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No meaningful drug interactions with doravirine, lamivudine and tenofovir disoproxil fumarate co-administration

(111/150 characters)

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ABSTRACT [250/250 words]

Background: Doravirine (DOR) is a novel non-nucleoside reverse transcriptase inhibitor available as a single tablet and a three-drug combination with lamivudine (3TC) and tenofovir disoproxil fumarate (TDF) to treat HIV-1 infection. These analyses assessed pharmacokinetic (PK) interactions with co-administration.

Methods: Two trials were conducted. Study 1: two-period, fixed-sequence; 8 healthy participants; Period 1, DOR 100 mg followed by ≥ 7 -day washout; Period 2, TDF 300 mg once daily for 18 days, co-administration of DOR 100 mg on day 14. Study 2: three-period, crossover, 15 healthy participants; Treatment A, DOR 100 mg; Treatment B, 3TC 300 mg + TDF 300 mg; Treatment C, DOR 100 mg + 3TC 300 mg + TDF 300 mg; ≥ 7 -day washout between periods.

Results: Study 1: geometric mean ratios (GMRs) (90% confidence interval [CI]) of DOR $AUC_{0-\infty}$ and C_{24h} (DOR + TDF / DOR) were 0.95 (0.80, 1.12) and 0.94 (0.78, 1.12), respectively. Study 2: GMRs (90% CI) of DOR $AUC_{0-\infty}$ and C_{24h} (DOR + 3TC + TDF / DOR) were 0.96 (0.87, 1.06) and 0.94 (0.83, 1.06), respectively. GMRs (90% CI) of 3TC and tenofovir $AUC_{0-\infty}$ (DOR + 3TC + TDF / 3TC + TDF) were 0.94 (0.88, 1.00) and 1.11 (0.97, 1.28), respectively. Study drugs were generally well tolerated.

Conclusions: Multiple doses of TDF did not have a clinically meaningful effect on DOR PK. The PK of DOR were similar when administered alone or in combination with 3TC and TDF. DOR had no meaningful effect on the PK of 3TC and tenofovir.

INTRODUCTION

Human immunodeficiency virus (HIV) continues to be a major global health challenge, infecting more than 36.9 million people worldwide [1]. In 2017, approximately 1.8 million people became infected with HIV, and 0.9 million people died from acquired immunodeficiency syndrome (AIDS)-related causes globally [1]. Antiretroviral therapy (ART) has been seminal in reducing the morbidity and mortality associated with HIV type 1 (HIV-1) infection. There are over 25 agents available for use in seven major mechanistic classes of ARTs: nucleoside reverse transcriptase inhibitors (NRTIs), non-nucleoside reverse transcriptase inhibitors (NNRTIs), protease inhibitors, a fusion inhibitor, a C-C chemokine receptor type 5 antagonist, a CD4-directed post-attachment inhibitor, and integrase strand transfer inhibitors [2]. Current guidelines generally recommend three antiviral agents from at least two different mechanistic classes, as treatment with agents across classes has demonstrated sustained virologic response [2, 3]. Despite the array of therapies currently available, no single antiviral agent or combination of agents is appropriate for every person living with HIV, and there are often additional challenges in finding the most suitable treatment including issues with resistance, tolerability, unfavourable drug–drug interaction (DDI) profiles, high pill burden and/or unfavourable dosing frequency [2, 3].

NNRTIs were formerly the cornerstone of front-line therapy; however, as protease inhibitors and integrase strand transfer inhibitors offer a greater barrier to resistance, improved tolerability and more rapid viral suppression for people living with HIV, they are no longer primarily recommended in major international guidelines [2, 3]. Although the NNRTIs efavirenz and rilpivirine remain as alternative treatment options under particular clinical circumstances, efavirenz has a relatively high rate of central nervous system-related adverse events (AEs),

limiting its tolerability; and rilpivirine has lower virological efficacy, particularly in patients with high baseline HIV-1 RNA (>100,000 copies/ml) and low CD4+ T-cell counts (<200 cells/mm³) [2]. As such, an unmet medical need exists for improved ART, including new NNRTI agents with improved tolerability and efficacy compared with currently available drugs in this class.

