BMJ Open Plasma concentrations of second-line antituberculosis drugs in relation to minimum inhibitory concentrations in multidrug-resistant tuberculosis patients in China: a study protocol of a prospective observational cohort study

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ABSTRACT

Introduction Individualised treatment through therapeutic drug monitoring (TDM) may improve tuberculosis (TB) treatment outcomes but is not routinely implemented. Prospective clinical studies of drug exposure and minimum inhibitory concentrations (MICs) in multidrug-resistant TB (MDR-TB) are scarce. This translational study aims to characterise the area under the concentration–time curve of individual MDR-TB drugs, divided by the MIC for *Mycobacterium tuberculosis* isolates, to explore associations with markers of treatment progress and to develop useful strategies for clinical implementation of TDM in MDR-TB.

Methods and analysis Adult patients with pulmonary MDR-TB treated in Xiamen, China, are included. Plasma samples for measure of drug exposure are obtained at 0, 1, 2, 4, 6, 8 and 10 hours after drug intake at week 2 and at 0, 4 and 6 hours during weeks 4 and 8. Sputum samples for evaluating time to culture positivity and MIC determination are collected at days 0, 2 and 7 and at weeks 2, 4, 8 and 12 after treatment initiation. Disease severity are assessed with a clinical scoring tool (TBscore II) and quality of life evaluated using EQ-5D-5L. Drug concentrations of pyrazinamide, ethambutol, levofloxacin, moxifloxacin, cycloserine, prothionamide and para-aminosalicylate are measured by liquid chromatography tandem-mass spectrometry and the levels of amikacin measured by immunoassay. Dried blood spot on filter paper, to facilitate blood sampling for analysis of drug concentrations, is also evaluated. The MICs of the drugs listed above are determined using custom-made broth microdilution plates and MYCOTB plates with Middlebrook 7H9 media. MIC determination of pyrazinamide is performed in BACTEC MGIT 960.

Ethics and dissemination This study has been approved by the ethical review boards of Karolinska Institutet, Sweden and Fudan University, China. Informed written consent is given by participants. The study results will be submitted to a peer-reviewed journal.

Strengths and limitations of this study

- To our knowledge, this is a novel study approach which fully investigates the distribution of drug exposure in relation to minimum inhibitory concentration (MIC) for *Mycobacterium tuberculosis* (Mtb) isolates from patients with multidrug-resistant tuberculosis (TB) along with biomarkers (eg, time to positivity), culture conversion and the clinical scoring tool TBscore II to assess treatment outcome.
- We used a translational approach with experts from research centres across the world to design a study protocol including both MIC determinations, drug exposure estimation using novel technology, as well as microbiological and clinical surrogate markers for improvement, to enable strategies for therapeutic drug monitoring use in TB treatment.
- The patients' drug exposure will be compared with individual Mtb MICs, exploring pharmacokinetics–pharmacodynamics indices in multidrug-resistant TB treatment.
- Dried blood spot as a method to simplify blood sampling by finger prick instead of venous sampling will be investigated.
- A limitation of the study is the low target number of patients for inclusion, due to a laborious and costly study protocol, which might partly be compensated for by using pharmacometric modelling.

Trial registration number NCT02816931; Pre-results.

INTRODUCTION

Despite programmatic management of tuberculosis (TB), the incidence of multidrug-resistant TB (MDR-TB), defined as

Mycobacterium tuberculosis (Mtb) resistant to rifampicin and isoniazid, is steadily increasing.¹ Inconsistent treatment, due to poor treatment adherence, lack of drugs, as well as subtherapeutic dosing are contributing factors. For many TB drugs, the administered dose is not predictive of the drug exposure and clinical effect in the patient.² A hollow-fibre study indicated that pharmacokinetic variability may be an underestimated cause of drug resistance development³ and low drug concentrations in the treatment of drug-susceptible TB have been associated with poor outcome in some prospective studies.^{4,5}

Therapeutic drug monitoring (TDM) is a strategy to personalise treatment by measuring systemic drug levels in blood/plasma as a guide for individual dose adjustments.⁶ Specifically for infectious diseases, the drug efficacy not only depends on the drug exposure but also on the susceptibility level of the bacteria, the minimum inhibitory concentration (MIC).⁷ The MIC is defined as the lowest concentration of a drug that inhibits visible growth of bacilli and should be exceeded to cure the infection.⁸ TDM has been recommended during MDR-TB treatment by several organisations, for example, the Infectious Disease Society of America.⁹ The pharmacokinetic studies that have been performed have shown that subtherapeutic drug levels in TB treatment are common, although with conflicting results regarding association between drug exposure and treatment outcome.^{10–12} However, studies on MDR-TB are limited and only a few studies have included drug concentrations as well as the individual MICs of the bacteria.^{13–15}

