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CHAPTER 5

**RIFAMPIN LEVELS IN DAILY
PRACTICE: THE ACCURACY
OF A SINGLE MEASUREMENT**

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ABSTRACT

BACKGROUND Measurement of rifampin levels is not part of routine practice. However, low levels are associated with failure of tuberculosis treatment. The clinical relevance of serum levels in daily practice is unclear. The objective was to evaluate rifampin serum concentrations and factors associated with insufficient concentrations.

METHODS Patients with at least one rifampin concentration drawn 3 hours after intake (C_3) between 2005 and 2014 were included. Data on demographic and clinical characteristics were collected, including side effects and dose adjustments. Two different criteria were used to define adequate concentrations (criterion 1: C_3 and $C_6 \geq 3$ mg/l; criterion 2: C_3 or $C_6 \geq 5$ mg/l).

RESULTS Of 63 patients, 66% and 76% had a sufficient level according to criterion 1 or 2, respectively. C_3 exceeded C_6 in most patients, while a late maximum was significantly associated with diabetes mellitus ($p=0.003$). A dose adjustment was made in 19% of cases, more frequently in patients with insufficient levels ($p=0.02$) or with ≥ 2 side effects ($p=0.03$).

CONCLUSION Rifampin levels varied but were mostly adequate and a single measurement at 3 hours after intake provided the required information in most cases, indicating that full AUC_{0-24} measurements could be limited to specific situations.

BACKGROUND

Tuberculosis (TB) remains one of the world's most important infectious threats, reflected by 1.8 million deaths in 2015, of which 0.4 million deaths among people living with HIV.¹ Hence, adequate treatment is paramount. Rifampin is a key drug in the first-line treatment of active or latent TB, due to its high activity against *Mycobacterium tuberculosis* with an MIC_{90} of ≤ 0.25 μ g/ml.

The treatment success rate, especially in new cases, is improving although treatment failure occurs in up to 14% of patients.⁵ While multiple factors, including poor treatment adherence, bacterial resistance and even drug quality, may contribute to treatment failure, drug dosage and insufficient concentrations are relevant in this regard. In a previous study, the risk of failure of long-term treatment was almost 9-fold higher in patients with low drug exposure, expressed as 24-hour area under the concentration time curve (AUC_{0-24}) for pyrazinamide, rifampin and/or isoniazid.⁶

That study and other data showed that insufficient serum concentrations may even result in development of drug resistance.^{6,7} Apart from the prescribed dose, drug exposure may be influenced by factors such as comorbidities, food intake and inter-individual differences in pharmacokinetics.⁷⁻¹² Therapeutic drug monitoring (TDM) of rifampin is not routinely performed and there is no consensus on adequate levels. In previous studies, rifampin serum concentrations at 2 hours (C_2) and at 6 hours (C_6) after intake have been used to approximate the peak level.¹³⁻¹⁵ A recent study found that the rifampin AUC_{0-24} in TB patients was predicted optimally using sampling at time points 1, 3, and 8 hours,¹⁶ which would be impractical for most outpatients or require availability of alternative methods such as dry blood spot analysis. During the past decades, a rifampin absorption test at our centre has consisted of measurement of serum concentrations at 0, 3 and 6 hours after intake, and only at the physician's request. The aim of the present study was to retrospectively evaluate the results of these absorption tests of rifampin regarding adequate levels, and factors associated with out of range serum concentrations.

STUDY POPULATION AND METHODS

Study population

The study population consisted of patients in whom one or more rifampin serum concentrations had been measured at Leiden University Medical Centre (LUMC), a tertiary care hospital, between October 2005 and May 2014. Demographic and clinical characteristics were collected from the medical charts, including age, sex, weight, country of origin, clinical diagnosis, comorbidity (HIV infection, present or past malignancy, liver disease, diabetes mellitus, chronic kidney failure, autoimmune disease(s) or other), pregnancy, concomitant medication, rifampin dose at the time of TDM, kidney and liver function, indication for TDM and side effects. Serum

concentrations of rifampin at 0, 3 and 6 hours after intake, time of blood sampling, possible dose change and results of possible repeated TDM were collected. Patients were excluded if only a trough level was available or if the clinical data could not be retrieved. The protocol of this retrospective study with anonymised data collection was evaluated by the Medical Ethics Committee of the LUMC and waived from the requirement of informed consent (protocol G16.017).