Doravirine (DOR, MK-1439) is a novel NNRTI designed to overcome the common resistance mutations which can reduce the effectiveness of other antiretrovirals in this class. Preclinical studies have demonstrated DOR to be active against wild-type HIV-1, as well as the two most prevalent NNRTI-associated mutant viruses (K103N and Y181C substitutions) [4]. In two Phase III studies, DOR demonstrated robust and durable efficacy, and was generally well tolerated [5, 6]. In the first of these, DOR co-administered with lamivudine (3TC)/tenofovir disoproxil fumarate (TDF) was associated with fewer treatment-emergent central nervous system AEs compared with the combination of efavirenz and emtricitabine/TDF [6]. In the second study, which compared DOR to ritonavir-boosted darunavir when both were co-administered with investigator-selected NRTIs (TDF and emtricitabine or abacavir and 3TC), there were no clinically relevant differences in the incidence of specific AEs, with the exception of a higher incidence of diarrhoea in the darunavir group [5]. In both studies, DOR combination therapy was associated with a more favourable lipid profile and similar antiviral efficacy over 48 weeks of treatment [5, 6]. DOR 100 mg administered once daily is indicated for the treatment of HIV-1 infection in combination with other ARTs, including 3TC and TDF, and is available for use as a single tablet or in a fixed dose combination tablet with 3TC and TDF [7, 8].

DOR is cleared primarily by oxidative metabolism via cytochrome P450 (CYP)3A [9]. Thus, drugs that induce or inhibit CYP3A may affect DOR elimination; this interaction has been confirmed in clinical DDI studies with the antibiotics rifabutin and rifampin, the antifungal

ketoconazole, and the antiretrovirals ritonavir and efavirenz [10-14]. DOR was also shown to be a substrate for P-glycoprotein (P-gp) [9]; however, studies conducted to date revealed that P-gp does not have a significant role in DOR absorption or elimination, suggesting that the likelihood of P-gp affecting DOR pharmacokinetics (PK) is minimal [9]. *In vitro* studies demonstrated that DOR is not expected to have a meaningful impact on the PK of other compounds, including substrates of all major CYPs and drug transporter [15]. Clinical drug-interaction studies with CYP3A and transporter substrates demonstrated no substantive interactions [16-19].

As a commonly used NRTI with a well-characterized PK profile, 3TC is eliminated primarily via urinary excretion by active organic cationic secretion and is not a known perpetrator of DDIs [20]. TDF is another commonly used NRTI which, following absorption, is rapidly converted to its active metabolite, tenofovir, and cleared by renal elimination [21, 22]. Although tenofovir has been shown to reduce CYP1A substrate concentrations, it is not a substrate, inducer or inhibitor of CYP3A [21]. Tenofovir DDIs have been reported with didanosine, resulting in increased didanosine concentrations after co-administration [23] and with ritonavir-boosted and unboosted atazanavir, with co-administration resulting in decreased atazanavir plasma concentrations and increased tenofovir concentration [21, 24].

Based on the metabolic profiles of DOR, 3TC and TDF, a meaningful PK DDI is unlikely. However, due to the use of these three agents in combination, and the unexpected effects seen with TDF when co-administered with other antiretroviral agents, two clinical trials were conducted to further explore potential DDIs.

METHODS

Study design

Study 1 (protocol MK-1439-003) was an open-label, two-period, fixed-sequence study in 8 healthy male participants, conducted between 19 September and 23 November 2011. In Period 1, all participants received a single oral dose of DOR 100 mg after an overnight fast. After a washout of ≥ 7 days, Period 2 began; all participants received a daily dose of TDF 300 mg for 18 days with co-administration of a single dose of DOR 100 mg on day 14. All doses of TDF alone were administered within 30 minutes prior to or after a standard meal; on day 14, study drugs were co-administered in the fasted state.

Study 2 (protocol MK-1439-038) was an open-label, single-dose, randomized, three-period crossover study in 15 healthy participants, conducted in January 2015. In the three treatment periods, participants received the following in a randomized manner: (A) a single oral dose of DOR 100 mg; (B) co-administration of single oral doses of 3TC 300 mg and TDF 300 mg; and (C) co-administration of single oral doses of DOR 100 mg, 3TC 300 mg and TDF 300 mg. Study drugs were administered after an overnight fast. The washout period between drug administrations was ≥ 7 days.

The studies were conducted in accordance with principles of Good Clinical Practice and were approved by the appropriate institutional review boards (Study 1: Thomas Jefferson University IRB, Philadelphia, PA, USA; Study 2: the Institutional Review Board of Optimum Clinical Research Inc., Oshawa, Ontario, Canada) and regulatory agencies.