An optimal estimation of drug exposure (ie, area under the concentration versus time curve (AUC)) traditionally requires multiple venous blood samples, often followed by prompt centrifugation and sample storage at -80°C . A simplified strategy for collection and transportation of blood samples needed for TDM would aid its implementation in clinical practice. Dried blood spot (DBS) allows minimal blood sampling by capillary finger pricking on filter paper, which can be transported without a cold chain, simplifying transportation and storage.¹⁶ DBS is a well-established and validated method, but has only been evaluated for a few second-line TB drugs, for example, moxifloxacin and linezolid.^{16–18} A clinical implementation of DBS could enable TDM for TB treatment in remote areas and reduce costs.¹⁶

There are scarce data regarding drug exposure and treatment outcome in MDR-TB treatment. Assessing end-of-treatment outcome in MDR-TB studies is cumbersome due to long treatment durations. Other strategies include using interim endpoints such as time to positivity (TTP) in liquid culture media, a surrogate of bactericidal activity,¹⁹ and sputum culture conversion after 2 or 3 months of treatment,²⁰ the latter commonly used in drug efficacy studies. A clinical composite scoring system, TBscore II, can be used as a surrogate marker for TB disease severity and to predict failure.²¹ Patients' quality of life can be objectified using the validated EQ-5D-5L tool assessing five different dimensions (mobility, self-care,

typical activity, pain/discomfort and anxiety/depression), an often overlooked tool in clinical treatment studies.

China has the second highest burden of MDR-TB in the world and has existing resources to perform TDM, thus making it an ideal setting for pharmacokinetic/pharmacodynamic (PK/PD) studies. The overall incidence of TB in China was 895 000 TB cases in 2017, of which 8.2% were MDR-TB.²²

We describe a new comprehensive approach to TDM studies, assessing drug exposure, individual MICs as well as clinical outcome markers. The primary aim of the study is to investigate the distribution of AUC/MIC and $C_{\text{max}}/\text{MIC}$ for MDR-TB drugs during MDR-TB treatment in China. Secondary aims are to analyse AUC/MIC in relationship to markers of clinical improvement, such as sputum culture conversion, TTP, TBscore II, body mass index (BMI) and qualitative measures of well-being (European Quality of Life scale, EQ-5D-5L). Signs of acquired resistance are assessed by investigating changes in MICs and genetic mutations, during the first 3 months of treatment. A clinical implementation of DBS as well as a method of simultaneous MIC determination are assessed to simplify the use of TDM in clinical practice.

METHODS AND ANALYSIS

Study design

We are conducting a prospective cohort study of TB drug exposure and MICs in patients with MDR-TB in Xiamen, China. This is a joint project between the School of Public Health Fudan University Shanghai, Department of Medicine Karolinska Institutet, Department of Pharmaceutical Biosciences University of Uppsala and the Public Health Agency of Sweden, in collaboration with the Centre for Disease Control (CDC) in Xiamen. The study protocol conforms with the Strengthening the Reporting of Observational Studies in Epidemiology Statement for cohort studies.²³

Patient and public involvement

The original study protocol by the coauthors was changed by reducing the number of blood samples after feedback from patients included in a pilot study. The result of the study can be obtained in Mandarin on request at the Xiamen CDC. Patients were not involved in the recruitment and the conduct of the study.

Study setting

The study is carried out in Xiamen, Fujian region in Southeast China, where the incidence of TB in 2016 was 42.4 cases/100 000 inhabitants and of the 1661 confirmed cases that year, there were 28 MDR-TB patients (1.7%).²⁴

The study hospital is the designated TB hospital in Xin Ling, Xiamen, a large teaching hospital with a specialised TB ward with 105 beds as well as a negative pressure ward (12 beds) with specialised TB physicians and nurses. Recruitment of patients is performed by the Xiamen CDC, which also keeps a screening log. Patients are routinely

admitted for 2 months of in-patient treatment. The study is registered at ClinicalTrials.gov (NCT02816931) and opened on 17 April 2016.

Study participants

A total number of 30 fully evaluable patients, according to the criteria below, will be included.