Criteria for interpretation of serum concentrations

As there are no uniform criteria for adequate rifampin levels, we used two different criteria. According to the original protocol used at our institution for several decades, the source of which could not be retrieved, serum levels of the sum of rifampin and desacetyl-rifampin ≥ 3 mg/l at 3 hours (C_3) and 6 hours (C_6) after intake were defined as adequate (criterion 1: C_3 and $C_6 \geq 3$) and clinical decisions therefore were only based on this criterion. As an alternative criterion, adequate absorption was defined as a single measurement of the sum of rifampin and desacetyl-rifampin ≥ 5 mg/l (criterion 2: C_3 or $C_6 \geq 5$) as is nowadays implemented in several institutions. The data were analysed according to both criteria.

Method of measurement of rifampin concentrations

Serum concentrations of rifampin and desacetyl-rifampin were measured by high performance liquid chromatography according to the method published by Chandhi et al.¹⁷ The method was linear in a concentration range of 0.5 mg/l up to at least 15 mg/l rifampin and/or desacetyl-rifampin. Accuracy was $>98.8\%$ and imprecision $<5.7\%$.

Statistics

Descriptive statistical parameters were used. To compare proportions or continuous values between two groups, two-way chi square tests (or Fisher's exact probability test in case of comparison of proportions including numbers <5), and ANOVA tests were used, respectively. Differences using two-sided testing were considered significant at $p < 0.05$. Statistical analysis was performed using IBM SPSS Statistics version 23.

RESULTS

Study population

Of 90 patients in whom rifampin levels had been determined, 63 met the inclusion criteria (15 were excluded because only a trough level had been measured and 12 because clinical data were unavailable). Patient characteristics are shown in *table 1*. The majority (42/63, 67%) were immigrants from TB endemic regions. Most patients

had one or more comorbidity, with autoimmune disease, chronic liver disease and malignancy being most frequent. The most frequent reason for TDM was control of compliance (52%), followed by suspected high (29%) or low concentration (6%). More than half of the patients had received rifampin for active TB and one-third for latent TB.

Serum rifampin concentrations

In 63 patients, a total of 138 rifampin concentrations (at 0, 3 and/or 6 hours) were available. Rifampin levels were not always available for all three time points (*table 2*). C_3 was available for all 63 patients, C_0 was available for 34/63 patients (54%) and C_6 for 41/63 patients (61%). According to the guidelines for TB treatment the standard dose of rifampin is 10 mg/kg, with a maximum of 600 mg. Most patients (45/63, 71.4%) were treated with a dose of 600 mg (*table 2*). The dose was 600 mg in 42/46 (91.3%) patients with a body weight ≥ 55 kg. The mean \pm SD dose per weight was 11.2 ± 3.9 mg/kg. Maximal rifampin levels did not differ according to dose per weight (data not shown). Maximal levels did not vary by any demographic or clinical parameter (*table 1*).