Study populations

Study 1 included healthy men, 18–50 years of age with a body mass index ≤ 35 kg/m². Study 2 included healthy men and women, 18–65 years of age with a body mass index of 19–33 kg/m². In both studies, participants with a history of clinically significant medical conditions, estimated creatinine clearance of ≤ 80 ml/min (based on Cockcroft–Gault equation), drug or alcohol abuse, recent smoking or positive test for HIV, or who were hepatitis B or C positive, were excluded. Concomitant medications were not permitted from 14 days or 5 half-lives prior to the start of the trials until trial completion (although participants could receive concomitant therapy and continue in the study if the sponsor and investigator agreed). Participants in both studies provided written, informed consent prior to any study-related procedures being performed.

Sample collection and plasma concentration determination

In Study 1, blood samples for assay of DOR plasma concentration were obtained pre-dose and up to 120 h following administration of DOR on day 1 (Period 1), and co-administration of DOR and TDF on day 14 (Period 2). In Study 2, blood samples were collected pre-dose and up to 72 h post-dose.

In both studies, DOR plasma concentrations were analysed by liquid–liquid extraction for analyte isolation followed by liquid chromatographic–tandem mass spectrometric (LC-MS/MS) detection using a validated method (MSD, Oss, Netherlands) [13]. The lower limit of quantitation was 1 ng/ml. The analytical range of the assay was 1.00–1,000 ng/ml. For Study 1, the inter-day accuracy of the quality control samples was 103.3–105.0%, and the inter-day precision was 3.3–5.2%. For Study 2, the inter-day accuracy was 97.0–99.5%, and the inter-day precision was 3.5–5.3%. In Study 2, following extraction, the plasma concentrations of 3TC and

tenofovir were determined by validated achiral LC-MS/MS detection methods (Pharma Medica Research, Inc., Mississauga, Ontario, Canada). The analytical ranges of the assays were 5.00–3,000 ng/ml for 3TC and 2.00–500 ng/ml for tenofovir. For 3TC, the inter-day accuracy of the quality control samples was 97.8–105.2%, and the inter-day precision was 0.9–3.3%. For tenofovir, the inter-day accuracy was 98.5–101.5%, and the inter-day precision was 1.0–2.2%.

PK evaluations

In Study 1, DOR area under the concentration–time curve from time 0 extrapolated to infinity ($AUC_{0-\infty}$), maximum plasma concentration (C_{max}), time to reach maximum plasma concentration (T_{max}) and the apparent terminal half-life ($t_{1/2}$, calculated as the quotient of the natural log of 2 (ln [2]) and apparent terminal elimination rate constant) were calculated using Phoenix[®] WinNonlin[®] (Version 6.3, Certara, Princeton, NJ, USA). The observed plasma concentrations at 24 h post-dose (C_{24h}) were obtained directly from plasma concentrations using SAS (Version 9.3; SAS Institute Inc., Cary, NC, USA). In Study 2, values of the same PK parameters as in Study 1 were calculated for DOR, 3TC and tenofovir using the non-compartmental approach in Phoenix[®] WinNonlin[®].

Safety and tolerability

Safety and tolerability were assessed in both studies by physical examinations, vital signs, laboratory assessments and AE monitoring.

Statistics

In both studies, the individual values of $AUC_{0-\infty}$, C_{max} and C_{24h} were ln-transformed prior to analysis and evaluated separately using a linear mixed-effect model. In Study 1, treatment was a

fixed effect and subject was a random effect. A two-sided 90% CI for the geometric mean ratio (GMR; DOR + TDF / DOR alone) was generated for DOR $AUC_{0-\infty}$, C_{max} and C_{24h} from the mixed-effect model. Tenofovir PK were not analysed. Descriptive statistics were provided for T_{max} and apparent $t_{1/2}$. Median values were reported for T_{max} while the geometric mean was reported for $t_{1/2}$.

In Study 2, $AUC_{0-\infty}$, C_{max} and C_{24h} were analysed using a linear mixed-effect model appropriate for a three-period, two-treatment crossover design with fixed-effect terms for treatment and period. An unstructured covariance matrix was used to allow for unequal treatment variances and to model the correlation between different treatment measurements within the same subject via the REPEATED statement SAS PROC MIXED. Kenward and Roger's method was used to calculate the denominator degrees of freedom for the fixed effects (DDFM=KR).

A two-sided 90% CI for the GMRs (DOR + 3TC + TDF / DOR alone) was generated for DOR $AUC_{0-\infty}$, C_{max} and C_{24h} .

In addition, 95% CIs were generated from the above mixed-effect model for geometric means by treatment for DOR $AUC_{0-\infty}$, C_{max} and C_{24h} . 3TC and tenofovir $AUC_{0-\infty}$, C_{max} and C_{24h} after co-administration of DOR 100 mg, 3TC 300 mg and TDF 300 mg were analysed in a similar manner.