Inclusion criteria

- ▶ Consenting adults (≥ 18 years) with a verified diagnosis of pulmonary MDR-TB, by routine drug susceptibility testing (DST) admitted to the Xin Ling TB Hospital, Xiamen.
- ▶ Eligible for and consent to MDR-TB treatment in Xiamen.

Exclusion criteria

- ▶ Pregnancy
- ▶ HIV infection
- ▶ Patients admitted to the intensive care unit
- ▶ Confirmed extensively drug-resistant TB by DST
- ▶ Ongoing medication for MDR-TB (ie, five active drugs or more for more than 1 day.)

Study outline

The overall study outline is shown in figure 1. After informed consent, a completed inclusion questionnaire with demographic and clinical information, baseline blood and sputum samples are collected from the patient by a designated study nurse. Treatment regimens adhere to WHO guidelines and are adjusted following DST results. The first day of MDR-TB treatment is defined as 'day 0'. Clinical data are collected at inclusion, day 2, weeks 1, 2, 4, 8 and week 12 after treatment initiation. Adverse events, routine blood tests and vital signs are closely monitored to ensure the safety of the study patients. The final treatment outcome is recorded at the end of MDR-TB treatment.

Drug concentrations of second-line TB drugs are measured at steady state at 2, 4 and 8 weeks after treatment initiation. In order to estimate the AUC, multiple blood samples for drug concentration analysis (ie, rich sampling) are collected at week 2 (0, 1, 2, 4, 6, 8 and 10 hours after drug intake). A sparse-sampling strategy is applied at weeks 4 and 8 (0, 4 and 6 hours). Whole blood samples are simultaneously collected and pipetted directly onto DBS cards. Finger prick blood samples are collected on DBS cards at week 2 (0, 4 and 6 hours after drug intake) (figure 2). The drug concentrations in plasma and DBS will be analysed using liquid-chromatography tandem mass spectrometry (LC-MS/MS) and immunoassay.²⁵ In order to assess delayed absorption and possible interactions, information of concomitant drugs is noted in the medical records. Additionally, detailed food intake is noted by the patient in a diary on the days of blood sample collection. Pharmacometric modelling and simulation will be performed in the analysis phase.

Sputum is collected at days 0 and 2 and weeks 1, 2, 4, 8 and 12 in order to evaluate changes in TTP. Whole

genome sequencing (WGS) and MIC determination for TB drugs (pyrazinamide (PZA), ethambutol, levofloxacin, moxifloxacin, ofloxacin, cycloserine, ethionamide, para-aminosalicylic acid (PAS), amikacin, kanamycin, rifampicin and isoniazid) are performed at baseline and for any positive culture after 1 month or more of treatment, to assess development of acquired resistance. Time to sputum culture conversion is defined as the day from starting treatment until the day of the first of two consecutive negative sputum cultures, collected at least 30 days apart.

Disease severity is estimated and monitored using inflammatory markers such as C reactive protein and erythrocyte sedimentation rate, presence of cavity on chest X-ray as well as the total score obtained in TBscore II, based on the following variables; cough, dyspnoea, chest pain, anaemia (pale lower conjunctivas), BMI and mid-upper arm circumference.²¹ Quality of life during the first 3 months of treatment is estimated using the validated EQ-5D-5L-5L (Mandarin version).²⁶ The patients are followed up until treatment completion or loss to follow-up, whichever occurs first, through information accessible from the TB-registry, Xiamen CDC.

LABORATORY METHODS

Drug concentration measurement

A combined assay for drug concentration analysis using LC-MS/MS is under development at the Xiamen CDC to measure the plasma concentrations of PZA, ethambutol, levofloxacin, moxifloxacin, cycloserine, prothionamide and PAS.^{27 28} The second-line injectable drug amikacin will be analysed with a commercial immunoassay kit (amikacin assay kit, Beckman Coulter). The collected venous blood samples will be centrifuged at 3500 rpm for 10 min within 1 hour from sampling. Aliquots of plasma are then frozen at -70°C awaiting analysis.

A puncture from the DBS card will be immersed in extraction solution as previously described¹⁸ and analysed through LC-MS/MS. Plasma concentrations will be compared with blood concentrations collected by DBS.¹⁸

Microbiology: TTP, DST and MIC

All microbiological tests are carried out at a biosafety laboratory level 3 at the Xiamen CDC, apart from routine DST testing which is partly performed in local hospital laboratories and WGS analysis performed at the Public Health Agency of Sweden.

