AMBER

A week-48 randomized phase-3 trial of darunavir/cobicistat/emtricitabine/tenofovir alafenamide in treatment-naïve HIV-1 patients

Eron JJ, Orkin C, Gallant J, Molina J-M, Negredo E, Antinori A, Mills A, Reynes J, Van Landuyt E, Lathouwers E, Hufkens V, Jezorwski J, Vanveggel S, and Opsomer M; on behalf of the AMBER study group.

***AIDS*. 2018;32:1431-1442.**

"Well powered, phase-3, double-blinded, randomized studies provide the most rigorous evidence to drive treatment guidelines. The week-48 virologic response rate (FDA-snapshot analysis) of 91.4% for D/C/F/TAF was among the highest achieved by a STR in phase-3 trials (range 80–93%) of ART-naïve patients and higher than in prior phase-3 trials with darunavir."

—Eron JJ, Orkin C, Gallant J, et al; on behalf of the AMBER study group

ART=antiretroviral therapy; C=cobicistat; D=darunavir; F=emtricitabine; STR=single-tablet regimen; TAF=tenofovir alafenamide.

INDICATION

SYMTUZA™ is indicated as a complete regimen for the treatment of human immunodeficiency virus type 1 (HIV-1) infection in adults:

- who have no prior antiretroviral treatment history or
- who are virologically suppressed (HIV-1 RNA less than 50 copies per mL) on a stable antiretroviral regimen for at least 6 months and have no known substitutions associated with resistance to darunavir or tenofovir.

IMPORTANT SAFETY INFORMATION

BOXED WARNING: POST TREATMENT ACUTE EXACERBATION OF HEPATITIS B

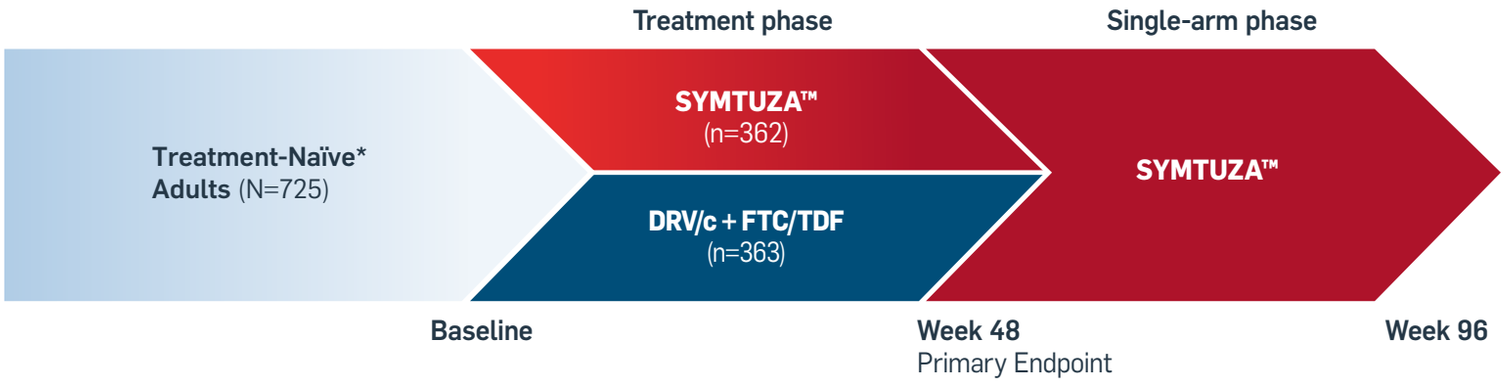
- **Severe acute exacerbations of hepatitis B (HBV) have been reported in patients who are coinfected with HIV-1 and HBV and have discontinued products containing emtricitabine and/or tenofovir disoproxil fumarate (TDF), and may occur with discontinuation of SYMTUZA™. Closely monitor hepatic function with both clinical and laboratory follow-up for at least several months in patients who are coinfected with HIV-1 and HBV and discontinue SYMTUZA™. If appropriate, anti-hepatitis B therapy may be warranted.**

Please see additional Important Safety Information throughout and full Prescribing Information, including Boxed WARNING, in pocket.

Symtuza™
darunavir/cobicistat/emtricitabine/tenofovir alafenamide tablets
800mg/150mg/200mg/10mg

Study Design

Phase 3, randomized, double-blind, active-controlled, international, multicenter, noninferiority study of SYMTUZA™ in treatment-naïve patients



Study Objectives

Primary endpoint:

- Proportion of patients with VL <50 copies/mL at Week 48 (noninferiority margin 10% by FDA Snapshot)

Selected secondary endpoints:

- Postbaseline HIV-1 genotypic resistance through 48 weeks
- Safety and tolerability

*Randomization was stratified by VL and CD4+.
DRV/c=darunavir/cobicistat; FTC/TDF=emtricitabine/tenofovir disoproxil fumarate; VL=viral load.

IMPORTANT SAFETY INFORMATION (continued)

CONTRAINDICATIONS

- Do not coadminister SYMTUZA™ and the following drugs due to the potential for serious and/or life-threatening events or loss of therapeutic effect: alfuzosin, carbamazepine, cisapride, colchicine (in patients with renal and/or hepatic impairment), dronedarone, elbasvir/grazoprevir, ergot derivatives (such as: dihydroergotamine, ergotamine, methylergonovine), lovastatin, lurasidone, oral midazolam, phenobarbital, phenytoin, pimozide, ranolazine, rifampin, St. John's wort *(Hypericum perforatum)*, sildenafil for pulmonary arterial hypertension, simvastatin, and triazolam.

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Patient Characteristics

	SYMTUZA™ (n=362)	Control (n=363)
Demographics, n (%), unless stated		
Median age, years (IQR)	34 (27-42)	34 (27-42)
>50	10% (36)	9% (32)
Gender		
Female	12% (44)	11% (41)
Male	88% (318)	89% (322)
Race		
White	83% (300)	83% (300)
Black	11% (40)	11% (40)
Other	6% (22)	6% (23)
Ethnicity		
Hispanic or Latino	14% (50)	12% (45)
Baseline disease characteristics		
Median time since diagnosis, months (IQR)	5.73 (2.53–25.59)	4.30 (2.07–17.74)
Median $\log_{10}$ VL, copies/mL (IQR)	4.44 (4.03–4.82)	4.57 (4.15–4.88)
VL ≥100,000 copies/mL	17% (60)	19% (70)
Median CD4+ count, cells/µl (IQR)	461.5 (342-617)	440.0 (325-594)
CD4+ count <200 cells/µl	6% (22)	8% (29)
Median $eGFR_{cr}$, mL/min Cockcroft–Gault (IQR)	119.3 (104.8–135.2)	118.4 (103.2–138.4)
Genotype at screening†	(n=361)‡	(n=362)‡
≥1 darunavir RAM	1% (3)	1% (4)
≥1 primary PI RAM	2% (7)	2% (8)
≥1 NRTI RAM	5% (18)	4% (16)
≥1 NNRTI RAM	15% (55)	17% (63)

†GenoSure® MG.
‡One patient in each group had failed screening genotypes and were enrolled based on local genotypes.

$eGFR_{cr}$=estimated glomerular filtration rate based on creatinine clearance; IQR=interquartile range; NNRTI=non-nucleoside reverse transcriptase inhibitor; NRTI=nucleoside reverse transcriptase inhibitor; PI=protease inhibitor; RAM=resistance-associated mutation; VL=viral load.

IMPORTANT SAFETY INFORMATION (continued)

WARNINGS AND PRECAUTIONS

- **Severe Acute Exacerbation of Hepatitis B in Patients Coinfected With HIV-1 and HBV:** Patients with HIV-1 should be tested for the presence of chronic HBV before initiating antiretroviral therapy. Severe acute exacerbations of hepatitis B (e.g., liver decompensation and liver failure) have been reported in patients who are coinfected with HIV-1 and HBV and have discontinued products containing emtricitabine and/or tenofovir disoproxil fumarate, and may occur with discontinuation of SYMTUZA™.

