Linezolid is a promising antimicrobial agent for the treatment of multidrug-resistant tuberculosis (MDR-TB), but its use is limited by toxicity. Therapeutic drug monitoring (TDM) may help to minimize toxicity while adequate drug exposure is maintained. Conventional plasma sampling and monitoring might be hindered in many parts of the world by logistical problems that may be solved by dried blood spot (DBS) sampling. The aim of this study was to develop and validate a novel method for TDM of linezolid in MDR-TB patients using DBS sampling. Plasma, venous DBS, and capillary DBS specimens were obtained simultaneously from eight patients receiving linezolid. A DBS sampling method was developed and clinically validated by comparing DBS with plasma results using Passing-Bablok regression and Bland-Altman analysis. This study showed that DBS analysis was reproducible and robust. Accuracy and between- and within-day precision values from three validations presented as bias and coefficient of variation (CV) were less than 17.2% for the lower limit of quantification and less than 7.8% for other levels. The method showed a high recovery of approximately 95% and a low matrix effect of less than 8.7%. DBS specimens were stable at 37°C for 2 months and at 50°C for 1 week. The ratio of the concentration of linezolid in DBS samples to that in plasma was 1.2 (95% confidence interval [CI], 1.12 to 1.27). Linezolid exposure calculated from concentrations DBS samples and plasma showed good agreement. In conclusion, DBS analysis of linezolid is a promising tool to optimize linezolid treatment in MDR-TB patients. An easy sampling procedure and high sample stability may facilitate TDM, even in underdeveloped countries with limited resources and where conventional plasma sampling is not feasible.

Linezolid is used as a second-line drug in the treatment of multidrug-resistant tuberculosis (MDR-TB) due to its efficacy in vitro (21), in vivo (9), and in patients (1, 2, 14, 17, 34) against Mycobacterium tuberculosis. The World Health Organization (WHO) classifies linezolid as a reserve antituberculosis drug for the treatment of multidrug-resistant/extensively drug-resistant tuberculosis (MDR/XDR-TB) (33). Linezolid is usually added to a treatment regimen consisting of antituberculosis drugs to which Mycobacterium tuberculosis is still susceptible. However, treatment with linezolid may be limited by toxicity, such as time- and dose-dependent neuropathy or myelosuppression (17, 29), which leads to dose reduction or cessation of treatment with linezolid. Therapeutic drug monitoring (TDM) can be used to implement dose reductions to limit toxicity while preventing inadequate exposure. Efficacy predicting pharmacokinetic/pharmacodynamic (PK/PD) parameters, such as the ratio of the area under the concentration-time curve from 0 to 24 h to the MIC (AUC0–24/MIC), might be helpful in evaluating linezolid dosages in a murine model (32), but evidence from human data is lacking. Further PK/PD data from TB programs or large studies are needed for the development of evidence-based PK/PD parameters, such as an AUC0–24/MIC target ratio.

Linezolid treatment has been evaluated for TB treatment in several case series (17, 23). However, neither drug susceptibility tests (DSTs) nor drug exposure assessment was performed for linezolid, making it difficult to draw conclusions about efficacy (5). For instance, drug interactions with other antimicrobial agents might have occurred and might have had an impact on linezolid exposure (6, 15). In addition, conventional drug exposure evaluation for TB drugs using plasma samples might have been hindered in these studies by logistical challenges (30). The use of dried blood spot (DBS) sampling may provide a helpful alternative to conventional plasma sampling through a simplified sampling procedure and increased sample stability. DBS sampling has been applied in the treatment of other infectious diseases like malaria and HIV (30). Other advantages of DBS samples may include the requirement of a lower blood sample volume and lower biohazard risk than for conventional plasma samples (12, 18, 30). Compared to conventional sampling, DBS sampling may be hindered by inter- and intrapatient hematocrit (Hct) variation which leads to different blood viscosity values, yielding a proportional analytical bias with Hct values. Furthermore, Hct may affect the drug blood/plasma partition ratio, complicating the comparison with conventional plasma samples. In the development of a bioanalytical method for linezolid using DBS analysis, important patient-related factors like blood spot volume, Hct value (3, 24), and difference between capillary and venous blood have to be assessed during validation (12, 18, 25, 30). To enable individualized linezolid treatment, the aim of this study was to develop and
validate a method for DBS analysis and evaluate it in MDR-TB and XDR-TB patients.