RESULTS

Study populations

A total of 8 healthy male participants were enrolled in Study 1; one participant discontinued on day 11 in Period 2 due to an AE that was not study-drug related. A total of 15 participants were

enrolled in Study 2, all of whom completed the study. Demographics for participants from both studies are summarized in Table 1.

PK evaluations

Study 1: mean plasma concentration–time curves for DOR alone or co-administered after multiple doses of TDF are shown in Figure 1A. DOR PK summary statistics are listed in Table 2. The GMRs (90% CI) of DOR $AUC_{0-\infty}$ and C_{24h} (DOR + TDF / DOR alone) were 0.95 (0.80, 1.12) and 0.94 (0.78, 1.12), respectively. The GMR (90% CI) of DOR C_{max} was 0.80 (0.64, 1.01). T_{max} and apparent $t_{1/2}$ were similar between the two treatment groups.

Study 2: the mean plasma concentration–time profiles for DOR, 3TC and tenofovir following DOR or 3TC + TDF administration or DOR + 3TC + TDF co-administration are shown in Figure 1B–D. DOR, 3TC and tenofovir PK summary statistics are listed in Tables 2 and 3. The GMRs (90% CI) of DOR $AUC_{0-\infty}$ and C_{max} (DOR + 3TC + TDF / DOR alone) were 0.96 (0.87, 1.06) and 0.97 (0.88, 1.07), respectively. The GMRs (90% CI) of 3TC $AUC_{0-\infty}$ and C_{max} (DOR + 3TC + TDF / 3TC + TDF) were 0.94 (0.88, 1.00) and 0.92 (0.81, 1.05), respectively. GMRs (90% CI) of tenofovir $AUC_{0-\infty}$ and C_{max} (DOR + 3TC + TDF / 3TC + TDF) were 1.11 (0.97, 1.28) and 1.17 (0.96, 1.42), respectively. Individual PK ratios and corresponding GMR plots of DOR, 3TC and tenofovir with and without co-administration of companion agents are shown in Figure 2.

Safety

All treatment combinations were generally well tolerated. There were no serious AEs, events of clinical interest or deaths reported during the studies. All AEs were mild in intensity, of limited duration and resolved by the end of the study.

In Study 1, 3 of the 8 participants (37.5%) reported a total of 10 AEs, 4 of which were considered to be related to study treatment (fatigue, [DOR alone], headache [DOR alone], rash [TDF] and somnolence [DOR + TDF]). Headache was the only AE reported by more than one participant. One participant was discontinued on day 11 of Period 2 due to an AE that was not study-drug related.

In Study 2, 5 of the 15 participants (33.3%) reported a total of 5 AEs. One incidence of somnolence (DOR alone) and one of headache (DOR + 3TC + TDF) were considered to be related to study treatment.

DISCUSSION

There is a continuing need for improved therapeutics for the treatment of HIV-1 infection. DOR is a novel HIV-1 NNRTI that is indicated for use in combination with other antiretroviral agents, and as a fixed-dose regimen with 3TC and TDF as a complete regimen, for the treatment of HIV-1 infection in adults with no prior antiretroviral treatment history [7, 8]. Although the metabolic profiles of these agents do not suggest that there would be meaningful DDIs with co-administration, clinical investigation was pursued to further evaluate potential DDIs.

Data from the two studies reported here demonstrate that neither co-administration of multiple doses of TDF nor co-administration with single doses of 3TC + TDF (at the recommended therapeutic dose of 300 mg each for 3TC and TDF) have a clinically meaningful impact on DOR PK. This is evidenced by a lack of a meaningful effect on DOR $AUC_{0-\infty}$, C_{max} and C_{24h} , with $AUC_{0-\infty}$ and C_{24h} GMRs close to unity and C_{max} reduced by 20% following multiple doses of TDF. The minor reduction in DOR C_{max} is not anticipated to have any meaningful impact on DOR efficacy or safety, as a DOR Phase IIb trial demonstrated similar efficacy to efavirenz

across a range of doses from 25 to 200 mg [25]. The single-dose DDI assessment in Study 2 further supports a lack of interaction, with DOR $AUC_{0-\infty}$, C_{max} and C_{24h} all without clinically meaningful changes.