  Patients coinfected with HIV-1 and HBV who discontinue SYMTUZA™ should be closely monitored with both clinical and laboratory follow-up for at least several months after stopping treatment. If appropriate, anti-hepatitis B therapy may be warranted, especially in patients with advanced liver disease or cirrhosis, since post-treatment exacerbation of hepatitis may lead to hepatic decompensation and liver failure.

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darunavir/cobicistat/emtricitabine/tenofovir alafenamide tablets
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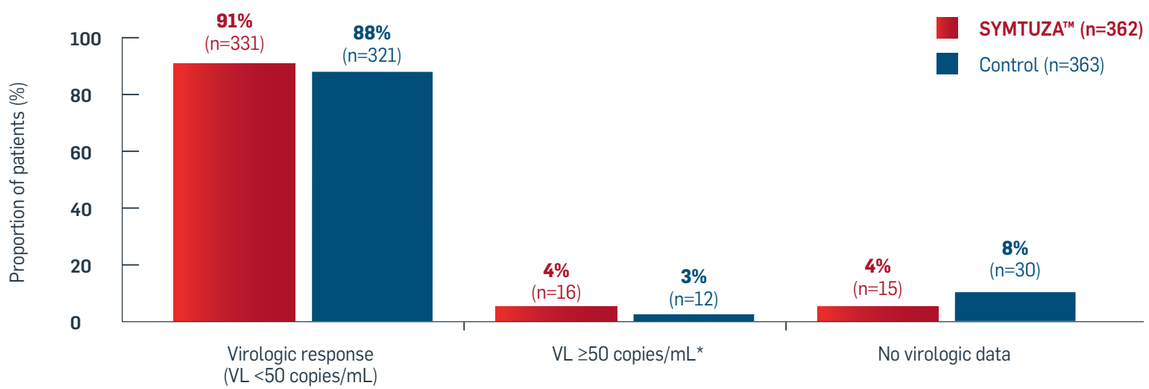
Results: 91% Virologic Response Achieved in Treatment-Naïve Patients

High rates of virologic response in treatment-naïve patients at Week 48 (FDA Snapshot analysis)



- Response rates were similar across subgroups, including age, gender, race, baseline CD4+ count, and baseline VL

0 patients developed treatment-emergent darunavir, primary PI, or TAF mutations

	DRV RAMs	Primary PI RAMs	TAF RAMs
SYMTUZA™ (n=8/362)†	0	0	0
Control (n=6/363)†	0	0	0

- Only one patient receiving SYMTUZA™ was found to have M184I/V; this patient also had a transmitted K103N mutation at screening
  - M184V was detected pretreatment by deep sequencing (Illumina MiSeq) as a minority variant (9.4%)

*Last viral load in the week-48 window at least 50 copies/mL, or discontinuations for efficacy reasons, or premature discontinuations not because of efficacy, adverse events, or death with a last viral load at least 50 copies/mL.
†Of patients with virologic failure, there were 7 in the SYMTUZA™ arm and 2 in the control with paired screening and postbaseline on-treatment genotypes available.
DRV=darunavir, NNRTI=non-nucleoside reverse transcriptase inhibitor; PI=protease inhibitor; RAM=resistance-associated mutations; TAF=tenofovir alafenamide; VL=viral load.

IMPORTANT SAFETY INFORMATION (continued)

- **Hepatotoxicity:** Drug-induced hepatitis and cases of liver injury, including some fatalities, have been reported in patients receiving darunavir, a component of SYMTUZA™. Patients with pre-existing liver dysfunction, including chronic active hepatitis B or C, have an increased risk for liver function abnormalities, including severe hepatic adverse reactions.

Please see additional Important Safety Information throughout and full Prescribing Information, including Boxed WARNING, in pocket.

Tolerability

Most common adverse events at least possibly related to study drug in treatment-naïve patients

(≥5% in either group)	SYMTUZA™ (n=362)	Control (n=363)
Diarrhea	9%	11%
Rash	6%	4%
Nausea	6%	10%

The majority of diarrhea episodes related to SYMTUZA™ were Grade 1 (7%) or Grade 2 (2%)

- Please refer to the full Prescribing Information for a complete list of adverse drug events

Rates of discontinuation due to adverse events

SYMTUZA™ (n=362)	Control (n=363)
2%	4%

- Only 1 patient in each group (0.3%) discontinued the study because of diarrhea
- There were no study drug-related CNS adverse events or discontinuations in >5% of patients

CNS=central nervous system.

IMPORTANT SAFETY INFORMATION (continued)

Appropriate laboratory testing should be conducted prior to initiating and during therapy with SYMTUZA™. Evidence of new or worsening liver dysfunction (including clinically significant elevation of liver enzymes and/or symptoms such as fatigue, anorexia, nausea, jaundice, dark urine, liver tenderness, and hepatomegaly) should prompt consideration of interruption or discontinuation of SYMTUZA™.

- **Severe Skin Reactions:** In patients receiving darunavir, a component of SYMTUZA™, severe skin reactions may occur. Stevens-Johnson syndrome was reported with darunavir coadministered with cobicistat in clinical trials at a rate of 0.1%. During darunavir postmarketing experience, toxic epidermal necrolysis, drug rash with eosinophilia and systemic symptoms (DRESS), and acute generalized exanthematous pustulosis have been reported.

  Discontinue SYMTUZA™ immediately if signs or symptoms of severe skin reactions develop These can include but are not limited to severe rash or rash accompanied with fever, general malaise, fatigue, muscle or joint aches, blisters, oral lesions, conjunctivitis, hepatitis, and/or eosinophilia.

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Important Safety Information (continued)

- **Risk of Serious Adverse Reactions or Loss of Virologic Response Due to Drug Interactions:** The concomitant use of SYMTUZA™ and other drugs may result in known or potentially significant drug interactions, some of which may lead to the loss of therapeutic effect of SYMTUZA™ and possible development of resistance or possible clinically significant adverse reactions from greater exposures of concomitant drugs.

  Consult the full Prescribing Information for potential drug interactions prior to and during SYMTUZA™ therapy, review concomitant medications during SYMTUZA™ therapy, and monitor for the adverse reactions associated with concomitant medications.

- **Immune Reconstitution Syndrome,** including the occurrence of autoimmune disorders with variable time to onset, had been reported in patients treated with combination antiretroviral therapy.

- **New Onset or Worsening Renal Impairment:** Renal impairment, including cases of acute renal failure and Fanconi syndrome, has been reported with the use of tenofovir prodrugs. SYMTUZA™ is not recommended in patients with creatinine clearance below 30 mL per minute. Patients taking tenofovir prodrugs who have impaired renal function and those taking nephrotoxic agents including nonsteroidal anti-inflammatory drugs are at increased risk of developing renal-related adverse reactions.

  In all patients, monitor serum creatinine, creatinine clearance, urine glucose, and urine protein prior to or when initiating SYMTUZA™ and during therapy. In patients with chronic kidney disease, also assess serum phosphorus. Discontinue SYMTUZA™ in patients who develop clinically significant decreases in renal function or evidence of Fanconi syndrome.

- **Sulfa Allergy:** Monitor patients with a known sulfonamide allergy after initiating SYMTUZA™.

- **Lactic Acidosis/Severe Hepatomegaly With Steatosis:** Lactic acidosis and severe hepatomegaly with steatosis, including fatal cases, have been reported with the use of nucleoside analogs, including emtricitabine, a component of SYMTUZA™, and tenofovir disoproxil fumarate (TDF), another prodrug of tenofovir, alone or in combination with other antiretrovirals. Discontinue SYMTUZA™ in any patient who develops clinical or laboratory findings suggestive of lactic acidosis or pronounced hepatotoxicity.

- **Diabetes Mellitus/Hyperglycemia:** New-onset or exacerbations of pre-existing diabetes mellitus and hyperglycemia have been reported in patients receiving protease inhibitors. Initiation or dose adjustments of insulin or oral hypoglycemic agents may be required.

- **Fat Redistribution:** Redistribution and/or accumulation of body fat have been observed in patients receiving antiretroviral therapy.

- **Hemophilia:** Increased bleeding in hemophiliacs has been reported in patients receiving protease inhibitors.

Please see additional Important Safety Information throughout and full Prescribing Information, including Boxed WARNING, in pocket.