**MATERIALS AND METHODS**

**Patients.** From September 2010 to March 2012, MDR-TB patients (≥18 years old) were recruited from the Tuberculosis Centre Beatrixoord, University Medical Center Groningen (Haren, The Netherlands). Eligible for inclusion were patients receiving treatment with antituberculosis drugs for which routine therapeutic drug monitoring was scheduled. Patients with bleeding disorders were excluded from the study. The study procedures were reviewed and approved by the local ethics committee. Patients receiving linezolid were included after providing written informed consent.

Sampling was performed at least 1 week after the start of linezolid treatment to ensure that the steady state was achieved. Venous blood samples were obtained before drug intake and at 1.2, 3, 4, and 8 h after dosing according to a previous study (2) and to local procedures for TDM of TB drugs to be able to calculate drug exposure and other PK parameters. Venous dried blood spot (VDBS) specimens were prepared by pipetting 50 μl of venous blood onto Whatman 31 ET CHR paper. The remaining venous blood was centrifuged at 3,000 rpm for 5 min at room temperature to obtain plasma, which was stored at −20°C until analysis. DBS specimens were obtained through a finger prick by dropping the blood directly on dried blood spot paper. DBS samples were obtained before drug intake and at 2 and 8 h after dosing, representing low, high, and medium linezolid blood levels, respectively. Both the VDBS and DBS samples were left to dry at room temperature and stored in sealed plastic bags with desiccant sachets at −20°C until analysis.

**DBS analysis.** To quantify DBS samples, an 8-mm-diameter disc was punched out of each blood spot. Extraction of these discs was performed by sonication with a frequency of 47 kHz for a period of 20 min using 500 μl of extracting solvent consisting of 0.3 mg/liter cyanoimipramine (internal standard) and 1g/liter EDTA in water. From this solution, a volume of 200 μl was added to 750 μl of acetonitrile. The samples were vortexed for 1 min and subsequently centrifuged at 11,000 rpm for 5 min. An injection volume of 5 μl was analyzed using a validated liquid chromatography-tandem mass spectrometry (LC-MS/MS) analysis method (16). The plasma samples were prepared and analyzed using the same method.

**DBS analytical method validation.** The DBS analytical method was validated in accordance with the U.S. Food and Drug Administration (FDA) guidance for bioanalytical method validation (27). For the validation, blood was prepared by mixing plasma and red blood cell and linezolid stock solution to achieve blood at the desired concentration and Hct level. Subsequently, the validation DBS samples were prepared by pipetting 50 μl of blood onto the paper. Linearity was assessed with 1/x2 weighting over a linezolid concentration range of 0.05 to 40 μg/liter. Clinical relevant concentrations were well within the range of the assay standards (2). The within-day and between-day accuracy and precision were evaluated on four validation levels, that is, lower limit of quantification (LLOQ), low, medium, and high, at linezolid concentrations of 0.05, 0.25, 15, and 30 mg/liter, respectively. Each validation level was analyzed on four validation levels, that is, lower limit of quantification (LLOQ), low, medium, and high, at linezolid concentrations of 0.05, 0.25, and 30 mg/liter, respectively. Each validation level was analyzed for 1 min and subsequently centrifuged at 11,000 rpm for 5 min. An injection volume of 5 μl was analyzed using a validated liquid chromatography-tandem mass spectrometry (LC-MS/MS) analysis method (16). The plasma samples were prepared and analyzed using the same method.

The DBS assay method showed linearity over the analytical concentration range. The pooled correlation coefficient, r2, was 0.9947. The regression equation is as follows: concentration = (0.1635 ± 0.0025) × response + (0.0001 ± 0.0003). Within-day and between-day accuracy and precision showed coefficients of variation (CVs) within accepted ranges. Within-day CVs ranged from 1.6% to 13.8%, and between-day CVs were from 3.5% to 10.2%. The mean measured concentration was within 98.7% to 106.3% of the nominal concentration. The bias caused by variable matrices, i.e., DBS and EDTA matrices, was less than 8.7%. The recovery of DBS extraction was between 94.1% and 97.2%. No significant linezolid deg-
radiation was observed after DBS samples were stored at 50°C for at least 1 week and at 37°C or ambient temperature for 2 months as biases were less than 15%.