Time to culture positivity

Sputum samples are treated according to Chinese National standards based on a WHO recommended protocol.²⁹ In short, NALC-NaOH is added to the sputum, then shaken using a vortex shaker until fully liquefied, followed by incubation for 15 min in room temperature. Phosphate buffer is added to reach a total volume of 45 mL, after which the solution is centrifuged for 15 min at 3000 g. The supernatant is removed and 1 mL phosphate buffer is

Figure 1 Study overview. Study patients are given a drug diary to record concomitant drugs and food intake during the first 12 weeks. Sputum samples are collected regularly during the study to assess time to positivity in BACTEC MGIT. Rich blood sampling is collected after 2 weeks of treatment and sparse blood sampling at week 6 and 8. Venous blood samples are collected as well as finger pricks on dried blood spot (DBS). The final treatment outcome is registered after treatment completion.

added. Finally, 0.5 mL of the solution is transferred using a pipette to two labelled MGIT tubes, gently tilted for 1 min and incubated in the BACTEC MGIT 960 machine at 37°C. TTP is done in duplicate and is automatically recorded by the BACTEC MGIT 960.

Drug susceptibility testing

Routine DST is performed according to Chinese National Guidelines with the proportion method on Lowenstein-Jensen (LJ) medium, according to WHO's recommendations.³⁰

Figure 2 Blood collected through finger prick onto dried blood spot filter paper.

Simultaneous MIC determination of Mtb using TREK Sensititre broth microdilution plates

Since MIC testing in BACTEC MGIT is labour-intensive and time-consuming, a high throughput broth microdilution plate has been developed to test up to 12 antibiotics simultaneously in Middlebrook 7H9 on a single MIC plate. We have designed a custom-made Sensititre plate for the drugs used at the study site, with concentration ranges including wild-type isolates, manufactured by Thermofisher (figure 3). The reference isolate H37Rv ATCC 27294 is always included in each test run and compared with previously published quality control target ranges for each drug.³¹ The Thermofisher Sensititre MYCOTB plate is used for internal validation of

ethionamide (range 0.5 mg/L–32 mg/L), which was not stable in the pretrial validation of the customised plate.

After positive culture of Mtb in the BACTEC MGIT 960 and recording of TTP, the isolates are stored at -80°C awaiting MIC determination. After thawing and reculturing on LJ media, bacterial suspensions are prepared from Mtb isolates which are no more than 2 weeks old. Bacterial suspension together with Middlebrook 7H9 stock solution are then added to each well, according to the manufacturer's instructions.³² The plates are sealed and left to incubate in 37°C . Manual reading is done after 10–21 days, depending on growth, assisted by an inverted mirror (figure 4).

Due to specific pH requirements, PZA susceptibility is determined using BACTEC MGIT 960 PZA Susceptibility Test, with a pH of 5.9 as previously described.³³ In short, colonies of Mtb no older than 2 weeks are suspended in Middlebrook 7H9 broth with phosphate-buffered saline. A bacterial suspension, corresponding to a McFarland turbidity of 0.5, is prepared. Following the test protocol provided by the manufacturer (Becton Dickinson Biosciences, Sparks, Maryland, USA), the suspension is thereafter diluted 1:5 (inoculum A), from which a 1:10 diluted control is prepared (inoculum B). From inoculums A and B, 0.5 mL is then added to the MGIT 960 PZA tubes and the proportional growth control tube, respectively. The tubes are incubated in 37°C and read automatically by the BACTEC MGIT.

	1	2	3	4	5	6	7	8	9	10	11	12
A	OFL 16	MOXI 4	RIF 8	AMI 16	CAP 16	LEVO 8	PAS 16	ETH 16	EMB 16	CYK 64	KAN 16	INH 4
B	OFL 8	MOXI 2	RIF 4	AMI 8	CAP 8	LEVO 4	PAS 8	ETH 8	EMB 8	CYK 32	KAN 8	INH 2
C	OFL 4	MOXI 1	RIF 2	AMI 4	CAP 4	LEVO 2	PAS 4	ETH 5	EMB 5	CYK 16	KAN 4	INH 1
D	OFL 2	MOXI 0,5	RIF 1	AMI 2	CAP 2,5	LEVO 1	PAS 2	ETH 4	EMB 4	CYK 8	KAN 2	INH 0,5
E	OFL 1	MOXI 0,25	RIF 0,5	AMI 1	CAP 2	LEVO 0,5	PAS 1	ETH 2	EMB 2	CYK 4	KAN 1	INH 0,25
F	OFL 0,5	MOXI 0,12	PTH 0,5	AMI 0,5	CAP 1	LEVO 0,25	PAS 0,5	ETH 1	EMB 1	PTH 1	KAN 0,5	INH 0,12
G	OFL 0,25	MOXI 0,06	PTH 0,25	AMI 0,25	CAP 0,5	LEVO 0,12	PAS 0,25	ETH 0,5	EMB 0,5	PTH 2	KAN 0,25	PTH 2,5
H	OFL 0,12	MOXI 0,03	PTH 0,12	AMI 0,12	CAP 0,25	LEVO 0,06	PAS 0,12	ETH 0,25	EMB 0,25	PTH 4	POS	POS