IMPORTANT SAFETY INFORMATION (continued)

ADVERSE REACTIONS

- The most common clinical adverse reactions (all grades) occurring in at least 2% of treatment-naïve patients were diarrhea, rash, nausea, fatigue, headache, abdominal discomfort, and flatulence. This is not a complete list of all adverse drug reactions reported with the use of SYMTUZA™. Please refer to the full Prescribing Information for a complete list of adverse drug reactions.

DRUG INTERACTIONS

- Consult the full Prescribing Information for SYMTUZA™ for information on significant drug interactions, including clinical comments.

USE IN SPECIFIC POPULATIONS

- **Pregnancy:** SYMTUZA™ is not recommended for use during pregnancy because of substantially lower exposures of darunavir and cobicistat during pregnancy.

  SYMTUZA™ should not be initiated in pregnant individuals. An alternative regimen is recommended for individuals who become pregnant during therapy with SYMTUZA™.

- **Renal Impairment:** SYMTUZA™ is not recommended in patients with severe renal impairment (creatinine clearance below 30 mL per minute).

- **Hepatic Impairment:** SYMTUZA™ is not recommended for use in patients with severe hepatic impairment (Child-Pugh Class C).

- Consult the full Prescribing Information for SYMTUZA™ for additional information on the Uses in Specific Populations.

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Summary

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In treatment-naïve patients:

- High virologic response
  - 91% of patients using SYMTUZA™ achieved HIV-1 VL <50 copies/mL at Week 48
- 0 patients developed treatment-emergent darunavir, primary PI, or TAF mutations*
- Discontinuations due to adverse events were lower for SYMTUZA™ (2%) vs control (4%)

*Only 1 patient receiving SYMTUZA™ was found to have M184I/V; this patient also had a transmitted K103N mutation at screening. M184V was detected pretreatment by deep sequencing (Illumina MiSeq) as a minority variant (9.4%).

PI=protease inhibitor; TAF=tenofovir alafenamide; VL=viral load.

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A week-48 randomized phase-3 trial of darunavir/cobicistat/emtricitabine/tenofovir alafenamide in treatment-naive HIV-1 patients

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Objectives: To investigate efficacy and safety of a single-tablet regimen of darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) 800/150/200/10 mg vs. darunavir/cobicistat plus emtricitabine/tenofovir disoproxyl fumarate (TDF) (control) in antiretroviral-treatment-naive, HIV-1-infected adults.

Design: Phase-3, randomized, active-controlled, double-blind, international, multicenter, noninferiority study (NCT02431247).

Methods: Seven hundred and twenty-five participants were randomized (1 : 1) to D/C/F/TAF (362) or control (363). The primary objective was to demonstrate noninferiority of D/C/F/TAF vs. control for percentage viral load less than 50 copies/ml (FDA-snapshot analysis) at 48 weeks (10% margin).

Results: At week 48, D/C/F/TAF was noninferior to control (91.4 vs. 88.4% achieved viral load <50 copies/ml, respectively; difference 2.7%; 95% CI −1.6 to 7.1; $P < 0.0001$), with 4.4 vs. 3.3% of patients, respectively, having viral load greater or equal to 50 copies/ml. No treatment-emergent mutations associated with darunavir or TAF/TDF resistance were observed in either group. One patient (D/C/F/TAF) was identified with M184I/V conferring resistance to emtricitabine. Incidences of grades 3 and 4 adverse events (5 vs. 6%), serious adverse events (5 vs. 6%) and adverse event-related discontinuations (2 vs. 4%) were low and similar between groups. Mean decrease in urine protein/creatinine ratio was greater with D/C/F/TAF than control (−22.42 vs. −10.34 mg/g, $P = 0.033$). Mean percentage change in bone mineral density with D/C/F/TAF vs. control was 0.21 vs. −2.73%, $P < 0.0001$ (hip), −0.68 vs. −2.38%, $P = 0.004$ (lumbar spine), and −0.26 vs. −2.97%, $P < 0.0001$ (femoral neck). Median change from baseline in total cholesterol/HDL-cholesterol ratio was 0.20 vs. 0.08, $P = 0.036$.

Conclusion: D/C/F/TAF achieved a high virologic suppression rate (91.4%) and was noninferior to darunavir/cobicistat with F/TDF. D/C/F/TAF also demonstrated the bone and renal safety advantages of TAF in combination with darunavir/cobicistat.

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Introduction

Combination antiretroviral therapy (ART) regimens for HIV-1-infected patients are now more effective, safe and convenient. However, treatment adherence, emergence of resistant virus with virologic failure, and tolerability remain important challenges [1]. Convenient once-daily, single-tablet regimens (STR) can facilitate treatment adherence and improve treatment effectiveness [2,3].

Since its initial approval in 2006, substantial clinical trial data and clinical experience with darunavir have accumulated, demonstrating the potent and durable virologic response, high genetic barrier to resistance, and favorable safety profile in ART-naive, HIV-1-infected patients [4,5]. A substantial proportion of newly diagnosed patients in the United States and Europe are treated with a boosted protease inhibitor [6,7], and darunavir is the recommended protease inhibitor in treatment guidelines [8–10]. United States guidelines recommend two nucleoside or nucleotide analogue reverse transcriptase inhibitors (NRTIs) combined with an integrase strand transfer inhibitor (INSTI), or in certain clinical situations boosted darunavir 800 mg once daily or a nonnucleoside reverse transcriptase inhibitor (NNRTI) [8,9]. Boosted darunavir is recommended for patients with uncertain adherence, those who require a regimen with a high-resistance barrier, or those patients without available resistance results [8]. European guidelines recommend two NRTIs combined with either an INSTI, boosted darunavir or an NNRTI for all ART-naive patients [10], with both darunavir and atazanavir as recommended protease inhibitors in the BHIVA guidelines [11].

Phase-3 studies have established the noninferior antiviral efficacy and improved renal and bone safety of ART regimens containing tenofovir alafenamide (TAF), a newer tenofovir prodrug, vs. tenofovir disoproxil fumarate (TDF), combined with different third agents [12,13], making TAF an optimal backbone component.

Darunavir/cobicistat/emtricitabine/tenofovir alafenamide (D/C/F/TAF) 800/150/200/10 mg is the first and only once-daily protease inhibitor-containing STR in development, combining the antiviral efficacy, and resistance barrier of darunavir with the safety of TAF. D/C/F/TAF was approved for use in Europe in September 2017, and is investigational and currently undergoing regulatory review in the United States. D/C/F/TAF is being evaluated in two international, randomized, phase-3 studies: AMBER (NCT02431247) in ART-naive, HIV-1-infected adults, and EMERALD (NCT02269917) in treatment-experienced adults with virologically suppressed HIV infection [14]. We present the 48-week primary analysis of AMBER, which evaluated D/C/F/TAF vs. darunavir/cobicistat in combination with emtricitabine/TDF (F/TDF).

Methods

Study design

AMBER (TMC114FD2HTX3001; ClinicalTrials.gov NCT02431247; EudraCT 2015–000754–38) is a phase-3, randomized, active-controlled, double-blind, noninferiority study being conducted at 121 sites across 10 countries in North America (USA, Canada) and Europe (Belgium, France, Germany, Italy, Poland, Russia, Spain, UK). The trial included a ~30-day screening period (up to ≤6 weeks) and a 48-week treatment period. In addition, all patients continue to receive D/C/F/TAF in an open-label, single-arm treatment phase up to week 96, and then in a roll-over extension phase.

Participants were randomized (1 : 1) using a computer-generated interactive web-response system to receive D/C/F/TAF 800/150/200/10 mg (q.d.) daily or darunavir/cobicistat 800/150 mg fixed-dose combination (FDC) co-administered with F/TDF 200/300 mg FDC daily (control). Participants received placebo tablets matching the alternative treatment – three tablets in total – and were instructed to take all study drugs and matching placebo tablets with food at approximately the same time each morning. Randomization was stratified by screening viral load (≤ or >100 000 copies/ml) and $CD4^+$ cell count (< or ≥200 cells/μl).

The trial was conducted in accordance with the principles of Good Clinical Practice and Declaration of Helsinki. The protocol and amendments were reviewed and approved by an institutional review board or independent ethics committee. All study participants provided written informed consent.

