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Pharmacokinetic Study of Tenofovir Disoproxil Fumarate Combined with Rifampin in Healthy Volunteers


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Tenofovir disoproxil fumarate (tenofovir DF) was studied in combination with rifampin in 24 healthy subjects in a multiple-dose, open-label, single-group, two-period study. All subjects were given tenofovir DF at 300 mg once a day (QD) from days 1 to 10 (period 1). From days 11 to 20 the subjects received tenofovir DF at 300 mg combined with rifampin at 600 mg QD (period 2). The multiple-dose pharmacokinetics of tenofovir (day 10 and 20) and rifampin (day 20) were assessed. The drug-related adverse events (AEs) experienced during this study were mostly mild. Only one grade 3 AE possibly or probably related to the treatment (raised liver enzyme levels) occurred during period 2; the subject was withdrawn from the study. Pharmacokinetic data for 23 subjects were thus evaluable. Point estimates for the mean ratios of tenofovir with rifampin versus tenofovir alone for the area under the concentration-time curve from time zero to 24 h (AUC0-24), the maximum concentration of drug in plasma (Cmax), and the minimum concentration of drug in plasma (Cmin) were 0.88, 0.84, and 0.85, respectively. The 90% classical confidence intervals for AUC0-24, Cmax, and Cmin were 0.84 to 0.92, 0.78 to 0.90, and 0.80 to 0.91, respectively, thus suggesting pharmacokinetic equivalence. Similarly, coadministration of rifampin and tenofovir DF did not result in changes in the values of the tenofovir pharmacokinetic parameters. For rifampin, the values of the pharmacokinetic parameters found in this study were comparable to those found in the literature, indicating that tenofovir DF has no effect on the pharmacokinetics of rifampin. In conclusion, adaptation of either the rifampin or the tenofovir DF dose for the simultaneous treatment of tuberculosis and human immunodeficiency virus (HIV) infection in HIV-infected patients is probably not required.

Coinfection with Mycobacterium tuberculosis and human immunodeficiency virus (HIV) is frequent, particularly in Africa and Asia (3, 14, 18). Simultaneous treatment of tuberculosis and HIV infection may lead to complex combination therapy. Rifampin is a drug of choice for the treatment of tuberculosis. Rifampin is known to have major pharmacokinetic interactions with HIV protease inhibitors and nonnucleoside reverse transcriptase inhibitors (8, 10, 12, 13, 16, 17). Tenofovir disoproxil fumarate (tenofovir DF) is the first drug from a new class of reverse transcriptase inhibitors that has been recently approved for use for the treatment of HIV infections in adults. However, no data are available regarding its pharmacokinetics in combination with tuberculosis drugs, in particular, rifampin. No influence of rifampin on the pharmacokinetics of tenofovir is expected, because both drugs are metabolized and eliminated in different ways. Tenofovir is eliminated unchanged by glomerular filtration and active tubular secretion (1, 6), while rifampin is extensively metabolized by intestinal and hepatic metabolism (4). However, a pharmacokinetic interaction cannot be excluded.

The clinical trial described here was designed to explore the pharmacokinetics of tenofovir DF with and without rifampin in an effort to establish whether there is a need to adjust the dosage of either medication when the two medications are used for the treatment of patients coinfected with M. tuberculosis and HIV.

MATERIALS AND METHODS

Study design. The present study was designed to evaluate the effect of 600 mg of rifampin on the pharmacokinetics of 300 mg of tenofovir DF and also to assess whether tenofovir DF has a substantial impact on steady-state exposure to rifampin. This study was a multiple-dose, open-label, single-group, two-period study with 24 healthy volunteers. First, the subjects received tenofovir DF at 300 mg once daily (QD) for 10 days (period 1). At study day 10, a steady-state 24-h pharmacokinetic curve was obtained for tenofovir. During the second period of the study (period 2), tenofovir DF at 300 mg was combined with rifampin at 600 mg QD, again for 10 days. At study day 20, 24-h steady-state pharmacokinetic curves were obtained for tenofovir and rifampin. During the study both tenofovir DF and rifampin had to be taken with breakfast. On the days prior to study days 9 and 19, the subjects reported to the study center for direct observation of dosing with the medications with a standardized breakfast. Subsequently, on the evenings of study days 9 and 19 the subjects remained at the study center for two overnight stays and remained at the study center until the mornings of study days 11 and 21, respectively. On days 9, 10, 11, 19, and 20 the subjects received a standardized breakfast of 550 kcal (two slices of white bread, 15 g of low-fat margarine, 14 g of jelly, 150 ml of orange juice, and 150 ml of skim milk). The medication was administered immediately after breakfast with 200 ml of tap water. All other meals and snacks on the pharmacokinetic study days were also standardized. When the subjects took the medication at home, study drugs were administered with breakfast (at least two and at most three slices of wheat bread).