Figure 3 Customised Sensititre broth microdilution plate (CML1FSWE). The wells are prefilled with antibiotics and Middlebrook 7H9 in predetermined concentrations (mg/L) for minimum inhibitory concentration determination. AMI, amikacin; CAP, capreomycin; CYK, cycloserine; EMB, ethambutol; ETH, ethionamide; INH, isoniazid; KAN, kanamycin; LEVO, levofloxacin; MOXI, moxifloxacin; OFL, ofloxacin; PAS, para-aminosalicylic acid; PTH, prothionamide; RIF, rifampicin.

Figure 4 Example of minimum inhibitory concentration (MIC) determination using microdilution plate. The red ring denotes the first well of amikacin (AMI) with no visible growth, that is, the MIC of AMI of this *Mycobacterium tuberculosis* isolates is 1 mg/L. Please refer to Figure 3 for plate lay-out. The two green rings represent growth controls.

Whole genome sequencing

All baseline study isolates, as well as any viable isolate after at least 1 month of treatment or more, will be analysed using WGS to detect new resistance mutations. In brief, DNA is extracted from Mtb LJ cultures using a chloroform/CTAB (N-cetyl-N,N,N-trimethyl ammonium bromide)-based protocol,³⁴ transported to Sweden and sequenced using Illumina technology (Illumina, San Diego, California, USA).

Mapping to a set of resistant genes from the Mtb H37Rv reference genome (GeneBank accession nr NC_000962.3) and extraction of variants are performed in CLC Genomics Workbench 8 (Qiagen, Hilden, Germany) using the following filters: minimum coverage: 10x; minimum count of reads calling variants: 2; minimum frequency of reads calling variants: 10%; minimum frequency of reads calling variants in each direction: 5%. In addition, pyro-error variants in homopolymer regions with a minimum length of 3 and a frequency below 0.8 are removed. The remaining variants are then compared with our in-house database of resistance mutations.

Data analysis plan

Regular study monitoring is performed quarterly by the Swedish and Chinese researchers as well as biweekly reports from the study site. Study data from the case report forms are entered in EpiData with a range check by two independent researchers and results compared for coherence.

The distribution of AUC/MIC and C_{max}/AUC will be presented and visualised in graphs. The agreement between drug exposure in plasma and DBS will be assessed. An exploratory analysis of the PK/PD indices for key TB drugs, such as fluoroquinolones, in relation to sputum culture conversion, TTP and changes in TBscore II during treatment will be performed. Pharmacometric

modelling will assess the relationship between dose, concentrations and effect and population models will be applied, using the non-linear mixed-effects modelling software NONMEM (Icon Development Solutions, Ellicott city). Time-to-event data with censoring will be analysed using the Cox regression model, whereas binary outcomes will be analysed with logistic regression and continuous outcome with linear regression, if data are normally distributed. The validated Chinese value set of the quality of life tool EQ-5D-5L will be used and quality of life perception described.

For analysis of trends in drug exposure over time, the dependent nature of the data will be taken into account using mixed-effect models. Missing values will not be imputed. A p value of <0.05 will be considered as statistically significant.

Information of potential confounders such as age, gender, BMI, concomitant treatment and comorbidities and disease severity assessed by TBscore II will be collected and evaluated during data analysis. As this is a feasibility and hypothesis-generating study, no power calculation was performed.

Ethics and dissemination

The study is performed in accordance with Good Clinical Practice and the Declaration of Helsinki. Ethical approval was obtained.

