNOTICES:

Copyright Notice: All materials on the University of the Witwatersrand, Johannesburg Library website are protected by South African copyright law and may not be distributed, transmitted, displayed or otherwise published in any format, without the prior written permission of the copyright owner.

Disclaimer and Terms of Use: Provided that you maintain all copyright and other notices contained therein, you may download material (one machine readable copy and one print copy per page) for your personal and/or educational non-commercial use only.

The University of the Witwatersrand, Johannesburg, is not responsible for any errors or omissions and excludes any and all liability for any errors in or omissions from the information on the Library website.