Trough levels were <2 mg/l in 31/34 patients (91.2%) and were 3.2 mg/l, 5.6 mg/l and 9.9 mg/l respectively in the remaining three patients. In the last of these three patients (patient 41 in *figure 1*), C_0 exceeded C_3 and C_6 and thus had most likely been measured after intake of rifampin. The average individual maximal concentration, which could be either at 3 or at 6 hours, was 8.9 mg/l (range 0.0 mg/l to 26.7 mg/l). With regard to criterion 1: C_3 and $C_6 \geq 3$, 41 patients could be evaluated. Criterion 1 was met in 27/41 (65.9%). Criterion 2: C_3 or $C_6 \geq 5$ was met in 48/63 patients (76.2%). There was no significant relation between age, sex, comorbidities, co-medication or indication for rifampin comorbidities and meeting the criteria or not. Levels in immigrant patients more frequently met criterion 2 than did those from native Dutch patients (86.4% vs 52.6%, $p=0.004$). *Figure 1* shows all individual rifampin concentrations, ranked by the value of C_3 which was available for all 63 patients. C_3 exceeded C_6 in all but 8 patients (case 2, 9, 12, 17, 18, 24, 46 and 53 in *figure 1*). C_6 was ≥ 5 mg/l and often even much higher in all of these eight patients with late maximal concentrations. In 7/8 patients criterion 1: C_3 and $C_6 \geq 3$ was also met. Of the eight patients with late maximal levels, four (50%) had diabetes mellitus and one additional patient suffered from systemic sclerosis. In the remaining three patients no factors associated with delayed absorption could be identified. The proportion of patients with diabetes in those with late maximal levels (4/8 patients with $C_6 > C_3$) was significantly different from that in patients with early maximal levels (1/33 patients with $C_3 > C_6$; Fisher's exact probability test $p=0.003$). In 12 patients (19%) rifampin measurements including at least C_3 were later repeated after a median interval of 11 days (range 1-50 days, and one outlier at 248 days) because of out of range first levels, newly experienced side effects and/or after adjustment of the dose based on initial levels. The results of paired individual maximal serum concentrations are shown in *figure 2*.

Side effects

At least one side effect was reported in 27/63 patients (42.8%). Side effects varied from mild to very severe, ranging from minor nausea to drug-induced hepatitis (table 2). The maximal rifampin level in patients experiencing side effects was not significantly different from that in patients without side effects. In the six patients with serum transaminases >100 IU/l, the maximal level was not different from that in patients without liver function disturbances.

Dose adjustments

Twelve out of 63 patients (19.0%) had a dose adjustment. Six of 15 patients (40%) who did not meet criterion 2 had a dose increase. Six of 48 patients (12.5%) meeting criterion 2 had a dose reduction. This difference in proportion with a dose adjustment was significant ($p=0.02$).

A dose adjustment was made in 5/13 patients who experienced ≥ 2 side effects, in 3/14 patients with one side effect and in 4/36 patients without side effects ($p=0.03$ for comparison of patients with ≥ 2 to those without side effects).

Of 12 patients who had a second measurement of the rifampin level, dose changes were reported in five (figure 2). In four of these, the maximal levels were adequate after a dose increase ($n=3$) or reduction ($n=1$).

Follow-up

None of the patients with active TB had treatment failure and none of the patients treated for latent TB infection and who later received immunosuppressive drugs had a TB reactivation during a follow-up time between two and ten years.

DISCUSSION

In the present study we retrospectively evaluated rifampin levels which had been determined in routine practice in a mixed population consisting mainly of patients treated for active or latent TB. The data showed considerable inter-individual variation but in the majority of patients serum levels were adequate as based on two different criteria, one of which had been in use for decades at our institution and an alternative criterion based on a single peak level of at least 5 mg/l, which is nowadays implemented in several Dutch institutions. Nevertheless, the dose was adjusted in 20% of patients because of either too low or very high levels. In most patients in whom both C_3 and C_6 were available, C_3 was highest and therefore most informative. Maximal serum levels were not affected by demographic parameters, the presence of comorbidities or use of co-medication.