Variation of blood spot volume between 30 μl to 90 μl had a minor impact on the assay accuracy as the bias ranged from −11.6% to 7.1%. The variation of Hct levels from 20% to 50% yielded biases within −7.6% to 6.8% and −12.5% to 5.7% for medium and high linezolid concentration levels. Larger biases of −17.8% to 11.9% were observed at the low concentration level (0.25 mg/liter) (Table 2).

**Comparisons of DBS, VDBS, and plasma analyses.** Significant proportional biases were observed in Passing-Bablok regressions in which the slope of regression line for linezolid concentrations between DBS samples and plasma was 1.28 (95% confidence interval [CI], 1.13 to 1.44) and between VDBS samples and plasma was 1.46 (95% CI, 1.40 to 1.54). The intercepts were −0.42 (95% CI, −1.72 to 0.17) and −0.67 (95% CI, −1.36 to −0.09), respectively (Fig. 1). In Bland-Altman analysis, the geometric mean linezolid concentration ratios in DBS and VDBS samples versus plasma were 1.20 (95% CI, 1.12 to 1.27) and 1.36 (95% CI, 1.32 to 1.40), respectively. The ratio of VDBS/plasma was higher than that of DBS/plasma (Wilcoxon signed-rank test, n = 24, P < 0.01). Limits of agreement of 95% were shown, with less than 5% of the values falling out of the ranges (Fig. 2).

**Pharmacokinetic and pharmacodynamic evaluation.** Median AUC_{0–12} values for linezolid in plasma were 50.9 (IQR, 50.5 to 54.9) mg · h/liter following a dose of 300 mg and 126 (IQR, 121.6 to 127.6) mg · h/liter following a dose of 600 mg. Linezolid pharmacokinetic parameters are shown in Table 3. The concentration–time curves for linezolid in plasma, DBS, and VDBS are presented in Fig. 3.
The AUC$_{0-12}$ values for linezolid in DBS and VDBS were calculated using the conversion factors 1.20 and 1.36 for DBS and VDBS, respectively. The subsequent results showed a good agreement with plasma. All the values were within the 95% limit of agreement (Fig. 4). The individual data for each patient for AUC$_{0-12}$ attained from plasma and converted (V)DBS concentrations and the respective AUC$_{0-24}$/MIC values are presented in Table 4. Patients that received a linezolid dose of 300 mg twice daily ($n = 3$) had a median AUC$_{0-24}$/MIC ratio in plasma of 236 (IQR, 219 to 322), and patients that received 600 mg twice daily ($n = 5$) had a median AUC$_{0-24}$/MIC ratio in plasma of 508 (IQR, 486 to 1,398).

**DISCUSSION**

This study showed that DBS analysis is an easy tool to individualize MDR-TB treatment with linezolid. In addition, this report presents a novel, validated method of analysis of linezolid in dried blood spots, with specimens that proved to be very stable over time.

In previous studies on DBS analysis of other drugs, several technical factors were pointed out that have to be considered when interpreting DBS analysis, such as the effect of Hct and blood spot volume (12, 13, 18, 30). For the analysis of linezolid in DBS samples, the effect of Hct seemed to be of minor concern. In this study, biases fell within accepted ranges for Hct values between 20 and 50%. These Hct values cover an even broader range than clinical Hct values found in TB patients in literature, i.e., 35.4% ± 6.7% (3) and in this study 37.4% ± 4.4%. Based on these findings, the standardization of Hct at 35% during DBS validation is acceptable. Furthermore, variation of blood spot volume between 30 and 90 μl had little effect as biases were within 15%.

Despite the minor influence of technical factors, i.e., Hct value and blood spot volume, physiological factors are also mentioned in literature to possibly limit the applicability and interpretation of DBS analysis (13). Such a factor might be differences between the concentration of linezolid in plasma and that in whole blood. This study shows that concentration of linezolid is higher in blood than in plasma. This is caused by different capacities of binding to plasma proteins and blood cells. Furthermore, concentrations of linezolid were higher in VDBS than in DBS samples. This might be caused by differences between the capillary and venous blood (13, 25, 30). Nevertheless, the linezolid concentrations in both DBS and VDBS specimens showed good correlation with the plasma concentration. To compensate for these differences, we propose conversion factors of 0.83 (1/1.20) for DBS and 0.74 (1/1.36) for VDBS to calculate corresponding plasma values. After the conversion, good agreement between the AUC$_{0-12}$ h values for linezolid in DBS samples and plasma was observed.