No crossover design was used in this study because rifampin could lead to considerable carryover effects, due to its long-lasting cytochrome P450-inducing effect. To eliminate this effect a longer washout period would be necessary, but this would have significantly prolonged the duration of the study and would have
Rifampin concentrations were determined by using a previously described high-performance liquid chromatography method. The concentrations of the quality controls used were 2.85, 9.5, and 24 mg/liter. The intra- and interassay coefficients of variation were less than 1.1% for all quality controls. The lower limit of quantification was 0.50 mg/liter. Samples from the same subject were analyzed by use of the same standard curve.

Pharmacokinetic analysis. Pharmacokinetic parameters for tenofovir and rifampin were calculated by noncompartmental methods by use of the WinNonlin software package (version 4.1; Pharsight Corporation, Mountain View, Calif.) and the log/linear trapezoidal rule. On the basis of the individual plasma concentration-time data, the following pharmacokinetic parameters were determined: the area under the plasma concentration-time curve (AUC) from time zero to 24 h (AUC$_{\text{0-24}}$, in milligram · hour per liter), the maximum concentration of drug in plasma (C$_{\text{max}}$, in milligrams per liter), the time to reach C$_{\text{max}}$ (T$_{\text{max}}$, in hours), the minimum concentration drug in plasma (C$_{\text{min}}$, in milligrams per liter), the apparent elimination half-life (t$_{1/2}$, in hours), and the apparent oral clearance (CL/F; in liters per hour). AUC$_{\text{0-24}}$, where C$_{\text{AUC}}$ is the last quantifiable concentration, was calculated for rifampin. C$_{\text{max}}$ and CL/F were not calculated for rifampin.

Statistical analysis. Statistical analyses were performed with SPSS software (version 11.0; SPSS Inc., 1989 to 1999). Descriptive statistics were calculated with Excel 2000 software (Microsoft Corporation, 1985 to 1999). Evaluation of the AUC$_{\text{0-24}}$ and the C$_{\text{max}}$ of tenofovir was the main objective of this trial. These parameters are considered the primary characteristics for the extent and the rate of drug absorption, respectively. The bioequivalence of tenofovir was determined by comparing the values of the relevant pharmacokinetic parameters obtained with the test treatment (tenofovir DF and rifampin on study day 20) to those obtained with the reference treatment (tenofovir DF alone on study day 10) by using the following statistical methods. The AUC$_{\text{0-24}}$, C$_{\text{max}}$, and C$_{\text{min}}$ of tenofovir were reported for study day 10 and study day 20 together by use of the ratios of the values on study day 20/values on study day 10. The arithmetic mean and standard deviations are given for study day 20 and study day 10. The geometric mean ratios and 90% classical confidence intervals for AUC$_{\text{0-24}}$, C$_{\text{max}}$, and C$_{\text{min}}$ were calculated. Treatments were considered bioequivalent if the respective 90% confidence intervals for AUC$_{\text{0-24}}$, C$_{\text{max}}$, and C$_{\text{min}}$ were included within the bioequivalence range of 0.8 to 1.25 (20). The values of the pharmacokinetic parameters for rifampin were compared with data from the literature by the use of descriptive statistics. The study was powered for the tenofovir C$_{\text{max}}$ by using nQuery software, and a sample size of 15 was required to achieve an 80% power to reject the null hypothesis that the two treatments are not equivalent in favor of the alternative hypothesis that the means of the two treatments are equivalent when the expected difference is 0.000. By this approach, a sample size of 15 would provide a 93% power for AUC$_{\text{0-24}}$. By considering the possibility that the subjects would drop out and/or that some difficulties with sample or pharmacokinetic analysis with some subjects would occur, 24 subjects were enrolled in this study.

RESULTS

Demographics. Twenty-four subjects (13 males, 11 females) were enrolled in this trial. One male subject was black; all other subjects were Caucasian. The mean age of the subjects was 41 years (range, 20 to 63 years). The mean body weight was 77 kg (range, 58 to 97 kg), and the mean height was 1.75 m (range, 1.59 to 1.88 m).