Despite the recognition that adequate rifampin concentrations are crucial for treatment success, TDM is not common practice. In addition, there are no clear criteria for the interpretation of concentrations. Studies in animals showed that the AUC_{0-24} in steady state divided by the MIC was the best predictive parameter for efficacy of rifampin.^{18,19} In humans, treatment failure has been associated with low AUC_{0-24} , and with development of bacterial resistance.⁶⁷ In a population pharmacokinetic model in patients with active TB, the rifampin AUC_{0-24} could be predicted with high precision using sampling at 0, 1, 3, and 8 hours after intake.¹⁶ However, such timing is not practical for most outpatients and the investment of the patient's time and the costs must be weighed against the value of the information thus obtained. In a previous study a single measurement of rifampin at four hours after intake gave the best estimate for AUC_{0-24} .²⁰ While precise AUC_{0-24} of rifampin is generally not needed, there are specific situations in which such information can be essential, such as in patients with extensive TB and a high bacillary load, or in patients with TB meningitis because of limited penetration. In general practice there may also be reasons to measure rifampin levels, however without the need for a precise AUC_{0-24} , e.g. if treatment adherence is doubted, if poor absorption is suspected or because of suspected high levels. In these situations it may suffice to measure the concentration at the time of expected peak concentration. Because there is a large inter-individual variation in pharmacokinetics the peak value can be missed if just one sample is used. However, the results of the present study showed that C_3 almost always exceeded C_6 . This is in agreement with a peak between 1 and 3 hours (occasionally 4 hours) after intake in studies in which multiple time points were used, the peak being closer to 2 hours if the drug was taken without food and closer to 3 hours if taken with a light meal.^{16,21} Thus, if full AUC_{0-24} is not required a single measurement at 2 to 3 hours after intake may provide sufficient information. In the limited number of patients in the present study in whom C_6 exceeded C_3 , more than half had a disorder associated with delayed gastric emptying such as diabetes mellitus, and including a later time point should thus be considered in that setting. In accordance with our finding, in a previous study in Indonesian patients the AUC_{0-6} was about 50% lower in patients with diabetes compared with nondiabetic²² TB patients. Trough levels were not informative and our data suggest that these could be omitted.

Combining data from the literature with those from the present study, we designed a simple and practical algorithm for the selection of time points for measurement of rifampin concentrations (figure 3). We think that testing rifampin concentrations at just one time point in most patients, and more frequently only on indication, could save time and money without loss of quality of care. In the LUMC, based on this study the single measurement is now implemented for routine practice, while AUC_{0-24} is available if needed. Regarding the standard rifampin dose of 600 mg it has been argued that the 600 mg dose is at the lower end of the dose-response curve.²³ An update of the TDM in the treatment of tuberculosis of rifampin suggests higher doses to be more effective.²⁴ The pharmacokinetic profile of rifampin

is nonlinear and a dose increase will result in a greater than proportional increase in AUC. Previous studies using a higher rifampin dose of 13 mg/kg or 20 mg/kg did not observe increased hepatotoxicity or other adverse events.^{23,25-29} In a recent study even a 1200 mg dose was well tolerated,³⁰ indicating that a higher dose can probably be given without increasing the risk of side effects. Higher rifampin doses were evaluated in large clinical trials targeting C_{max} values ≥ 8 mg/l. Higher doses were associated with a better outcome and/or no increase of toxicity.³¹⁻³³ Boeree *et al.* even described a possibility of a shorter regimen of tuberculosis treatment with a higher dose (up to 35 mg/kg) of rifampin.³² A limitation of our study was the retrospective nature and the probable selection bias because rifampin levels were not routinely measured.

CONCLUSIONS

The results of this study show that in most cases a single rifampin level measured at 3 hours after intake provided sufficient information regarding adequacy of treatment. In the presence of risk factors for delayed absorption sampling at a later time point had added value. We think that a complete AUC₀₋₂₄ measurement can be limited to specific situations. Our findings could contribute to a cost-effective, rapid and patient-friendly approach to TDM of rifampin and to effective treatment. However, further studies in different populations and settings are needed to assess the generalisability of our findings.

