Switch to Atazanavir-RTV with Either Raltegravir or TDF-FTC HARNESS Trial



Switch to Atazanavir-ritonavir with Either Raltegravir or TDF-FTC HARNESS: Study Design

Study Design: HARNESS

- Background: Open label, prospective, randomized, parallel group trial evaluating switching from stable ARV regimen (2 NRTIs + 3rd agent, excluding atazanavir) to ritonavirboosted atazanavir with either raltegravir 400 mg BID or TDF-FTC.
- Inclusion Criteria (n = 109)
 - Age ≥18 years
 - HIV RNA <50 copies/mL for ≥3 months
 - Single HIV RNA <40 copies/mL in past 30 days
 - No history of virologic failure or resistance
- Treatment Arms
 - Atazanavir + RTV QD + Raltegravir 400 mg BID
 - Atazanavir + RTV QD + TDF-FTC QD

Atazanavir QD + Ritonavir QD + Raltegravir 400 mg BID (n = 72)

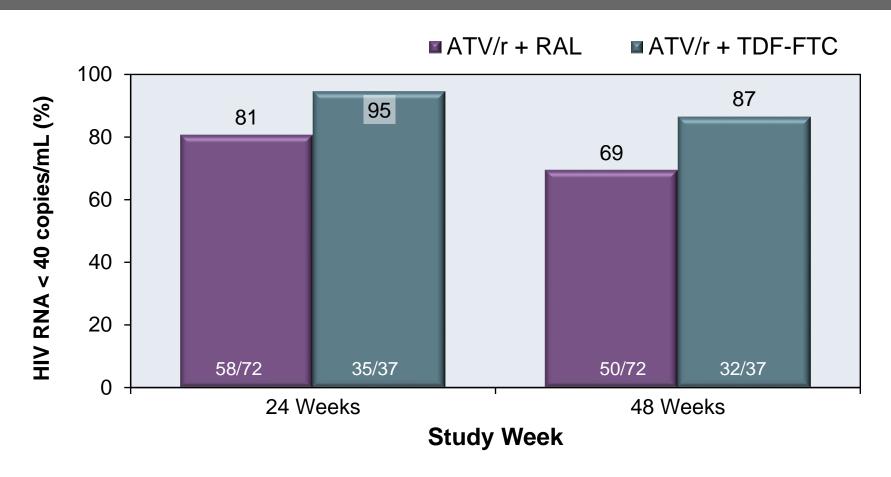
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Atazanavir QD + Ritonavir QD + TDF-FTC QD (n = 37)



Switch to Atazanavir-ritonavir with Raltegravir or TDF-FTC HARNESS: Results

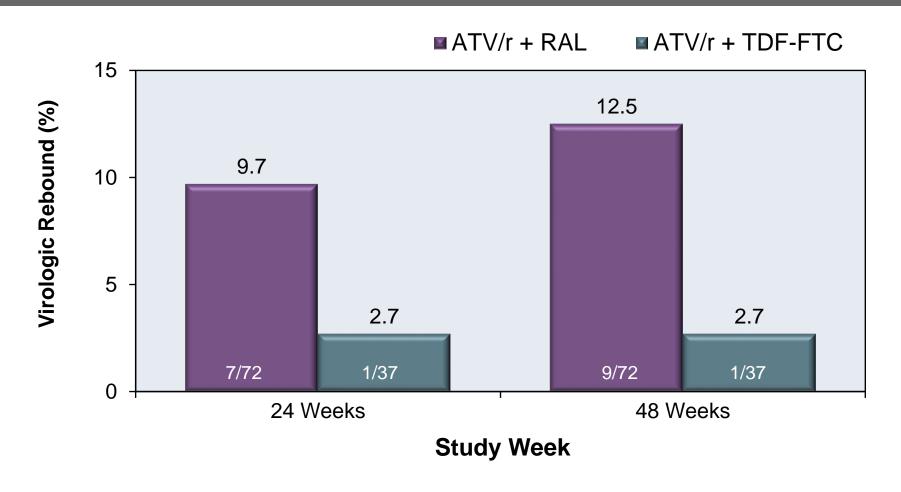
Week 24 and 48: Virologic Response (Intent-to-Treat Analysis)





Switch to Atazanavir-ritonavir with Raltegravir or TDF-FTC HARNESS: Results

Week 24 and 48: Virologic Rebound





Switch to Atazanavir-ritonavir with Raltegravir or TDF-FTC HARNESS: Conclusions

Conclusion: "In conclusion, switching to ATV/r + RAL resulted in a higher virological rebound rate than switching to ATV/r plus tenofovir disoproxil fumarate/emtricitabine."



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