3TC and TDF are commonly administered together, without evidence of a meaningful interaction when co-administered [20, 21]. As such, Study 2 was designed with co-administration of 3TC + TDF without evaluation of each of the separate components. Data showed the lack of a meaningful effect of DOR on either 3TC or tenofovir PK. Tenofovir exposure and C_{max} increased slightly (by 11% and 17%, respectively) with co-administration of DOR. These changes are not clinically meaningful, based on drug-interaction effects and dosing recommendations for TDF [21]. The cause of the effect is unknown. It has been noted that tenofovir is a substrate of P-gp and breast cancer-resistant protein (BCRP) transporters [21]. Atazanavir and other HIV protease inhibitors, which are P-gp and BCRP inhibitors, modestly increase tenofovir plasma concentrations, although not to a clinically meaningful level, likely secondary to P-gp and BCRP inhibition [21, 26, 27]. However, *in vitro* observations with DOR have shown that it is not an inhibitor of P-gp, indicating that interactions between DOR and P-gp are unlikely to be the cause of the increases to plasma tenofovir levels in the current study [15].

Study 1 was designed to assess the impact of TDF at steady state on DOR PK to maximize any potential inductive or time-dependent effects of TDF. While Study 2 was conducted with single-dose administration only, no inductive effects by DOR, 3TC or tenofovir were anticipated and there is no time dependence for the PK of DOR, 3TC or tenofovir. Moreover, with Study 1 data demonstrating minimal effect of TDF on DOR PK, a single-dose assessment was considered an appropriate approach for Study 2 and is anticipated to be predictive of multiple-dose behaviour [28]. The results of these studies did not demonstrate any substantive effect and indicate that,

with multiple-dose administration, there would not be a meaningful PK DDI between these agents.

Administration of DOR, 3TC and TDF individually and in combination was generally well tolerated, providing further evidence of the tolerability of DOR alone and in combination with 3TC and TDF. The most common treatment-related AEs in both Study 1 and Study 2 were headache and somnolence, which have also been reported in DOR Phase III studies [5, 6]. The lack of DDIs between DOR and 3TC + TDF supports the fixed-dose, three-drug, single-tablet regimen (MK-1439A [DOR 100 mg/3TC 300 mg/TDF 300 mg]) that has been developed [29], and which was the formulation used in a recently reported Phase III study (discussed in the Introduction) [6].

In summary, multiple doses of TDF co-administered with a single dose of DOR did not have a clinically meaningful effect on the PK of DOR. DOR, 3TC and tenofovir PK were similar when administered alone or co-administered. Consequently, co-administration of the three drugs without dose adjustment is supported.

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DISPLAY ITEMS**Table 1.** Study population demographics

Characteristic	Study 1 (N=8)	Study 2 (N=15)
Gender, n (%)		
Male	8 (100)	7 (46.7)
Female	0	8 (53.3)
Age (years)		
Mean (range)	44.3 (36–50)	44 (23–56)
Body mass index (kg/m²)		
Mean (\pm standard deviation)	29.2 \pm 2.0	26.2 \pm 2.9
Race, n (%)		
Asian	0	2 (13.3)
Black or African American	6 (75.0)	3 (20.0)
White	2 (25.0)	10 (66.7)

Table 2. Plasma PK of DOR 100 mg when administered alone, with multiple-dose TDF 300 mg administered once daily for 14 days, or with single-dose TDF 300 mg and 3TC 300 mg to healthy participants

Study 1: DOR administered alone or with multiple-dose TDF									
PK parameter	DOR + TDF			DOR			DOR + TDF / DOR		
	N	GM	95% CI	N	GM	95% CI	GMR	90% CI	rMSE ^a
AUC _{0-∞} (h·μM) ^b	7	33.4	25.9, 43.2	8	35.3	27.5, 45.3	0.95	0.80, 1.12	0.162
C _{24h} (nM) ^b	7	547	430, 697	8	584	463, 738	0.94	0.78, 1.12	0.171
C _{max} (nM) ^b	7	1,310	965, 1,780	8	1,630	1,210, 2,190	0.80	0.64, 1.01	0.216
T _{max} (h) ^c	7	3.0	1.0, 7.9	8	2.5	0.5, 5.0			
t _{1/2} (h) ^d	7	15.4	25.0	8	14.4	24.7			

Study 2: DOR administered with single-dose TDF and 3TC									
PK parameter	DOR + 3TC + TDF			DOR			DOR + 3TC + TDF / DOR		
	N	GM	95% CI	N	GM	95% CI	GMR	90% CI	Intra-subject %CV ^e
AUC _{0-∞} (h·μM) ^b	15	37.7	28.7, 49.4	15	39.1	31.5, 48.6	0.96	0.87, 1.06	15.2
C _{24h} (nM) ^b	15	507	332, 774	15	541	390, 750	0.94	0.83, 1.06	19.6
C _{max} (nM) ^b	15	2,030	1,720, 2,400	15	2,090	1,810, 2,420	0.97	0.88, 1.07	15.1
T _{max} (h) ^c	15	2.0	1.0, 6.0	15	3.0	1.0, 4.0			
t _{1/2} (h) ^d	15	13.5	40.6	15	13.8	31.9			