REFERENCES

- 1 WHO. Global tuberculosis report 2016. WHO, Geneva, Switzerland;2017.
- 2 Bemer-Melchior P, Bryskier A, Drugeon HB. Comparison of the *in vitro* activities of rifapentine and rifampicin against Mycobacterium tuberculosis complex. *J Antimicrob Chemother.* 2000;46:571-6.
- 3 Heifets LB, Iseman MD. Determination of *in vitro* susceptibility of mycobacteria to ansamycin. *Am Rev Respir Dis.* 1985;132:710-1.
- 4 Luna-Herrera J, Reddy MV, Gangadharam PR. In-vitro and intracellular activity of rifabutin on drug-susceptible and multiple drug-resistant (MDR) tubercle bacilli. *J Antimicrob Chemother.* 1995;36:355-63.
- 5 Cox HS, Morrow M, Deuschmann PW. Long term efficacy of DOTs regimens for tuberculosis: systematic review. *BMJ.* 2008;336:484-7.
- 6 Pasipanodya JG, McIlleron H, Burger A, et al. Serum drug concentrations predictive of pulmonary tuberculosis outcomes. *J Infect Dis.* 2013;208:1464-73.
- 7 Srivastava 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.
- 8 Becker C, Dressman JB, Junginger HE, et al. Biowaiver monographs for immediate release solid oral dosage forms: rifampicin. *J Pharm Sci.* 2009;98:2252-67.
- 9 Peloquin CA, Jaresko GS, Yong CL, et al. Population pharmacokinetic modeling of isoniazid, rifampin, and pyrazinamide. *Antimicrob Agents Chemother.* 1997;41:2670-9.
- 10 Ruslami R, Nijland HM, Alisjahbana B, et al. Pharmacokinetics and tolerability of a higher rifampin dose versus the standard dose in pulmonary tuberculosis patients. *Antimicrob Agents Chemother.* 2007;51:2546-51.
- 11 Smythe W, Khandelwal A, Merle C, et al. A semimechanistic pharmacokinetic-enzyme turnover model for rifampin autoinduction in adult tuberculosis patients. *Antimicrob Agents Chemother.* 2012;56:2091-8.
- 12 Wilkins JJ, Savic RM, Karlsson MO, et al. Population pharmacokinetics of rifampin in pulmonary tuberculosis patients, including a semimechanistic model to describe variable absorption. *Antimicrob Agents Chemother.* 2008;52:2138-48.
- 13 Magis-Escurra C, van den Boogaard J, Ijdema D, et al. Therapeutic drug monitoring in the treatment of tuberculosis patients. *Pulm Pharmacol Ther.* 2012;25:83-6.
- 14 Peloquin CA. Therapeutic drug monitoring in the treatment of tuberculosis. *Drugs.* 2002;62:2169-83.
- 15 Prah JB, Johansen IS, Cohen AS, et al. Clinical significance of 2 h plasma concentrations of first-line anti-tuberculosis drugs: a prospective observational study-authors' response. *J Antimicrob Chemother.* 2015;70:321-2.
- 16 Sturkenboom MG, Mulder LW, de Jager A, et al. Pharmacokinetic Modeling and Optimal Sampling Strategies for Therapeutic Drug Monitoring of Rifampin in Patients with Tuberculosis. *Antimicrob Agents Chemother.* 2015;59:4907-13.
- 17 Chandi LS, van der Sijs IH, Guchelaar H-J. Bepaling van rifampicine en desacetyl rifampicine. *Ziekenhuisfarmacie.* 1998;1:71-2.