Study population

Eligible patients were treatment-naive, HIV-1-infected adults (≥18 years) with a screening plasma viral load at least 1000 copies/ml, $CD4^+$ cell count greater than 50 cells/μl, genotypic sensitivity to darunavir, emtricitabine, and tenofovir (GenoSure MG HIV-1 protease/reverse transcriptase genotype assay; Monogram Biosciences, South San Francisco, California, USA), and an estimated glomerular filtration rate based on serum

creatinine ($eGFR_{cr}$) at least 70 ml/min (Cockcroft–Gault formula) [15]. Main exclusion criteria included diagnosis of a new AIDS-defining condition within 30 days prior to screening, hepatitis B or C coinfection, clinically significant disease (e.g. malignancy, severe infections), and pregnancy or breastfeeding in women. Medications or herbal supplements known or suspected to have drug interactions with the investigational medications were disallowed.

Main study assessments and outcomes

Study visits were at baseline, weeks 2, 4, 8, and 12, and then every 12 weeks until week 96. Adverse events were graded according to the Division of AIDS grading table [16] and coded using the Medical Dictionary for Regulatory Activities (version 19.1). At each visit, urine and blood samples were collected for plasma viral load (COBAS AmpliPrep/COBAS TaqMan HIV-1 Test, V2.0; Roche Diagnostics, Basel, Switzerland) and $CD4^+$ cell count determinations, biochemistry, hematology, urinalysis and urine chemistry, serum cystatin C for calculating $eGFR_{cyst}$ [Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) formula] [17], and serum creatinine for calculating $eGFR_{cr}$ (Cockcroft–Gault formula and CKD-EPI formula) [15,17]. Treatment adherence was monitored at each visit (except week 2) by drug accountability (pill count and patient log booklet). The renal proteinuria biomarkers, urinary retinol-binding protein (RBP), and beta-2-microglobulin were measured at baseline, weeks 2, 4, 12, 24, and 48 in the fasted state. Fasted metabolic profile assessments (total, high-density lipoprotein [HDL]-cholesterol and low-density lipoprotein [LDL]-cholesterol, triglycerides) were performed at baseline, weeks 24 and 48. Pharmacokinetic sampling was performed at weeks 2, 4, 8, 12, 24, 36, 48, and study endpoint.

Protocol-defined virologic failure (PDVF) was defined as virologic nonresponse (viral load <1 $\log_{10}$ reduction from baseline and ≥ 50 copies/ml at week 8, confirmed at next visit) or virologic rebound (confirmed viral load ≥ 50 copies/ml after confirmed, consecutive viral load <50 copies/ml or confirmed viral load >1 $\log_{10}$ increase from the nadir) and/or viremia at the final time point (viral load ≥ 400 copies/ml at study endpoint or study discontinuation after week 8). Post screening resistance testing (PhenoSense GT) was performed on samples from patients with PDVF and viral load greater or equal to 400 copies/ml at time of failure (preferably confirmed, or otherwise unconfirmed) or at later time points.

The primary objective was the noninferiority evaluation of D/C/F/TAF vs. darunavir/cobicistat co-administered with F/TDF in the proportion of patients with viral load less than 50 copies/ml (response rate) by the Food and Drug Administration (FDA)-snapshot analysis at week 48.

Secondary outcomes included proportion of patients with viral load <20 and <200 copies/ml (FDA-snapshot analysis) and viral load <50 copies/ml (time-to-loss-of-virologic-response algorithm) at week 48; changes from baseline in $\log_{10}$ viral load and $CD4^+$ cell count; antiretroviral resistance development in PDVFs; safety and tolerability through 48 weeks; changes from baseline at week 48 in serum creatinine, $eGFR_{cr}$, $eGFR_{cyst}$, and ratios of total urine protein, urine albumin, urine RBP, and beta-2-microglobulin to creatinine (UPCR, UACR, RPB:Cr, and B2M:Cr, respectively).

Bone investigation substudy

The bone investigation substudy was performed at selected study sites in consenting participants from both randomization groups. Endpoints at weeks 24 and 48 were percentage changes from baseline in spine, hip, and femoral neck bone mineral density (BMD; measured by dual-energy X-ray absorptiometry scans); changes in associated T score (normal BMD defined as a T score ≥ -1; osteopenia as a T score from ≥ -2.5 to <-1; and osteoporosis as a T score <-2.5); and changes in bone biomarkers, alkaline phosphatase (ALP), procollagen type N-terminal propeptide (P1NP), C-type collagen sequence (CTX), parathyroid hormone (PTH), and 25-hydroxy vitamin (25[OH]D), measured in the fasted state.

Statistical analysis

The week-48 primary analysis was performed on the intent-to-treat population (constituting all patients who were randomized and received at least one dose of study drug). A per-protocol analysis was also performed, excluding patients with major protocol violations or other predefined criteria that potentially affected efficacy. Data analysis was performed using SAS software (SAS Institute, Inc, Cary, North Carolina, USA) version 9.2.

Assuming a response rate of 80% at week 48 (FDA-snapshot analysis) for both treatment groups, 335 patients needed to be enrolled in each group to establish noninferiority of D/C/F/TAF to control, with a noninferiority margin of 10% at 90% power and a one-sided significance level of 2.5%. For the bone investigation substudy, at least 85 patients per treatment group were required to detect an absolute difference between groups in BMD of at least 2% with 90% power, assuming a 4% inter-subject variability and a one-sided significance level of 2.5%.

Noninferiority of D/C/F/TAF to control would be demonstrated if the lower limit of the two-sided 95% confidence interval (CI) of the stratum-adjusted (viral load $\leq 100\,000$ or $>100\,000$ copies/ml and $CD4^+$ cell count <200 or ≥ 200 cells/μl) Mantel–Haenszel difference between treatment groups (D/C/F/TAF minus control) in the week-48 response rate was greater than

−10%. Superiority would be established if the lower limit of the 95% CI was greater than 0.

The difference between groups in least square mean (LSM) change from baseline at week 48 in $CD4^+$ cell count and associated 95% CIs were constructed using analysis of covariance (ANCOVA), including $CD4^+$ cell count at baseline as a continuous covariate. In patients who discontinued, $CD4^+$ cell count values after discontinuation were imputed with the baseline value (noncompleter = failure). For other missing values, the last observation was carried forward.

Baseline and postbaseline HIV-1 genotypes were analyzed for protease resistance-associated mutations (RAMs) [including International Antiviral Society (IAS)–USA primary PI RAMs] and reverse transcriptase RAMs (including IAS-USA NRTI RAMs and IAS–USA NNRTI RAMs), as well as specific RAMs to the study drugs [18]. Antiretroviral sensitivity, based on the genotype/phenotype report, was also assessed.

Within-treatment comparisons of mean changes from baseline in renal and bone biomarkers, and fasting lipids were performed using the Wilcoxon signed-rank test. Between-treatment comparisons were assessed using the Wilcoxon rank-sum test. Between-treatment differences in change from baseline in serum creatinine, eGFR, and BMD were tested using ANCOVA, including treatment as a factor and corresponding baseline values as covariates.

Results

Patient disposition and baseline characteristics

The study began on 6 July 2015, and the cut-off date for the week-48 primary analysis was 13 March 2017. Of 866 screened patients, 725 were randomized and included in the intent-to-treat population (Fig. 1); 362 received D/C/F/TAF and 363 received darunavir/cobicistat with F/TDF.

Through 48 weeks, 93.6% (339/362) of patients in the D/C/F/TAF group and 92.3% (335/363) in the control group completed therapy (Fig. 1). The most common reasons for discontinuing the study, as reported by the investigators, were adverse events, withdrawn consent, and loss to follow-up.