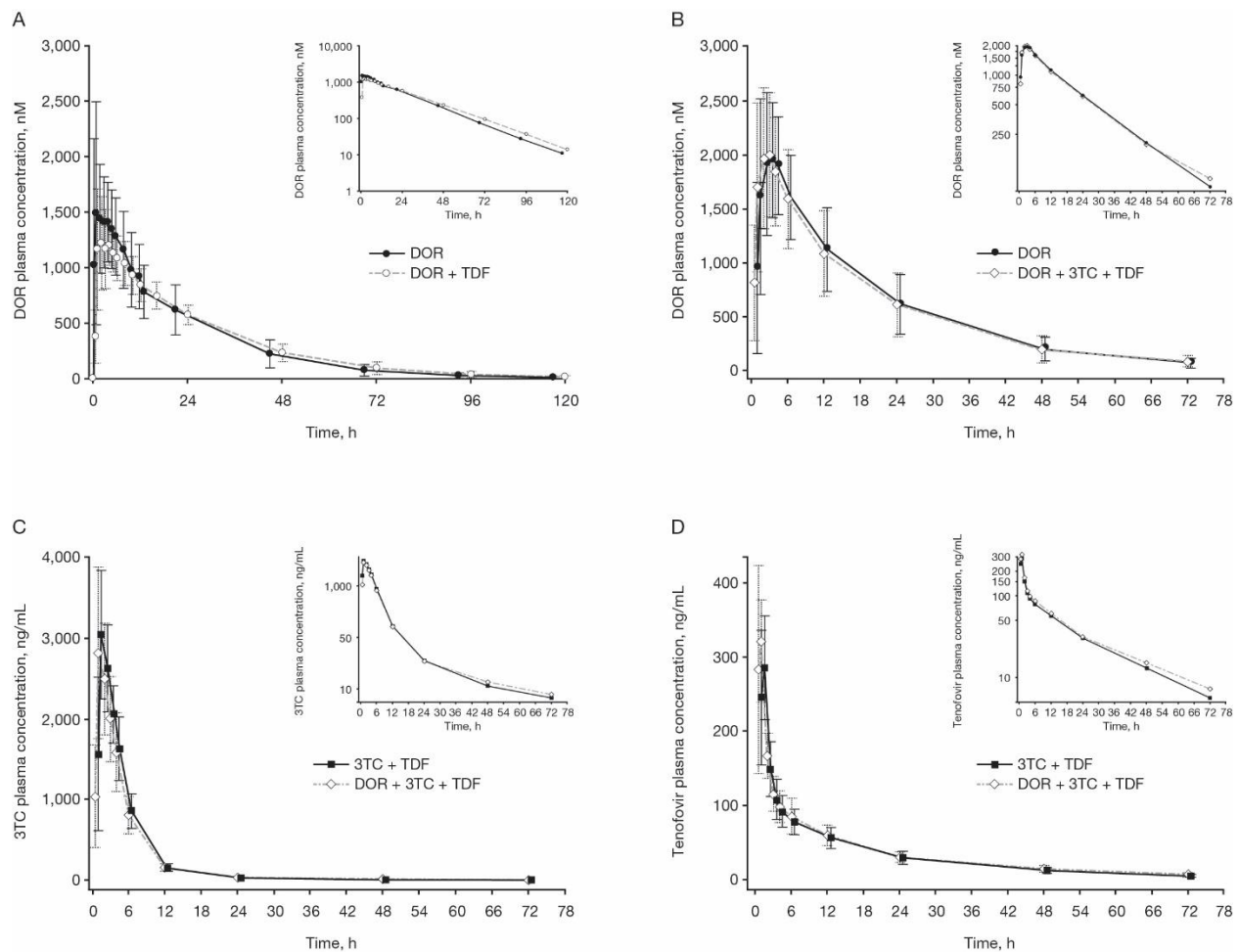
^arMSE (residual error) from the linear mixed-effect model. When multiplied by 100 approximates the within-subject %CV on the raw scale. ^bBack-transformed LSM and CI from linear mixed-effects model performed on natural log-transformed values. ^cMedian (minimum, maximum) reported for T_{max}. ^dGM and % geometric CV reported for t_{1/2}. ^eEstimated based on the elements of the variance-covariance matrix as: $CV(\%) = 100 \cdot \sqrt{(\sigma_A^2 + \sigma_B^2 - 2 \cdot \sigma_{AB})/2}$. 3TC, lamivudine; AUC_{0-∞}, area under the concentration–time curve from time 0 to infinity; C_{max}, maximum plasma concentration; C_{24h}, concentration of analyte in plasma 24 h after administration; CI, confidence interval; CV, coefficient of variation; DOR, doravirine; GM, geometric mean; GMR, geometric mean ratio; LSM, least-squares mean; PK, pharmacokinetic; rMSE, root mean square error; t_{1/2}, apparent elimination half-life; T_{max}, time to reach maximum plasma concentration; TDF, tenofovir disoproxil fumarate.