- 18 Gumbo T, Louie A, Deziel MR, et al. Concentration-dependent Mycobacterium tuberculosis killing and prevention of resistance by rifampin. *Antimicrob Agents Chemother.* 2007;51:3781-8.
- 19 Jayaram R, Gaonkar S, Kaur P, et al. Pharmacokinetics-pharmacodynamics of rifampin in an aerosol infection model of tuberculosis. *Antimicrob Agents Chemother.* 2003;47:2118-24.
- 20 Magis-Escurra C, Later-Nijland HM, Alffenaar JW, et al. Population pharmacokinetics and limited sampling strategy for first-line tuberculosis drugs and moxifloxacin. *Int J Antimicrob Agents.* 2014;44:229-34.
- 21 Acocella G. Pharmacokinetics and metabolism of rifampin in humans. *Rev Infect Dis.* 1983;5 Suppl 3: S428-S32.
- 22 Nijland HM, Ruslami R, Stalenhoef JE, et al. Exposure to rifampicin is strongly reduced in patients with tuberculosis and type 2 diabetes. *Clin Infect Dis.* 2006;43:848-54.
- 23 van Ingen J, Aarnoutse RE, Donald PR, et al. Why Do We Use 600 mg of Rifampicin in Tuberculosis Treatment? *Clin Infect Dis.* 2011;52:e194-e9.
- 24 Alsultan A, Peloquin CA. Therapeutic drug monitoring in the treatment of tuberculosis: an update. *Drugs.* 2014;74:839-54.
- 25 Acocella G, Pagani V, Marchetti M, et al. Kinetic studies on rifampicin. I. Serum concentration analysis in subjects treated with different oral doses over a period of two weeks. *Chemotherapy.* 1971;16:356-70.
- 26 Curci G, Bergamini N, Delli VF, et al. Half-life of rifampicin after repeated administration of different doses in humans. *Chemotherapy.* 1972;17:373-81.
- 27 Furesz S, Scotti R, Pallanza R, Mapelli E. Rifampicin: a new rifamycin. 3. Absorption, distribution, and elimination in man. *Arzneimittelforschung.* 1967;17:534-7.
- 28 Lobue P, Menzies D. Treatment of latent tuberculosis infection: An update. *Respirology.* 2010;15:603-22.
- 29 Nitti V, Delli VF, Ninni A, Meola G. Rifampicin blood serum levels and half-life during prolonged administration in tuberculosis patients. *Chemotherapy.* 1972;17:121-9.
- 30 Aarnoutse RE, Kibiki GS, Reither K, et al. Pharmacokinetics, tolerability and bacteriological response of 600, 900 and 1200 mg rifampicin daily in patients with pulmonary TB. *Antimicrob Agents Chemother.* 2017;61(11) pii: e01054-17.
- 31 Peloquin CA, Velasquez GE, Lecca L, et al. Pharmacokinetic Evidence from the HIRIF Trial To Support Increased Doses of Rifampin for Tuberculosis. *Antimicrob Agents Chemother.* 2017;61.
- 32 Boeree MJ, Heinrich N, Aarnoutse R, et al. High-dose rifampicin, moxifloxacin, and SQ109 for treating tuberculosis: a multi-arm, multi-stage randomised controlled trial. *Lancet Infect Dis.* 2017;17:39-49.
- 33 Boeree MJ, Diacon AH, Dawson R, et al. A dose-ranging trial to optimize the dose of rifampin in the treatment of tuberculosis. *Am J Respir Crit Care Med.* 2015;191:1058-65.