Baseline characteristics were balanced between the two groups (Table 1). Median age was 34 years, 88% were

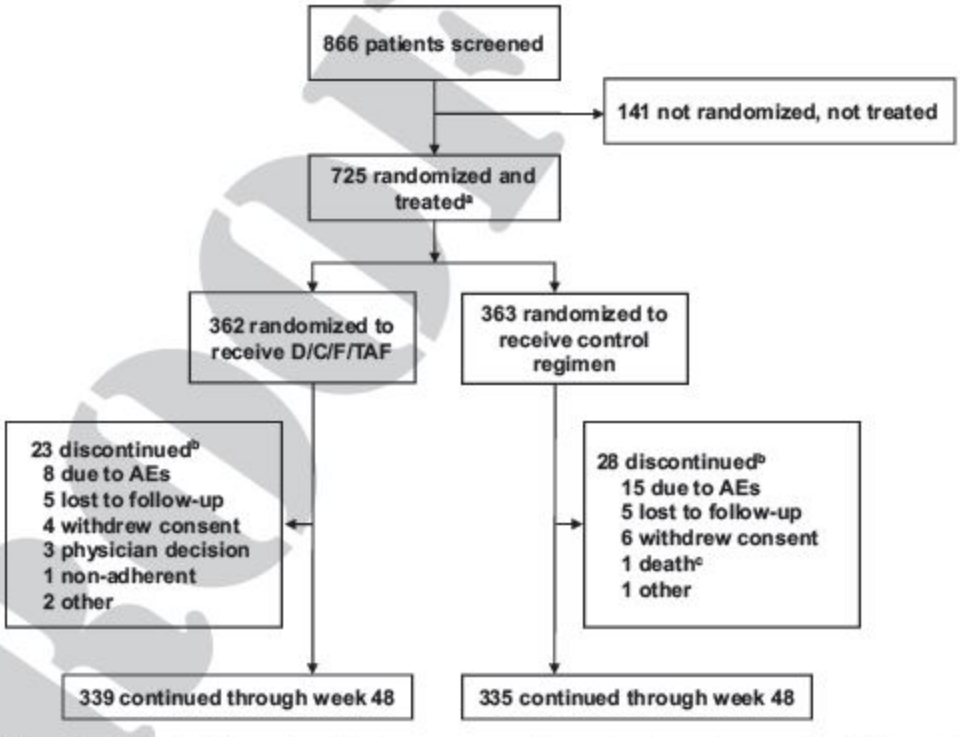

Fig. 1. Patient disposition through 48 weeks. AE, adverse event; Control regimen, darunavir/cobicistat with emtricitabine/tenofovir disoproxil fumarate once daily; D/C/F/TAF, darunavir/cobicistat/emtricitabine/tenofovir alafenamide once daily. [a]Received at least one dose of study medication. [b]Based upon the 'Trial Termination' electronic case report form page as reported by the investigators. Reasons for discontinuation may not match those reported in Supplementary Table 1, http://links.lww.com/QAD/B260 because patients may have viral load data within the week-48 window, which was used to determine the FDA-snapshot category. [c]Occurred in the follow-up phase (11 days after last study drug intake). FDA, Food and Drug Administration.

Table 1. Patient baseline demographics and disease characteristics.

Demographics, *n* (%), unless stated	D/C/F/TAF 800/150/200/10 mg once daily, *N* = 362	Control regimen, *N* = 363	Total, *N* = 725
Median age (IQR), years	34 (27–42)	34 (27–42)	34 (27–42)
More than 50	36 (10)	32 (9)	68 (9)
Gender			
Female	44 (12)	41 (11)	85 (12)
Male	318 (88)	322 (89)	640 (88)
Race			
White	300 (83)	300 (83)	600 (83)
Black/African-American	40 (11)	40 (11)	80 (11)
Other	22 (6)	23 (6)	45 (6)
Ethnicity			
Hispanic or Latino	50 (14)	45 (12)	95 (13)
Baseline disease characteristics			
Median (IQR) time since diagnosis, months	5.73 (2.53–25.59)	4.30 (2.07–17.74)	4.83 (2.33–21.62)
Median (IQR) $\log_{10}$ viral load, copies/ml	4.44 (4.03–4.82)	4.57 (4.15–4.88)	4.52 (4.10–4.87)
Viral load at least 100 000 copies/ml, *n* (%)	60 (17)	70 (19)	130 (18)
Median (IQR) $CD4^+$ cell count, cells/μl	461.5 (342–617)	440.0 (325–594)	453.0 (333–601)
$CD4^+$ cell count less than 200 cells/μl, *n* (%)	22 (6)	29 (8)	51 (7)
Median (IQR) $eGFR_{cr}$, ml/min (Cockcroft–Gault)	119.3 (104.8–135.2) $N = 361$[b]	118.4 (103.2–138.4) $N = 362$[b]	119.1 (104.4–136.5) $N = 723$
Genotype[a] at screening, n (%) [18]			
At least one darunavir resistance-associated mutation	3 (1)	4 (1)	7 (1)[c]
At least one primary protease inhibitor resistance-associated mutation	7 (2)	8 (2)	15 (2)
At least one NRTI resistance-associated mutation	18 (5)	16 (4)	34 (5)[d]
At least one NNRTI resistance-associated mutation	55 (15)	63 (17)	118 (16)[e]

Control regimen, darunavir/cobicistat plus emtricitabine/tenofovir disoproxil fumarate once daily; D/C/F/TAF, darunavir/cobicistat/emtricitabine/tenofovir alafenamide once daily; $eGFR_{cr}$, estimated glomerular rate based on serum creatinine; IQR, interquartile range; NNRTI, nonnucleoside analogue reverse transcriptase inhibitor; NRTI, nucleoside or nucleotide analogue reverse transcriptase inhibitor.
[a]GenoSureMG.
[b]One patient in each group had failed screening genotypes and were enrolled based on local genotypes.
[c]Six V111, one L33F.
[d]The most prevalent NRTI mutation: A62V: 21/725 (2.9%).
[e]The most prevalent NNRTI mutation: K103N 26/725 (3.6%).

men, 83% were white, and 18% had viral load at least 100 000 copies/ml. Median baseline $CD4^+$ cell count was 453 cells/μl.

As depicted by the protocol, at screening, all enrolled participants demonstrated genotypic sensitivity to darunavir, emtricitabine, and tenofovir based on the genotype report. Few had viruses with at least one darunavir RAMs (1%) or primary PI RAMs (2%) (Table 1). No RAMs related to emtricitabine or TDF/TAF were detected. NNRTI and NRTI RAMs were detected in 16 and 5% of patients, respectively (Table 1).

Efficacy

In the primary analysis of virologic response at week 48 (FDA-snapshot analysis), noninferiority of D/C/F/TAF [91.4% (331/362)] vs. control [88.4% (321/363)] was demonstrated (difference 2.7%; 95% CI −1.6 to 7.1; $P < 0.0001$; Fig. 2a and Supplemental Table S1, http://links.lww.com/QAD/B260). A low proportion of participants in the D/C/F/TAF group [4.4% (16/362)] and control group [3.3% (12/363)] had a viral load greater or equal to 50 copies/ml at week 48 (FDA-snapshot analysis).

Results from the per-protocol analysis confirmed noninferiority of D/C/F/TAF [94% (327/348)] to control [92.2% (317/344)] (difference 1.5%; 95% CI −2.3 to 5.2; $P < 0.0001$), as did other sensitivity analyses (Supplementary Table 2, http://links.lww.com/QAD/B260). Week-48 response rates (FDA-snapshot analysis) were consistent across a range of patient subgroups (Fig. 2b).

A similar proportion of patients in each group also achieved a viral load <200 or <20 copies/ml (FDA-snapshot analysis) at week 48 (Supplementary Table 2, http://links.lww.com/QAD/B260). LSM increases ($P < 0.0001$) from baseline in $CD4^+$ cell count (noncompleter = failure) at week 48 were 190.5 cells/μl for D/C/F/TAF vs. 172.0 cells/μl for control ($P = 0.213$ between groups; Supplementary Table 2, http://links.lww.com/QAD/B260).