Table 3. Plasma PK of 3TC and tenofovir following single-dose administration of 3TC 300 mg + TDF 300 mg or DOR 100 mg + 3TC 300 mg + TDF 300 mg to healthy participants

3TC									
PK parameter	DOR + 3TC + TDF			3TC + TDF			DOR + 3TC + TDF / 3TC + TDF		
	N	GM	95% CI	N	GM	95% CI	GMR	90% CI for GMR	Intra-subject %CV^a
AUC _{0-∞} (h·ng/ml) ^b	15	14,200	12,400, 16,200	15	15,000	13,800, 16,500	0.94	0.88, 1.00	9.7
C _{max} (ng/ml) ^b	15	2,910	2,460, 3,450	15	3,150	2,760, 3,600	0.92	0.81, 1.05	19.4
T _{max} (h) ^c	15	1.00	1.0, 2.0	15	1.00	0.5, 2.0			
t _{1/2} (h) ^d	15	15.9	57.6	15	15.7	32.2			
Tenofovir									
PK parameter	DOR + 3TC + TDF			3TC + TDF			DOR + 3TC + TDF / 3TC + TDF		
	N	GM	95% CI	N	GM	95% CI	GMR	90% CI for GMR	Intra-subject %CV^a
AUC _{0-∞} (h·ng/ml) ^b	15	2,790	2,470, 3,150	15	2,500	2,090, 2,990	1.11	0.97, 1.28	20.5
C _{max} (ng/ml) ^b	15	338	286, 400	15	289	237, 352	1.17	0.96, 1.42	29.7
T _{max} (h) ^c	15	1.0	0.5, 2.0	15	1.00	0.5, 1.0			
t _{1/2} (h) ^d	15	20.9	18.5	15	19.7	12.6			