TABLE 1 Clinical characteristic and rifampin levels in 63 patients.

Characteristic	Categories	No. (%)	Maximal rifampin level (average \pm SD) in mg/l	P value
Sex	Men	37 (58.7)	8.6 \pm 4.9	0.5
	Women	26 (41.3)	9.5 \pm 6.0	
Age (range in years)	0-15	11 (17.5)	9.2 \pm 5.0	0.6
	16-30	13 (20.6)	9.5 \pm 4.9	
	31-45	12 (19.0)	9.0 \pm 5.9	
	46-60	14 (22.2)	7.8 \pm 4.3	
	61-75	11 (17.5)	10.4 \pm 6.9	
	>75	2 (3.2)	3.5 \pm 4.9	
Immigration	No	19 (30.2)	7.65 \pm 6.4	0.2
	Yes	44 (69.8)	9.5 \pm 4.8	
Region of origin	Western Europe	19 (30.2)	7.6 \pm 6.4	0.4
	Eastern Europe/Russia	4 (6.3)	5.6 \pm 2.7	
	Africa	19 (30.2)	9.8 \pm 5.6	
	Middle East	7 (11.1)	10.9 \pm 2.2	
	Asia (other than Middle East)	11 (17.5)	10.5 \pm 4.7	
	North and Central America	2 (4.5)	4.4 \pm 2.8	
	South America	1 (2.3)	9.9	
Comorbidities	None	8 (12.7)	7.4 \pm 3.5	0.4
	≥ 1	55 (87.3)	9.2 \pm 5.5	
	HIV	4 (6.3) ^a	4.5 \pm 1.8	
	Malignancy	13 (20.6)	11.4 \pm 6.4	
	Chronic liver disease	10 (15.9)	9.4 \pm 5.4	
	Diabetes mellitus	6 (9.5)	7.5 \pm 2.2	
	Pregnancy	4 (6.3)	9.6 \pm 7.0	
	Chronic kidney failure	3 (4.8)	9.3 \pm 2.3	
	Autoimmune disease	20 (31.7)	8.5 \pm 4.8	
	Other	29 (46.0)	9.6 \pm 6.4	
	No. of comorbidities ^b	0	16 (25.4)	
1		35 (55.6)	9.1 \pm 5.8	
2		11 (17.5)	9.2 \pm 4.5	
3		1 (1.6)	6.9	
Indication for rifampin	Active tuberculosis	35 (55.6)	9.3 \pm 6.0	0.9
	Latent tuberculosis	20 (31.7) ¹	8.3 \pm 4.5	
	IV catheter-related infection	6 (9.5)	8.8 \pm 5.6	
	Other	2 (3.2)	9.1 \pm 0.6	

HIV=human immunodeficiency virus; IV=intravascular. a: The sum of the comorbidities exceeds 63 (100%) as patients could have more than one comorbidity; b: based on the reported seven specific comorbidities as listed in this table, thus excluding the category of other comorbidities.

TABLE 2 Dose, side effect, available concentrations, interpretation and dose adjustments.

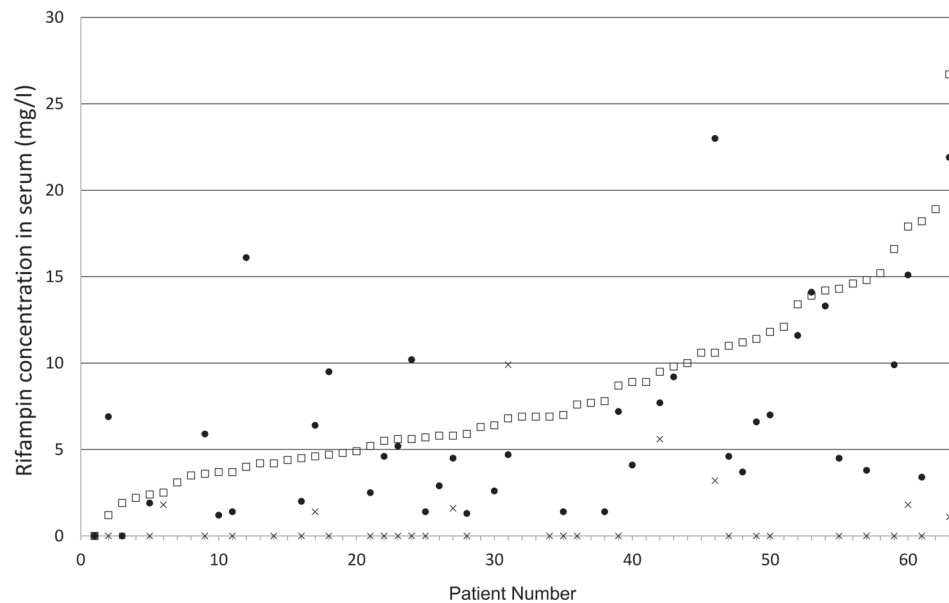
Parameter	Category	No. (%) ^a	
Dose (mg)	600	44 (70.1)	
	450	6 (9.7)	
	300	3 (4.8)	
	Other	9 (14.5)	
Side effects	≥ 1 side effect	27 (42.8)	
	≥ 2 side effects	13 (20.6)	
	General symptoms	19 (30.2) ^b	
	Gastrointestinal complaints	7 (11.1)	
	Drug induced hepatitis	6 (9.5)	
	Skin involvement	5 (7.9)	
	Headache	2 (3.2)	
Available rifampin levels	Neurological symptoms	1 (1.6)	
	Other	6 (9.5)	
	Only C ₃	18 (28.6)	
	Only C ₃ and C ₆	11 (17.5)	
	Only C ₀ and C ₃	4 (6.3)	
Criterion	C ₀ , C ₃ and C ₆	30 (47.6)	
	Yes	27/41 (65.9)	
	C ₃ and C ₆	\rightarrow dose change	2/27 (7.4)
≥ 3 mg/l ^c	No	14/41 (34.1)	
	\rightarrow dose change	3/14 (21.4) p =n.s.	
Criterion	Yes	48 (76.2)	
	C ₃ or C ₆	\rightarrow dose change	6/48 (12.5)
	≥ 5 mg/l	No	15 (23.8)
	\rightarrow dose change	6/15 (40.0) p =0.02	