Prior to the study start, a designated study team of nurses, doctors and laboratory staff participated in training workshops of the study protocol and ethical considerations, led by the main study investigators from Fudan University and Karolinska Institutet. Patients are informed about the study orally and in writing, including information that neither study participation nor study termination will result in any changes in their treatment. An informed consent is signed or, in the case of illiteracy, a fingerprint given under observation by a witness. A travel grant to enable follow-up is offered to all the study participants. The sum was set so as not to create financial motivation to accept study participation. All patients are treated according to standard of care at the designated MDR-TB hospital and patients' safety ensured by regular monitoring.

Extensive blood sampling is a sensitive issue in China and should be avoided in severely ill patients. Therefore, the number of blood samples collected have been reduced to a minimum for the estimation of the AUC. Moreover, extensive blood sampling should be minimised in severely ill patients. An intravenous line is inserted to minimise patient discomfort. The increased sputum sample collection is a potential hazardous risk for other patients, hospital and laboratory staff. Therefore, biosafety and awareness training, as well as an upgrade of biosafety equipment, have been implemented.

Dissemination

We aim to present our data in international conferences and to publish our results in a peer-reviewed journal.

regardless of study results. Any significant protocol amendments will be reported to the respective ethical boards in Sweden and China.

DISCUSSION

In this prospective observational cohort study, we present a comprehensive, translational approach to TDM studies in MDR-TB, likely to be of benefit in future trials in the area. Multiple blood sampling and individual MIC determination will enable exploration of AUC/MIC for MDR-TB drugs, a poorly investigated research area. In a key study, the level of peak drug concentrations and AUC of PZA, rifampicin and isoniazid strongly influenced treatment outcome, although no comparison with the Mtb MICs was performed.⁴ Bacterial MIC has also been found to influence treatment outcome of patients with MDR-TB, with a sixfold increased odds of failure when comparing MIC of gatifloxacin of ≤ 0.25 mg/L to 1 mg/L, although both concentrations are still regarded as susceptible.³⁵ There are very few tentative targets for most second-line TB drugs, although an AUC/MIC >100 for fluoroquinolones has been suggested.³⁶ So far, the tentative targets have not been correlated with clinical outcome. To our knowledge, this is the first study to assess both AUCs and individual MICs for the most commonly used second-line drugs in MDR-TB regimen.

Not only have optimal PK/PD targets not been established, the critical concentrations used for DST are poorly validated.³⁷ MIC determination provides more information of the level of the resistance, but it has the drawback of being time-consuming. Fortunately, commercial MIC plates are available, facilitating fast MIC determination and will be assessed in this study.³⁸ When interpreting individual MICs, the clinician should bear in mind the innate variability may be up to ± 1 twofold MIC dilution step, but often less in a meticulous laboratory, which impacts on PK/PD indices estimates. Furthermore, when results of MIC testing are reported, it is important to note that there is no reference method for MIC testing of Mtb.

A limitation of this study protocol is the limited target number of included patients, common with other studies in the field, mainly due to MDR-TB incidence in Xiamen and costly and cumbersome sampling procedures. This may not allow us to perform extensive analysis of PK/PD indices in relation to treatment outcome, especially since all patients are treated with multiple drugs. However, pharmacometric mathematical modelling and simulation enables reduced sample sizes in clinical trials³⁹ and may partly compensate for the limited number of study patients in our study. Also, we use markers of early clinical improvement using microbiological surrogate measurements, such as TTP and sputum culture conversion, due to the nature of long treatment periods and follow-up. Sputum culture conversion is an imperfect surrogate marker of the final treatment outcome but it is, nevertheless, commonly used in clinical trials and is a sign of clinical improvement.⁴⁰

This study will provide useful insights of the PK/PD relations in MDR-TB treatment and highlight the importance of individualised treatment, taking both drug concentrations and MICs and innovative surrogate markers of improvement into account. With a simultaneous method for drug concentration analysis and blood sample collection simplified through DBS, TDM would be more feasible in clinical practice, including low-resource and high-endemic settings. We hope that this study will inspire future randomised controlled studies for TDM for both drug susceptible and MDR-TB, including treatment groups such as children, pregnant women, diabetic and HIV-infected patients who are prone to altered PK characteristics.

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Contributors LDF, KN, YH, RZ, XZ, JW, JP, USHS, EE, J-WA, SH, BX, TS and JB designed the study. RK, WC, YH, CH, YL, YG, JW, XZ, TS, LDF, KN and MM developed the plan for the microbiological part of the study. USHS, EE, TS, XZ, YH, LDF, KN, JB, JK, WC and J-WA developed the pharmacokinetic part of the study. LDF wrote the first draft of the manuscript together with KN and YH. All authors contributed and approved the final version.