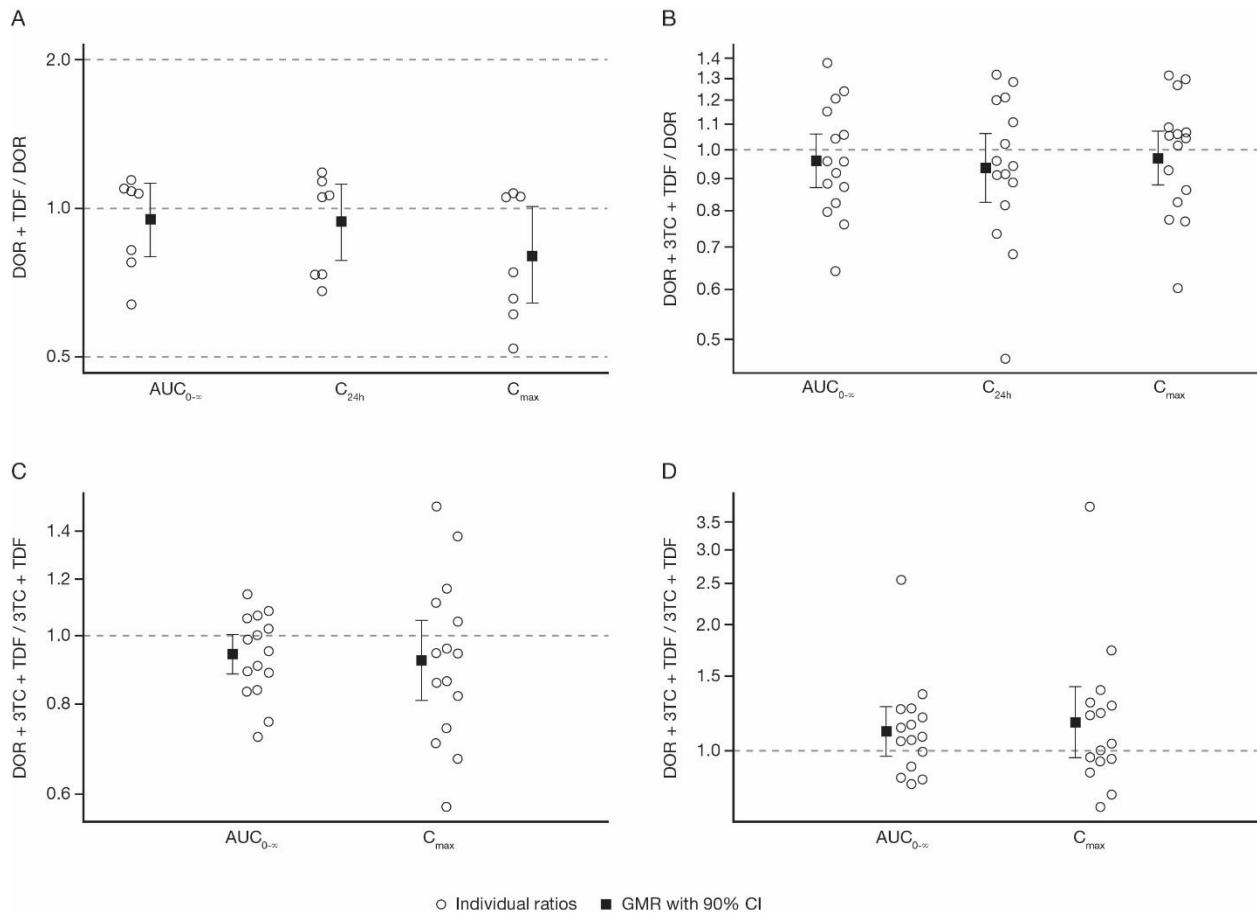
^aEstimated based on the elements of the variance-covariance matrix as: $CV(\%) = 100 \cdot \sqrt{[(\sigma_A^2 + \sigma_B^2 - 2 \cdot \sigma_{AB})/2]}$. ^bBack-transformed LSM and CI from linear mixed-effects model performed on natural log-transformed values. ^cMedian (minimum, maximum) reported for T_{max}. ^dGM and % geometric CV reported for t_{1/2}. 3TC, lamivudine; AUC_{0-∞}, area under the concentration–time curve from time 0 to infinity; C_{max}, maximum plasma concentration; CI, confidence interval; CV, coefficient of variation; DOR, doravirine; GM, geometric mean; GMR, geometric mean ratio; LSM, least-squares mean; PK, pharmacokinetic; t_{1/2}, apparent elimination half-life; T_{max}, time to reach maximum plasma concentration; TDF, tenofovir disoproxil fumarate.

Figure 1. Arithmetic mean (\pm standard deviation) plasma concentration–time profiles of (A) single-dose DOR 100 mg alone and co-administered with TDF 300 mg after 14 days of once-daily TDF administration ($N=8$, inset=semi-log scale); (B) single-dose DOR 100 mg alone and co-administered with single doses of 3TC 300 mg + TDF 300 mg; (C) single-dose 3TC 300 mg following administration of 3TC 300 mg + TDF 300 mg and DOR 100 mg + 3TC 300 mg + TDF 300 mg and (D) tenofovir following administration of 3TC 300 mg + TDF 300 mg and DOR 100 mg + 3TC 300 mg + TDF 300 mg ($N=15$, inset=log-linear scale for B, C and D)



3TC, lamivudine; DOR, doravirine; h, hours; TDF, tenofovir disoproxil fumarate.

Figure 2. Individual plasma pharmacokinetic ratios and corresponding geometric mean ratios with 90% confidence intervals for (A) single-dose DOR 100 mg with and without multiple-dose TDF 300 mg ($n=8$), (B) single-dose DOR 100 mg with and without single-dose 3TC 300 mg + TDF 300 mg ($n=15$), (C) 3TC after administration of single-dose 3TC 300 mg + TDF 300 mg, with DOR 100 mg versus without DOR ($n=15$) and (D) tenofovir after administration of 3TC 300 mg + TDF 300 mg, with DOR 100 mg versus without DOR ($n=15$)



3TC, lamivudine; AUC, area under the concentration–time curve; $AUC_{0-\infty}$, AUC from time 0 to infinity; C_{max} , maximum concentration; C_{24} , concentration of analyte in plasma 24 h after administration; DOR, doravirine; TDF, tenofovir disoproxil fumarate.