a. Denominator was 63 unless otherwise specified

b. the sum of the side effects exceeds 27 as patients could have more than one side effect;

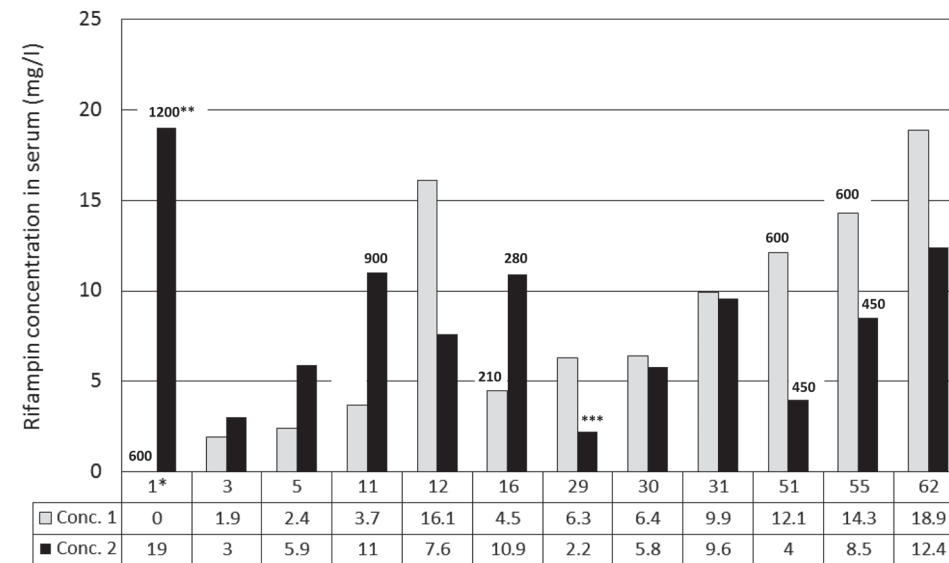
c. this criterion could only be tested for 41 patients for whom at least C₃ and C₆ were available

FIGURE 1 Distribution of rifampin levels in 63 patients, ranked by the concentration at 3 hours after intake.



Trough value is indicated by x; C₃ (concentration 3 hours after intake) is indicated by □; C₆ is indicated by •

FIGURE 2 Maximal rifampin levels in 12 patients in whom rifampin concentrations were measured twice.



Dose changes are indicated above the bars as dose in mg.

* The top row indicates the patient numbers corresponding to those used in figure 1.

** In patient 1 with initial undetectable rifampin concentrations, the maximal concentration was very high after doubling the dose, which suggested that rifampin may not have been taken at the time of first TDM.

*** In patient 29 the dose was increased from 500 mg to 600 mg based on the results of the repeated level.

FIGURE 3 The dotted line reflects the authors' opinion that retesting is generally not necessary if the clinical course is favourable but can be considered depending on the specific clinical situation.