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Competing interests None declared.

Patient consent Not required.

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REFERENCES

- WHO, 2016. Global tuberculosis report. <http://apps.who.int/medicinedocs/documents/s23098en/s23098en.pdf>
- Eliasson E, Lindh JD, Malmström RE, *et al.* Therapeutic drug monitoring for tomorrow. *Eur J Clin Pharmacol* 2013;69(Suppl 1):25–32.
- Shrivastava S, Pasipanodya JG, Meek C, *et al.* Multidrug-resistant tuberculosis not due to noncompliance but to between-patient pharmacokinetic variability. *J Infect Dis* 2011;204:1951–9.
- Pasipanodya JG, McIlleron H, Burger A, *et al.* Serum drug concentrations predictive of pulmonary tuberculosis outcomes. *J Infect Dis* 2013;208:1464–73.
- Chideya S, Winston CA, Peloquin CA, *et al.* Isoniazid, rifampin, ethambutol, and pyrazinamide pharmacokinetics and treatment outcomes among a predominantly HIV-infected cohort of adults with tuberculosis from Botswana. *Clin Infect Dis* 2009;48:1685–94.
- Zuur MA, Bolhuis MS, Anthony R, *et al.* Current status and opportunities for therapeutic drug monitoring in the treatment of tuberculosis. *Expert Opin Drug Metab Toxicol* 2016;12:509–21.
- Mouton JW, Brown DF, Apfalter P, *et al.* The role of pharmacokinetics/pharmacodynamics in setting clinical MIC breakpoints: the EUCAST approach. *Clin Microbiol Infect* 2012;18:E37–E45.
- Andrews JM. Determination of minimum inhibitory concentrations. *J Antimicrob Chemother* 2001;48(Suppl 1):5–16.
- Nahid P, Dorman SE, Alipanah N, *et al.* Official American Thoracic Society/Centers for Disease Control and Prevention/Infectious Diseases Society of America Clinical Practice Guidelines: treatment of drug-susceptible tuberculosis. *Clin Infect Dis* 2016;63:e147–e195.
- Wilby KJ, Ensom MH, Marra F. Review of evidence for measuring drug concentrations of first-line antitubercular agents in adults. *Clin Pharmacokinet* 2014;53:873–90.
- McCallum AD, Sloan DJ. The importance of clinical pharmacokinetic-pharmacodynamic studies in unraveling the determinants of early and late tuberculosis outcomes. *International Journal of Pharmacokinetics* 2017;2:195–212.
- Guiastrennec B, Ramachandran G, Karlsson MO, *et al.* Suboptimal antituberculosis drug concentrations and outcomes in small and HIV-coinfected children in India: recommendations for dose modifications. *Clin Pharmacol Ther* 2017.
- Mpagama SG, Nduiso N, Stroup S, *et al.* Plasma drug activity in patients on treatment for multidrug-resistant tuberculosis. *Antimicrob Agents Chemother* 2014;58:782–8.
- Gumbo T, Pasipanodya JG, Wash P, *et al.* Redefining multidrug-resistant tuberculosis based on clinical response to combination therapy. *Antimicrob Agents Chemother* 2014;58:6111–5.
- Peloquin CA, Hadad DJ, Molino LP, *et al.* Population pharmacokinetics of levofloxacin, gatifloxacin, and moxifloxacin in adults with pulmonary tuberculosis. *Antimicrob Agents Chemother* 2008;52:852–7.
- Vu DH, Alffenaar JW, Edelbroek PM, *et al.* Dried blood spots: a new tool for tuberculosis treatment optimization. *Curr Pharm Des* 2011;17:2931–9.
- Vu DH, Bolhuis MS, Koster RA, *et al.* Dried blood spot analysis for therapeutic drug monitoring of linezolid in patients with multidrug-resistant tuberculosis. *Antimicrob Agents Chemother* 2012;56:5758–63.
- Vu DH, Koster RA, Alffenaar JW, *et al.* Determination of moxifloxacin in dried blood spots using LC-MS/MS and the impact of the hematocrit and blood volume. *J Chromatogr B Analyt Technol Biomed Life Sci* 2011;879:1063–70.
- Epstein MD, Schluger NW, Davidow AL, *et al.* Time to detection of Mycobacterium tuberculosis in sputum culture correlates with outcome in patients receiving treatment for pulmonary tuberculosis. *Chest* 1998;113:379–86.