Virology

Through week 48, eight (D/C/F/TAF) and six (control) participants had PDVF, with paired screening and postbaseline on-treatment genotypes available for seven vs. two patients, respectively. No darunavir, primary protease inhibitor, or TDF/TAF RAMs emerged in any patient. An M184V/I mutation associated with phenotypic resistance to emtricitabine and lamivudine was identified in one patient receiving D/C/F/TAF. This patient harbored a K103N mutation at screening,

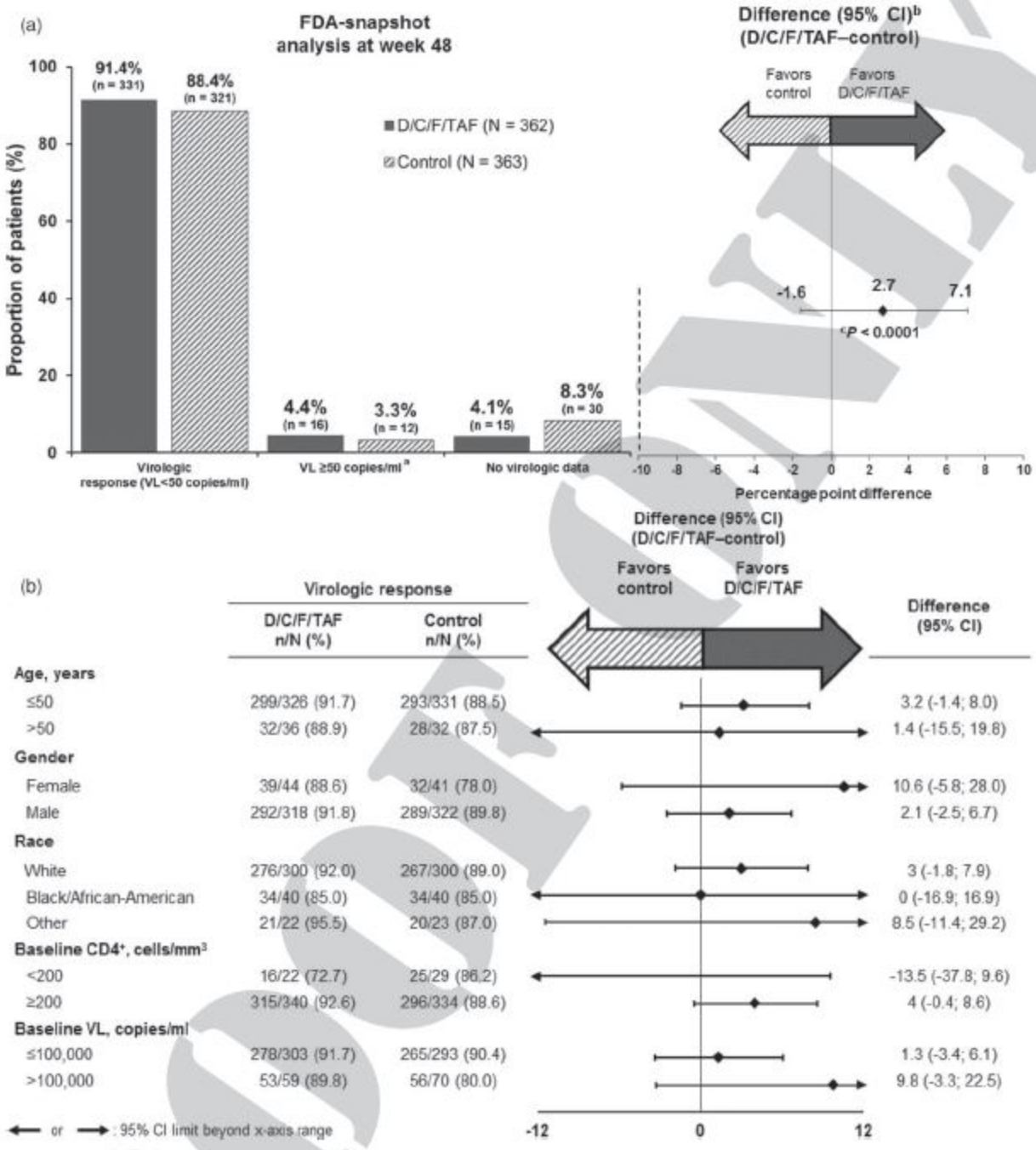

Fig. 2. Week-48 Food and Drug Administration-snapshot analysis (<50 copies/ml). (a) Virologic outcomes overall and (b) subgroup analyses of week-48 response rates. CI, confidence interval; control regimen, darunavir/cobicistat with emtricitabine/tenofovir disoproxil fumarate once daily; D/C/F/TAF, darunavir/cobicistat/emtricitabine/tenofovir alafenamide once daily; VL, viral load. [a]Last viral load in the week-48 window at least 50 copies/ml, or discontinuations for efficacy reasons, or premature discontinuations not because of efficacy, adverse events or death with a last viral load at least 50 copies/ml. [b]Calculated with Mantel–Haenszel test adjusting for screening viral load (≤ or >100 000 copies/ml) and CD4+ cell count (< or ≥200 cells/μl). [c]*P* value for noninferiority at 10% margin.

indicating transmitted NNRTI (efavirenz and nevirapine) resistance. Although the patient appeared to have good adherence (≥95% based on pill count), darunavir plasma concentrations were low [32–192 ng/ml, except at week 4 (1440 ng/ml)], indicating nonadherence that resulted in the patient being discontinued from the study after week 48. All other participants had virus that remained susceptible to all drugs in the treatment regimens.

Table 2. Treatment-emergent adverse events and laboratory abnormalities through 48 weeks.

	D/C/F/TAF 800/150/200/10 mg once daily, $N = 362$	Control regimen, $N = 363$
Any adverse event regardless of causality	312 (86)	307 (85)
Any study drug-related adverse event	126 (35)	151 (42)
Any grade 3 or 4 adverse event regardless of causality	19 (5)	22 (6)
Any serious adverse event regardless of causality[a]	17 (5)	21 (6)
Adverse events leading to permanent discontinuation[b]	7[c] (2)	16 (4)
Death[d]	0	0
Most common adverse events regardless of causality (≥5% of patients in either group)		
Diarrhea[e]	71 (20)	66 (18)
Headache	47 (13)	32 (9)
Nasopharyngitis	40 (11)	31 (9)
Rash	32 (9)	25 (7)
Nausea	28 (8)	45 (12)
Upper respiratory tract infection	20 (6)	21 (6)
Fatigue	19 (5)	18 (5)
Syphilis	17 (5)	19 (5)
Osteopenia	17 (5)	27 (7)
Bronchitis	14 (4)	19 (5)
Adverse events at least possibly related to study drug (≥5% of patients in either group)		
Diarrhea[e]	31 (9)	40 (11)
Rash	22 (6)	14 (4)
Nausea	20 (6)	36 (10)
Median (IQR) change from baseline in fasting lipids at week 48		
Total cholesterol (mg/dl)	28.6 (12.8 to 47.2)[f]	10.4 (−8.0 to 29.8)
HDL-cholesterol (mg/dl)	4.3 (−1.2 to 12.0)[f]	1.5 (−3.9 to 8.1)
LDL-cholesterol (mg/dl)	17.4 (2.9 to 32.9)[f]	5.0 (−10.8 to 19.0)
Triglycerides (mg/dl)	23.9 (−3.0 to 58.5)[g]	14.2 (−12.0 to 40.7)
Total cholesterol/HDL-cholesterol ratio	0.20 (−0.28 to 0.67)[h]	0.08 (−0.41 to 0.53)

Data are *n* (%) unless otherwise stated. Control regimen, darunavir/cobicistat plus emtricitabine/tenofovir disoproxil fumarate once daily; D/C/F/TAF, darunavir/cobicistat/emtricitabine/tenofovir alafenamide once daily; HDL, high-density lipoprotein; IQR, interquartile range; LDL, low-density lipoprotein.

[a]Considered study drug-related in zero (D/C/F/TAF group) vs. six patients (1.7% control; rash and toxic skin eruption in two patients each, and bone marrow edema and Stevens–Johnson syndrome in one patient each).

[b]D/C/F/TAF ($n = 7$): rash ($n = 4$), generalized rash, maculopapular rash, diarrhea ($n = 1$ each); control ($n = 16$): rash/erythema ($n = 7$), toxic skin eruption ($n = 2$), neoplasms ($n = 2$), Stevens–Johnson syndrome, diarrhea, bone marrow edema, increased beta-2-microglobulin, and arthralgia ($n = 1$ each).

