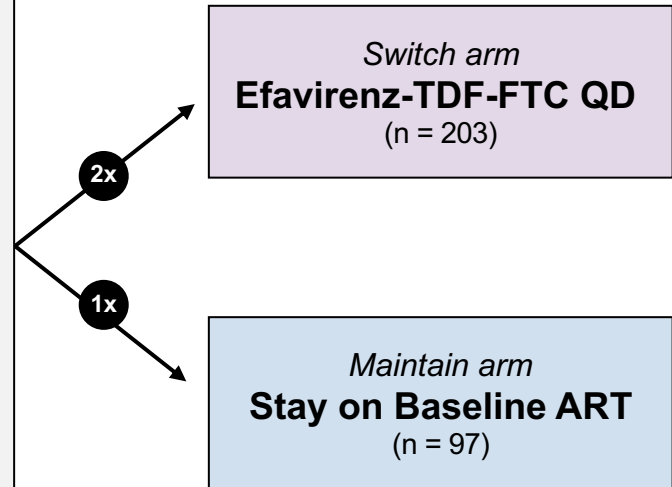


Simplification to Efavirenz-TDF-FTC
STUDY 073

Simplification to Efavirenz-TDF-FTC

Study 073: Study Design

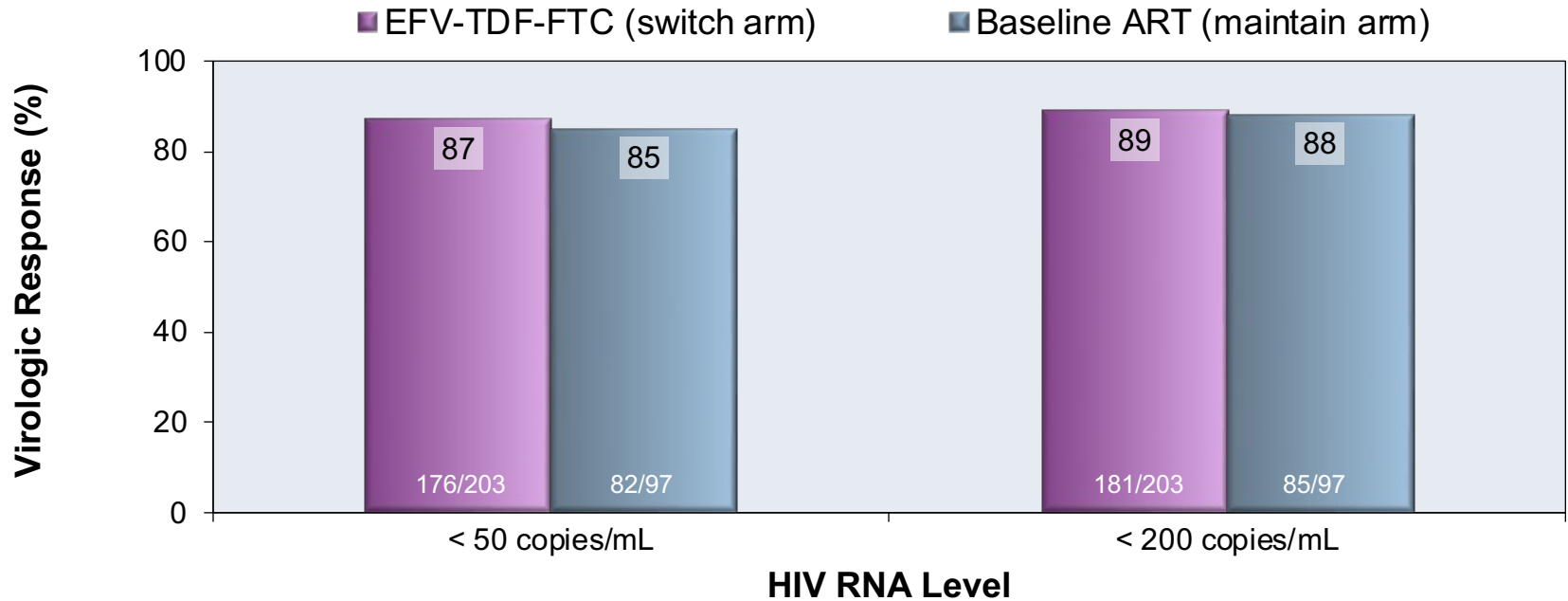
- **Background:** Randomized, controlled, open-label, phase 3 trial evaluating a simplification strategy for patients suppressed on baseline ART by switching to a single-tablet regimen of efavirenz-tenofovir DF-emtricitabine once daily
- **Inclusion Criteria (n = 300)**
 - Age ≥ 18 years
 - HIV RNA ≤ 200 copies/mL for ≥ 3 months
 - No new AIDS-defining conditions in past 30 days
 - On 1st ART regimen or viral suppression on previous protease inhibitor-based therapy at time of prior therapy switch
- **Treatment Arms**
 - Switch arm: Efavirenz-tenofovir DF-emtricitabine
 - Stay arm: maintain baseline ART



