Maraviroc versus Efavirenz in Treatment-Naïve MERIT (A4001026) Trial



Maraviroc versus Efavirenz, both with Zidovudine-Lamivudine MERIT (A4001026): Study Design

Study Design: MERIT Study

- Background: Randomized, double-blind, doubledummy, phase 2b/3 study evaluating the efficacy and safety of maraviroc versus efavirenz as part of ART for treatment-naïve persons with HIV infection
- Inclusion Criteria (n = 721 treated/analyzed)
 - Age ≥16
 - Antiretroviral-naïve patients
 - R5-tropic virus
 - HIV RNA ≥2000 copies/mL
 - No resistance to zidovudine, lamivudine, or efavirenz
- Treatment Arms
 - Maraviroc 300 mg BID + ZVD-3TC BID
 - Efavirenz 600 mg QD+ ZVD-3TC BID

MVC 300 mg twice daily + ZVD-3TC twice daily (n = 360)

EFV once daily + ZVD-3TC twice daily

(n = 361)

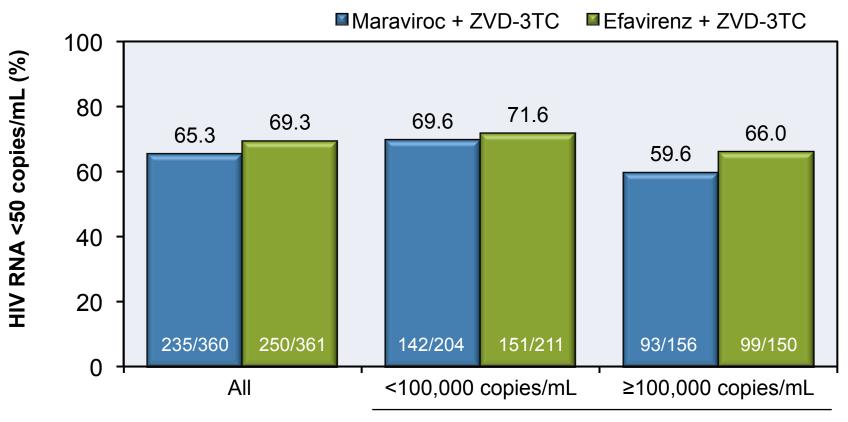
MVC 300 mg once daily + ZVD-3TC twice daily arm (n = 174)
DISCONTINUED at interim analysis

MERIT = Maraviroc versus Efavirenz in Treatment-Naive



Maraviroc versus Efavirenz, both with Zidovudine-Lamivudine MERIT (A4001026): Result

Week 48: Virologic Response (Primary Analysis)

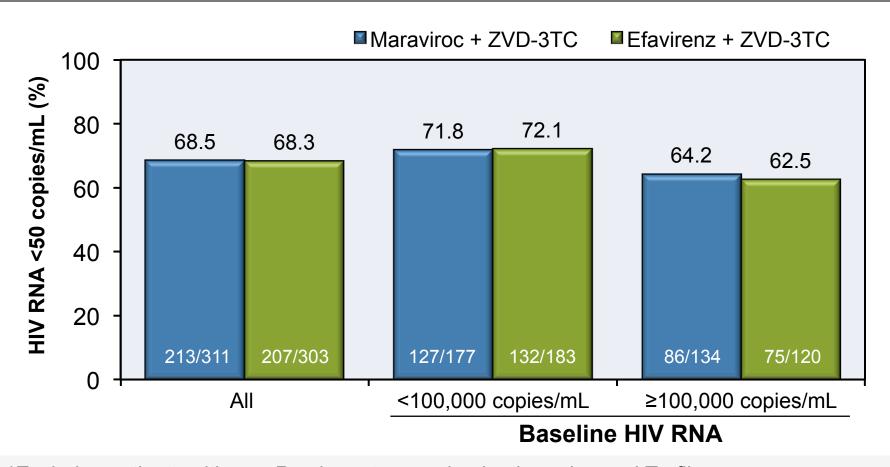


Baseline HIV RNA



Maraviroc or Efavirenz, both with Zidovudine-Lamivudine MERIT (A4001026): Result

Week 48: Virologic Response (Post-hoc Reanalysis*)



^{*}Excludes patients with non-R5 virus at screening by the enhanced Trofile assay



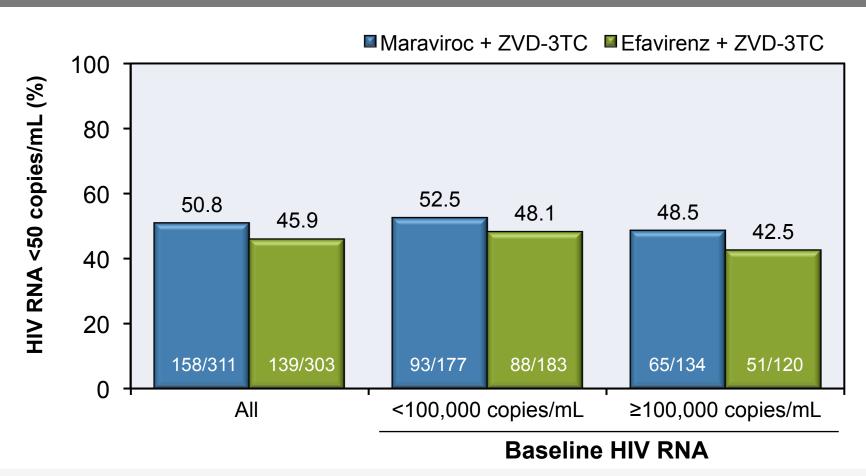
Maraviroc versus Efavirenz, both with Zidovudine-Lamivudine MERIT (A4001026): Conclusions

Conclusions: "Twice-daily maraviroc was not noninferior to efavirenz at <50 copies/mL in the primary analysis. However, 15% of patients would have been ineligible for inclusion by a more sensitive screening assay. Their retrospective exclusion resulted in similar response rates in both arms."



Maraviroc versus Efavirenz, plus Zidovudine-Lamivudine MERIT (A4001026): Result

Week 240 (Year 5): Virologic Response



*Excludes patients with non-R5 virus at screening by the enhanced Trofile assay

Source: Cooper DA, et al. AIDS. 2014;28:717-25.



Acknowledgment

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