Pharmacokinetics. The pharmacokinetic evaluation was based on data sets for subjects that completed the study on both study days (study days 10 and 20). Data for 23 subjects were included in the pharmacokinetic analysis of tenofovir and rifampin. Table 1 provides a summary of the values of the pharmacokinetic parameters for tenofovir, including the arithmetic means, geometric mean ratios, and 90% confidence interval estimates for the pharmacokinetic parameters for tenofovir alone (study day 10) and tenofovir in combination with rifampin (study day 20). The tenofovir AUC$_{\text{0-24}}$, C$_{\text{max}}$, and C$_{\text{min}}$ were lower in period 2 when tenofovir DF was coadministered with rifampin. However, the magnitudes of these differences were small, with geometric mean ratios (90% confidence intervals) of 0.88 (0.84 to 0.92), 0.84 (0.78 to 0.90), and
The data are from reference 15 and are for subjects who received rifampin with breakfast.

Values are means (standard deviations).

a The data are from reference 15 and are for subjects who received rifampin with breakfast.

b Values are arithmetic means ± standard deviations (geometric means), unless indicated otherwise.

c Values are medians (ranges).

FIG. 1. Plasma tenofovir concentrations. ■, tenofovir concentration on study day 10 (n = 23) after administration of 300 mg QD; ◆, tenofovir concentration on study day 20 (n = 23) after administration of 300 mg combined with rifampin at 600 mg QD. Data are presented as means, and error bars indicate standard deviations.
0.78 to 0.90 for $C_{\text{max}}$. By definition, bioequivalence was proven for AUC and $C_{\text{min}}$ but was only suggested for $C_{\text{max}}$.

The tenofovir DF dose used in this study (300 mg QD) is the dose recommended for the treatment of HIV infection in adults (11). The rifampin dose used (600 mg QD) is an accepted regimen for the treatment of tuberculosis in patients weighing more than 50 kg (7). A previous study (2) has shown that steady-state conditions for rifampin are generally achieved after the sixth daily dose of rifampin at 600 mg. To ensure the achievement of steady-state pharmacokinetics, subjects were given tenofovir DF combined with rifampin for 10 days before pharmacokinetic assessment.

The reason for the lower observed tenofovir levels is unknown. Several mechanisms could contribute to this interaction. Because tenofovir is not metabolized and is eliminated unchanged by a combination of glomerular filtration and active tubular secretion (1, 6), it is unlikely that the inducing effect of rifampin on hepatic and intestinal cytochrome P450 enzymes (especially CYP3A4) (8) is the mechanism responsible for this effect. This is supported by no apparent changes in the tenofovir $t_{1/2}$ and no clinically relevant effects of rifampin on the tenofovir $C_{\text{min}}$.

Similarly, as tenofovir minimally binds to proteins in human plasma or serum (<0.7 and 7.2%, respectively) (11), altered distribution is also probably not the mechanism responsible for the pharmacokinetic differences observed. As the decrease in the tenofovir $C_{\text{max}}$ was 16% while the decrease in AUC$_{0–24}$ was 12%, the cause may be in the process of tenofovir DF or tenofovir absorption. Rifampin has been shown to be an inducer of the efflux transporter P-glycoprotein (9). No information exists in the literature that P-glycoprotein plays a role in the process of absorption of tenofovir in vivo. However, van Gelder et al. (19) have described the transport of tenofovir DF by a P-glycoprotein-related efflux mechanism in the Caco-2 system.

AEs led to one discontinuation in this study; grade 3 elevations in hepatic enzyme levels were reported after the medication was stopped during period 2, when tenofovir DF was combined with rifampin. Liver disturbance is a well-known side effect of rifampin. Gastrointestinal disorders are well-known AEs of both tenofovir and rifampin and occurred in a total of 46% of the study subjects during both study periods. During period 2 all subjects reported discoloration of their urine, which is a well-known AE of rifampin (7).

Some additional considerations are important for the extrapolation of the results of this study to patients. First, it should be noted that all the participants in this study were healthy subjects. It cannot be excluded that the pharmacokinetics of tenofovir and rifampin are different in HIV-infected patients coinfected with $M. tuberculosis$ due to one or both of the diseases. Second, 23 of the 24 subjects of this study were Caucasian. Race might have an effect on the values of the pharmacokinetic parameters for tenofovir, although the available pharmacokinetic data do not indicate substantial differences with regard to race (11). Finally, the subjects in this study were given tenofovir DF and rifampin only, while HIV-infected patients coinfected with $M. tuberculosis$ are treated with other antiretroviral and tuberculostatic drugs, which can cause interactions.

In conclusion, the data from this study demonstrate that the addition of rifampin to tenofovir DF is well tolerated, and the small decrease in plasma tenofovir levels during combination treatment suggests that these drugs can be coadministered without the need for dose adjustments. This implies that standard doses should be a starting point for the use of these medications by HIV-infected patients. Additional pharmacokinetic studies in a clinical setting are warranted to confirm the findings of this study.

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