[c]One fewer patient in the D/C/F/TAF group (compared with Fig. 1) had an adverse event assessed as leading to discontinuation as data are taken from the adverse event electronic case report form (whether or not the drug was withdrawn), and the patient had interrupted treatment.

[d]One death occurred in the control arm, but in follow-up (not considered related to study drug).

[e]The majority of episodes of diarrhea were mild: grade 1: 16 vs. 13% (related: 7 vs. 9%) and grade 2: 4 vs. 5% (related: 2 vs. 2%).

[f]$P < 0.0001$ (total cholesterol, HDL-cholesterol, LDL-cholesterol).

[g]$P = 0.001$ (triglycerides).

[h]$P = 0.036$ (total cholesterol/HDL-cholesterol ratio) for D/C/F/TAF group vs. control group.

Adherence

Through week 48, 88.3% (264/299) vs. 88.3% (271/307) of patients in the D/C/F/TAF and control groups, respectively, were at least 95% adherent as measured by pill count (all patients took three tablets daily based on the study design).

Safety

Safety profiles were similar between groups (Table 2). Most adverse events regardless of causality were grade 1 or 2. The most common (≥5% in either group) study drug-related adverse events through week 48 were diarrhea, rash, and nausea (Table 2). All episodes of study drug-related diarrhea were mild or moderate (grade 1 or 2) and mostly transient in duration. Only one patient in each group (0.3%) discontinued the study because of diarrhea. There were no nervous system study drug-related adverse events greater than 5% nor discontinuations in either group.

Renal adverse events regardless of causality occurred in 2% (7/362) of D/C/F/TAF vs. 6% (21/363) of control patients. No renal adverse events were suggestive of treatment-emergent proximal renal tubulopathy and no renal adverse events led to discontinuation.

Grades 3 and 4 adverse events regardless of causality, serious adverse events, and adverse event-related discontinuations were rare (Table 2). The only grade 4 adverse event reported for at least two patients was suicide attempt, reported in two (0.6%) patients in the control group. There were no deaths during the treatment phase in either group (Table 2). However, one patient in the control group died following grade 4 sepsis in the follow-up phase (11 days after last study drug intake), which was not considered related to study drug (Fig. 1). Incidences and types of laboratory abnormalities were similar in both treatment groups, being mostly grade 1 or 2.

Median changes from baseline at week 48 for fasting lipid parameters were higher for D/C/F/TAF than control (Table 2 and Supplementary Figure 1, http://links.lww.com/QAD/B260). Changes in HDL-cholesterol favored D/C/F/TAF and remaining lipid increases favored control, with a small, statistically significant difference in the change from baseline in total cholesterol/HDL-cholesterol ratio between groups. Six (1.7%) vs. two (0.6%) patients, respectively, initiated a lipid-lowering drug during the treatment period ($P=0.1770$ between groups).

Serum creatinine increased from baseline to week 48 in the D/C/F/TAF group (4.8 μmol/l), consistent with cobicistat inhibition of creatinine tubular secretion [19], but less so than in the control group (8.2 μmol/l; $P<0.0001$, ANCOVA D/C/F/TAF vs. control). Consequently, the mean decrease in $eGFR_{cr}$ (CKD-EPI formula) at week 48 was less for D/C/F/TAF than control (−5.9 vs. −9.3 ml/min per 1.73 m^2, respectively; $P<0.0001$, ANCOVA; Fig. 3a), although mean $eGFR_{cr}$ was within normal limits. However, mean $eGFR_{cyst}$ (CKD-EPI formula) actually increased at week 48, and the increase was greater for D/C/F/TAF than control (5.3 vs. 2.9 ml/min per 1.73 m^2, respectively; $P=0.001$, ANCOVA) (Fig. 3b).

At week 48, all quantitative measures demonstrated less proteinuria for D/C/F/TAF vs. control, as determined by mean changes from baseline in UPCR [−22.42 mg/g (SD 71.98) vs. −10.34 mg/g (118.18), respectively; $P=0.033$], UACR [−2.45 mg/g (23.81) vs. −0.58 mg/g (68.93); $P=0.003$], RBP:Cr [16.84 μg/g (317.31) vs. 401.12 μg/g (2688.91); $P<0.0001$], and B2M:Cr [−100.58 μg/g (788.60) vs. 837.63 μg/g (6122.87); $P<0.0001$].

Baseline characteristics in the bone investigation substudy were well balanced between the D/C/F/TAF ($N=113$) and control ($N=99$) groups (Supplementary Table 3, http://links.lww.com/QAD/B260). Hip, lumbar spine, and femoral neck BMD from baseline to week 48 were stable with D/C/F/TAF (mean percentage change 0.21, −0.68, and −0.26% at each site, respectively; Fig. 3), whereas they decreased significantly at week 48 in the control group [−2.73, −2.38, and −2.97%, respectively; $P<0.0001$ (hip and femoral neck) and $P=0.004$ (spine) for between-treatment comparisons]. Fewer patients receiving D/C/F/TAF had at least 3% decreases from baseline in BMD at each site than in the control group. More patients had at least 3% increases in the D/C/F/TAF group (Supplementary Table 4, http://links.lww.com/QAD/B260). A similar trend was seen for at least 5 and at least 7% increases or decreases in BMD (Supplementary Table 4, http://links.lww.com/QAD/B260). At week 48, a greater proportion of participants receiving D/C/F/TAF had improvements in T score at each site than in the control group, and a smaller proportion of participants receiving D/C/F/TAF had worsening BMD status (Supplementary Table 4, http://links.lww.com/QAD/B260). Fractures occurred infrequently and were not different between groups [1.1% (4/362) D/C/F/TAF vs. 0.6% (2/363) control; $P=0.451$]; all were traumatic and none were suspected to be osteoporotic. New antiosteoporotic treatment was started by 9/362 (2.5%) vs. 16/363 (4.4%) patients, respectively, during the treatment phase. Changes from baseline in bone biomarker levels (ALP, P1NP, CTX, and PTH) suggested less bone turnover for D/C/F/TAF than control (Supplementary Figure 2, http://links.lww.com/QAD/B260). 25[OH]D levels increased from baseline in both groups.

Discussion

In this investigational phase-3, double-blinded, randomized, controlled trial, the D/C/F/TAF once-daily STR was virologically noninferior to darunavir/cobicistat co-administered with F/TDF in ART-naive patients. Response rates were similar across age, sex, race, and baseline HIV characteristics including $CD4^+$ cell count less than 200 cells/μl and viral load greater than 100 000 copies/ml. Although INSTI-based regimens have rapidly moved up in global treatment guidelines [8–10], there are still many patients who might benefit from the established characteristics of the protease inhibitor darunavir, such as high genetic barrier to resistance, efficacy in the face of resistance and uncertain adherence, provider comfort, and experience. Well powered, phase-3, double-blinded, randomized studies provide the most rigorous evidence to drive treatment guidelines. The week-48 virologic response rate (FDA-snapshot analysis) of 91.4% for D/C/F/TAF was among the highest achieved by a STR in phase-3 trials (range 80–93%) of ART-naive patients [12,20–26], and higher than in prior phase-3 trials with darunavir [4,23,27,28].