- Wallis RS, Pai M, Menzies D, *et al.* Biomarkers and diagnostics for tuberculosis: progress, needs, and translation into practice. *Lancet* 2010;375:1920–37.
- Rudolf F, Lemvik G, Abate E, *et al.* TBscore II: refining and validating a simple clinical score for treatment monitoring of patients with pulmonary tuberculosis. *Scand J Infect Dis* 2013;45:825–36.
- WHO, 2017. WHO tuberculosis country profiles. <http://www.who.int/tb/country/data/profiles/en/>
- von Elm E, Altman DG, Egger M, *et al.* Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. *BMJ* 2007;335:806–8.
- Prevention CISfDCa. 2017.
- Veringa A, Sturkenboom MGG, Dekkers BGJ, *et al.* LC-MS/MS for Therapeutic Drug Monitoring of anti-infective drugs. *TrAC Trends in Analytical Chemistry* 2016;84:34–40.
- Luo N, Liu G, Li M, *et al.* Estimating an EQ-5D-5L value set for China. *Value Health* 2017;20:662–9.
- U.S. Department of Health and Human Services FaDA, Center for Drug Evaluation and Research (CDER), Center for Veterinary Medicine (CVM), 2018. Bioanalytical method validation - guidance for industry 2018. <https://www.fda.gov/downloads/drugs/guidances/ucm070107.pdf>
- European Medicines Agency, 2011. Guideline on bioanalytical method validation. http://www.ema.europa.eu/ema/index.jsp?curl=pages/regulation/general/general_content_001280.jsp&mid=WC0b01ac0580032ec5
- National health and family planning commission of China, 2008. Diagnostic criteria for tuberculosis. <http://www.nhfpcc.gov.cn/zhuz/s9491/200801/38801.shtml>
- Canetti G, Fox W, Khomenko A, *et al.* Advances in techniques of testing mycobacterial drug sensitivity, and the use of sensitivity tests in tuberculosis control programmes. *Bull World Health Organ* 1969;41:21–43.
- Kaniga K, Cirillo DM, Hoffner S, *et al.* A multilaboratory, multicountry study to determine MIC quality control ranges for phenotypic drug susceptibility testing of selected first-line antituberculosis drugs, second-line injectables, fluoroquinolones, clofazimine, and linezolid. *J Clin Microbiol* 2016;54:2963–8.
- Lee J, Armstrong DT, Ssengooba W, *et al.* Sensititre MYCOTB MIC plate for testing Mycobacterium tuberculosis susceptibility to first- and second-line drugs. *Antimicrob Agents Chemother* 2014;58:11–18.
- Werngren J, Alm E, Mansjö M. Non-*pncA* gene-mutated but pyrazinamide-resistant mycobacterium tuberculosis: why is that? *J Clin Microbiol* 2017;55:1920–7.
- van Soolingen D, Hermans PW, de Haas PE, *et al.* Occurrence and stability of insertion sequences in Mycobacterium tuberculosis complex strains: evaluation of an insertion sequence-dependent DNA polymorphism as a tool in the epidemiology of tuberculosis. *J Clin Microbiol* 1991;29:2578–86.
- Rigouts L, Coeck N, Gumusboga M, *et al.* Specific *gyrA* gene mutations predict poor treatment outcome in MDR-TB. *J Antimicrob Chemother* 2016;71:314–23.
- Ghimire S, Van't Bovenind-Vrubleuskaya N, Akkerman OW, *et al.* Pharmacokinetic/pharmacodynamic-based optimization of levofloxacin administration in the treatment of MDR-TB. *J Antimicrob Chemother* 2016;71:2691–703.
- Schön T, Jurén P, Giske CG, *et al.* Evaluation of wild-type MIC distributions as a tool for determination of clinical breakpoints for Mycobacterium tuberculosis. *J Antimicrob Chemother* 2009;64:786–93.
- Reynolds J, Heysell SK. Understanding pharmacokinetics to improve tuberculosis treatment outcome. *Expert Opin Drug Metab Toxicol* 2014;10:813–23.
- Karlsson KE. *Benefits of pharmacometric model-based design and analysis of clinical trials*: Uppsala University, 2010.
- Kurbatova EV, Cegielski JP, Lienhardt C, *et al.* Sputum culture conversion as a prognostic marker for end-of-treatment outcome in patients with multidrug-resistant tuberculosis: a secondary analysis of data from two observational cohort studies. *Lancet Respir Med* 2015;3:201–9.