A meta-analysis showed that a $\geq$600-mg linezolid daily dose resulted in lower frequency of either an adverse event or adverse events necessitating treatment discontinuation than a dose of

![FIG 2 Bland-Altman plot of linezolid concentration ratios in DBS and VDBS samples versus plasma: solid line, mean ratio; dashed line, limit of agreement (mean ratio ± 1.96 × standard deviation of the ratio).](http://aac.asm.org/)

![FIG 3 Concentration-time curves of linezolid in plasma (●, solid line), VDBS samples (○, dashed line), and DBS samples (□, dotted line). Plasma and VDBS data are presented as means and standard deviations. For visual purposes, the DBS data are presented as means without error bars.](http://aac.asm.org/)

![FIG 4 Bland-Altman plot of AUC$_{0-12}$ for linezolid from corrected DBS versus AUC$_{0-12}$ plasma samples. Solid line, mean difference; dashed line, limit of agreement (mean difference ± 1.96 × standard deviation of the difference).](http://aac.asm.org/)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value for the parameter by linezolid dose$^a$</th>
</tr>
</thead>
<tbody>
<tr>
<td>AUC$_{0-12}$ (mg · h/liter)</td>
<td>50.9 (50.5–54.9) 126.9 (121.6–127.6)</td>
</tr>
<tr>
<td>C$_{max}$ (mg/liter)</td>
<td>8.8 (7.8–8.9) 16.5 (14.4–16.5)</td>
</tr>
<tr>
<td>T$_{max}$ (h)</td>
<td>1.9 (1.9–4.8) 1.9 (1.7–3.0)</td>
</tr>
<tr>
<td>t$_{1/2}$ (h)</td>
<td>4.6 (4.0–6.9) 7.5 (7.3–7.9)</td>
</tr>
<tr>
<td>Cl (liters/h)</td>
<td>4.9 (3.8–5.1) 3.1 (3.0–3.1)</td>
</tr>
<tr>
<td>V$_{d}$ (liters)</td>
<td>32.6 (29.4–34.4) 34.8 (32.9–41.6)</td>
</tr>
</tbody>
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$^a$ Data are presented as medians (IQR). All patients ($n = 8$) received linezolid twice daily.
>600 mg daily (8). Among the published data, the lowest rate of adverse effects was observed with a dose of 300 mg once daily (17). Nevertheless, lowering the dose clearly results in lower exposure to the drug (2, 19). In addition, interpatient variability and possible drug-drug interactions may lead to under- or overexposure. Therefore, treatment with a fixed dose may be questionable (6, 11, 22, 26). The application of TDM for linezolid can help avoid under- or overexposure, which may occur in 30 to 40% of the cases (20).

In this study, all patients had Mycobacterium tuberculosis isolates with a linezolid MIC of ≤0.5 mg/liter. With a dose of 600 mg (n = 5) twice daily, very high AUC_{0-24}/MIC ratios were reached (10), so dose reductions could be implemented to prevent time- and dose-dependent toxicity. Furthermore, a high correlation of AUC_{0-24}/MIC values between converted values for DBS and plasma (Spearman’s rho = 0.976, n = 8) was observed. This suggests that TDM using DBS may result in interventions identical to those with conventional plasma sampling. Therefore, adaptive dosing of linezolid to prevent potential toxicity and to ensure therapeutic exposure is feasible using DBS sampling.

The high stability of DBS specimens can minimize the logistical burden of conventional sampling in limited-resource areas. With simple instructions, the DBS samples can be obtained easily and sent to equipped facilities for analysis by mail (12, 30). This could allow TDM in TB programs worldwide, including in resource-limited settings where an MDR/XDR-TB epidemic is a growing problem. TDM using DBS sampling for MDR/XDR-TB should be limited settings where an MDR/XDR-TB epidemic is a growing problem. TDM using DBS sampling for MDR/XDR-TB should be limited settings where an MDR/XDR-TB epidemic is a growing problem. TDM using DBS sampling for MDR/XDR-TB should be limited settings where an MDR/XDR-TB epidemic is a growing problem. TDM using DBS sampling for MDR/XDR-TB should be limited settings where an MDR/XDR-TB epidemic is a growing problem.

Since treatment of MDR/XDR-TB is long and complicated by adverse drug reactions, TDM of linezolid with DBS sampling could be used to optimize drug exposure during treatment. In conclusion, this study presents a novel, validated analysis of linezolid levels in critically ill patient receiving concomitant linezolid and rifampin. Pharmacotherapy 27:476–479.

## REFERENCES