Simplification to Efavirenz-TDF-FTC

Study 073: Results

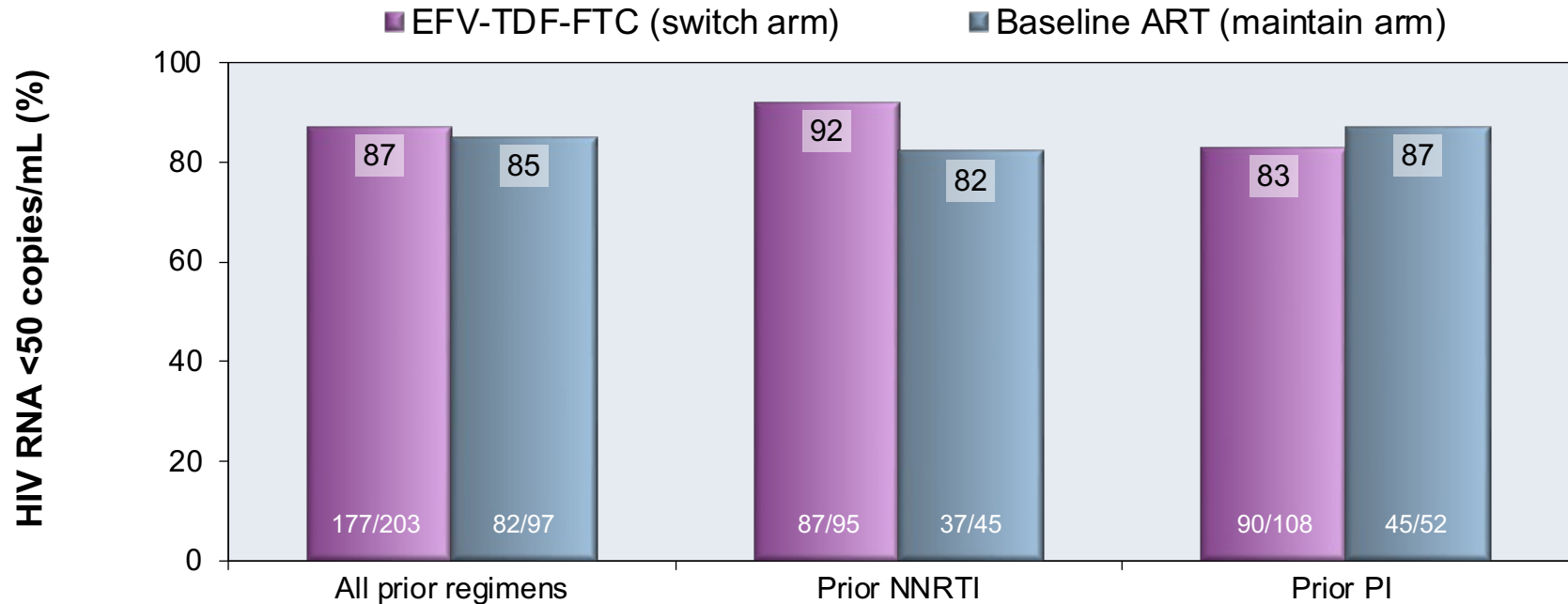
Week 48 Virologic Response by RNA Threshold



Simplification to Efavirenz-TDF-FTC

Study 073: Subgroup Analysis Result

Week 48 Virologic Response, by Baseline Regimen (Intent-to-Treat, Noncompletion = Failure)



Simplification to Efavirenz-TDF-FTC

Study 073: Conclusions

Interpretation: “Simplification to EFV/FTC/TDF maintained high and comparable rates of virologic suppression versus stay on baseline regimen (SBR) through 48 weeks.”

Acknowledgments

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