No treatment-emergent mutations associated with darunavir or tenofovir resistance were observed. Only one patient (D/C/F/TAF) was found to have M184I/V, conferring resistance to emtricitabine; this patient also had a transmitted K103N mutation at screening. M184V was detected pretreatment by deep sequencing (Illumina MiSeq) as a minority variant (9.4%). In addition, for this patient, darunavir plasma concentrations were low and much lower than the steady-state predose concentration (~692 ng/ml), indicating potential nonadherence, which in fact resulted in discontinuation from the study. The observation of no darunavir phenotypic resistance and the genotypic results are consistent with previous darunavir studies [4,5,27,29], confirming the high resistance barrier of darunavir-based initial ART with no emergence of DRV resistance. D/C/F/TAF is the only STR in development that combines the high barrier to resistance of darunavir with the F/TAF backbone. In this context, D/C/F/TAF may have an important role for treating

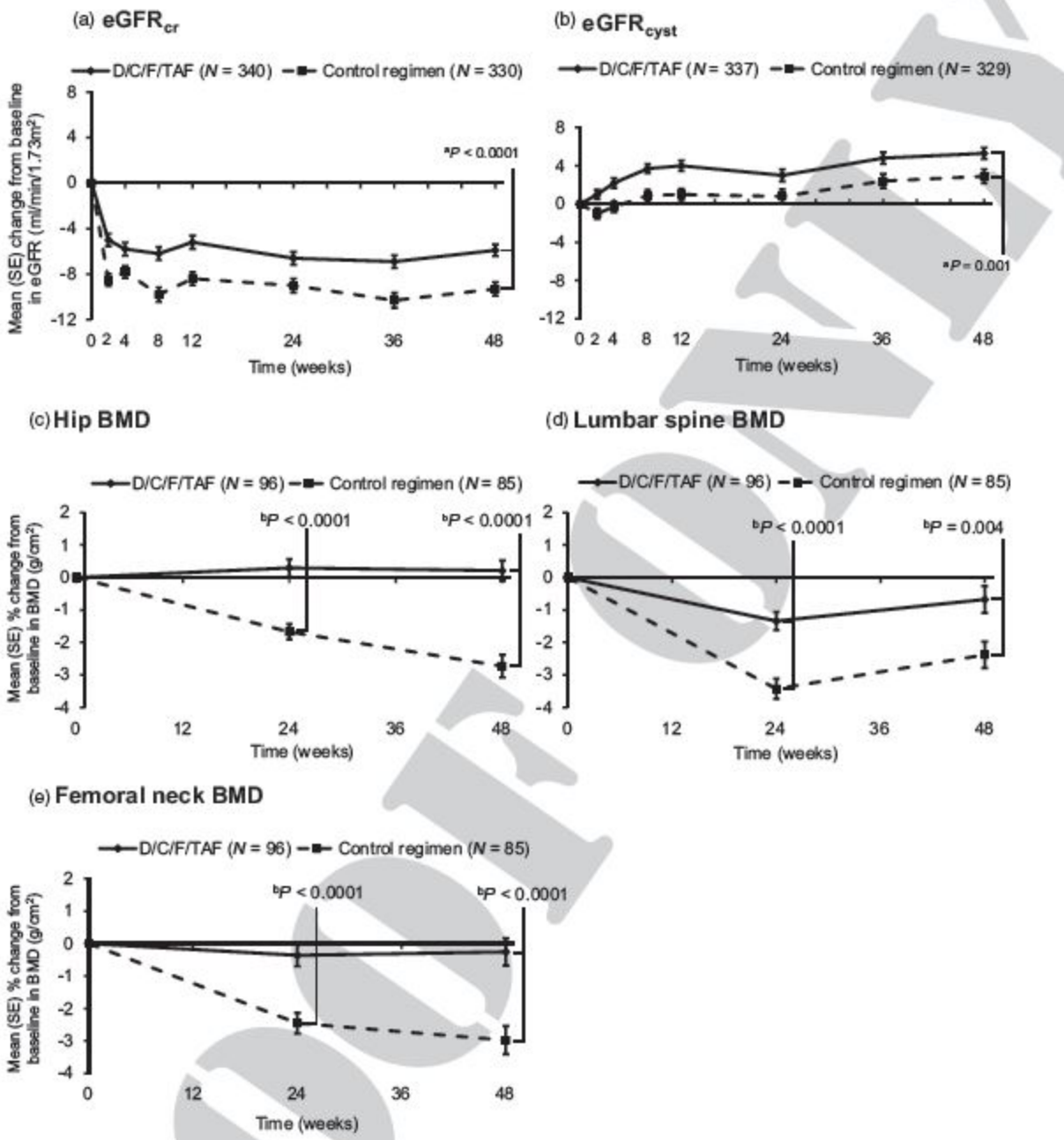

Fig. 3. Mean change from baseline to week 48 in kidney and bone parameters. Mean change in (a) eGFR$_{cr}$ and (b) eGFR$_{cyst}$ was based on serum concentrations and the Kidney Disease Epidemiology Collaboration formula. BMD of the (c) hip, (d) lumbar spine, and (e) femoral neck was analyzed with dual energy X-ray absorptiometry. Bars show SE. ANCOVA, analysis of covariance; BMD, bone mineral density; Control regimen, darunavir/cobicistat with emtricitabine/tenofovir disoproxil fumarate once daily; D/C/F/TAF, darunavir/cobicistat/emtricitabine/tenofovir alafenamide once daily; eGFR$_{cr}$, estimated glomerular filtration rate based on serum creatinine; eGFR$_{cyst}$, estimated glomerular filtration rate based on serum cystatin C; SE, standard error. [a]*P*-value for between-treatment comparison estimated using ANCOVA, including treatment as a factor and corresponding baseline eGFR as a covariate; within-group changes from baseline at week 48, $P < 0.0001$ (both groups for eGFR$_{cr}$ and eGFR$_{cyst}$). [b]*P* value for between-treatment comparison estimated using ANCOVA, including treatment as a factor and baseline BMD as a covariate; Within-group changes from baseline at week 48 at each site, $P < 0.0001$ (D/C/F/TAF); *P* = nonsignificant (control).

patients with uncertain adherence or who plan to start treatment prior to the availability of resistance-testing results [8]. Patients with transmitted NNRTI and NRTI resistance were included in the study. As D/C/F/TAF does not require HLA B*5701 screening or hepatitis or resistance testing before treatment initiation, it is currently being evaluated in a rapid initiation protocol (NCT03227861). These characteristics suggest

D/C/F/TAF is a highly feasible option in a test and treat setting or for very early treatment-naive patients where rapid combination ART initiation could be warranted.

Safety profiles were similar between the two treatment groups. However, adverse event-related discontinuations were lower for D/C/F/TAF (2%) than control (4%), and similar to those reported in phase-3 studies of other recently approved STRs [12,20–26]. The low incidences and similar types of adverse events, grade 3 or 4 adverse events, and serious adverse events between groups reflects the well characterized safety profiles for darunavir and cobicistat reported previously [4,27,29]. Given the low incidence of nervous system adverse events, D/C/F/TAF may be an important treatment option for ART-naive patients at risk of nervous system adverse events, such as insomnia and depression.

Less renal tubular proteinuria, and more favorable hip and spine BMD for D/C/F/TAF compared with control are consistent with TAF vs. TDF effects [12,13,29–31]. The improvement in $eGFR_{cyst}$ could reflect ART-related improvement in HIV-associated renal impairment, as was seen in the START study [32]. The favorable renal tubular and BMD outcomes at the 48-week time point are reassuring, given the fact that the cumulative adverse effects of TDF on renal and bone outcomes have been greater whenever TDF was combined with boosted protease inhibitors [33]. Median increases from baseline in fasting lipids were higher for D/C/F/TAF vs. control, with the increase in HDL-cholesterol favoring D/C/F/TAF and remaining lipid increases favoring control. There was a small, statistically significant difference in the total cholesterol/HDL-cholesterol ratio between groups. Differences in lipid profiles were likely because of the loss of the lipid-lowering effect of TDF rather than an adverse effect of TAF or any other of the components on lipids [12,13].

As in other recent phase-3 trials in ART-naive patients [20–26], study limitations were inclusion of more than 80% white patients and a comparatively small proportion of female or older (>50 years) participants or who had high viral loads. The latter most likely reflects earlier initiation of ART based on current guideline recommendations [8–11]. Phase-3 studies often lack power to detect rare clinical safety events; however, the large clinical safety database for darunavir and substantial clinical experience counterbalance this limitation. Renal and bone safety were assessed using surrogate markers rather than clinical events, and bone safety was assessed in a smaller number of patients.

In conclusion, D/C/F/TAF was noninferior to a regimen of darunavir/cobicistat co-administered with F/TDF at week 48, with a high virologic response (91.4%) in ART-naive, HIV-1-infected adults. D/C/F/TAF was associated with a better bone and renal safety profile than control, with few moderate, severe, or serious adverse events. Changes in HDL-cholesterol favored D/C/F/TAF and remaining lipid increases favored control. D/C/F/TAF is a novel STR that combines the known efficacy and high-genetic barrier to resistance of darunavir with the safety advantages of TAF to provide a new option for the treatment of ART-naive, HIV-1-infected patients.

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