

Atazanavir + Ritonavir + ABC-3TC
SHARE Trial

Atazanavir + Ritonavir + ABC-3TC in Treatment-Naïve SHARE: Study Design

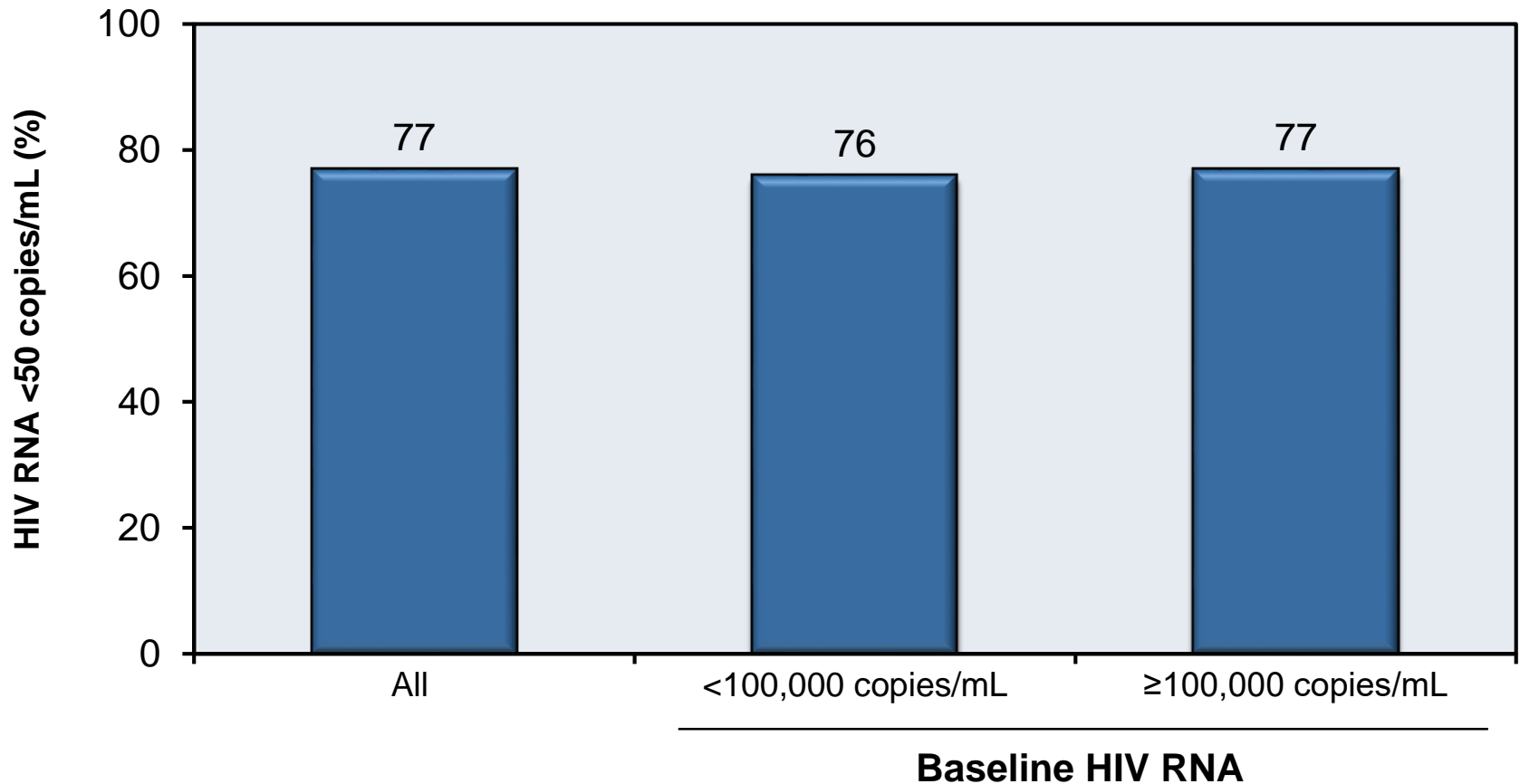
Study Design: SHARE

- **Background:** Nonrandomized, open-label, single-arm pilot study evaluating the efficacy and safety of once-daily abacavir-lamivudine plus ritonavir-boosted atazanavir in treatment-naïve adults with HIV infection
- **Inclusion Criteria (n = 112)**
 - Age ≥ 18
 - Antiretroviral treatment-naïve
 - HIV RNA ≥ 5000 copies/mL
 - No AIDS-defining illnesses in previous 30 days
- **Treatment Arms** (All Once daily)
 - Atazanavir 300 mg + Ritonavir 100 mg + Abacavir-Lamivudine 600/300 mg

Atazanavir + ritonavir + ABC-3TC
(n = 111)

Atazanavir + Ritonavir + ABC-3TC in Treatment-Naïve SHARE: Results

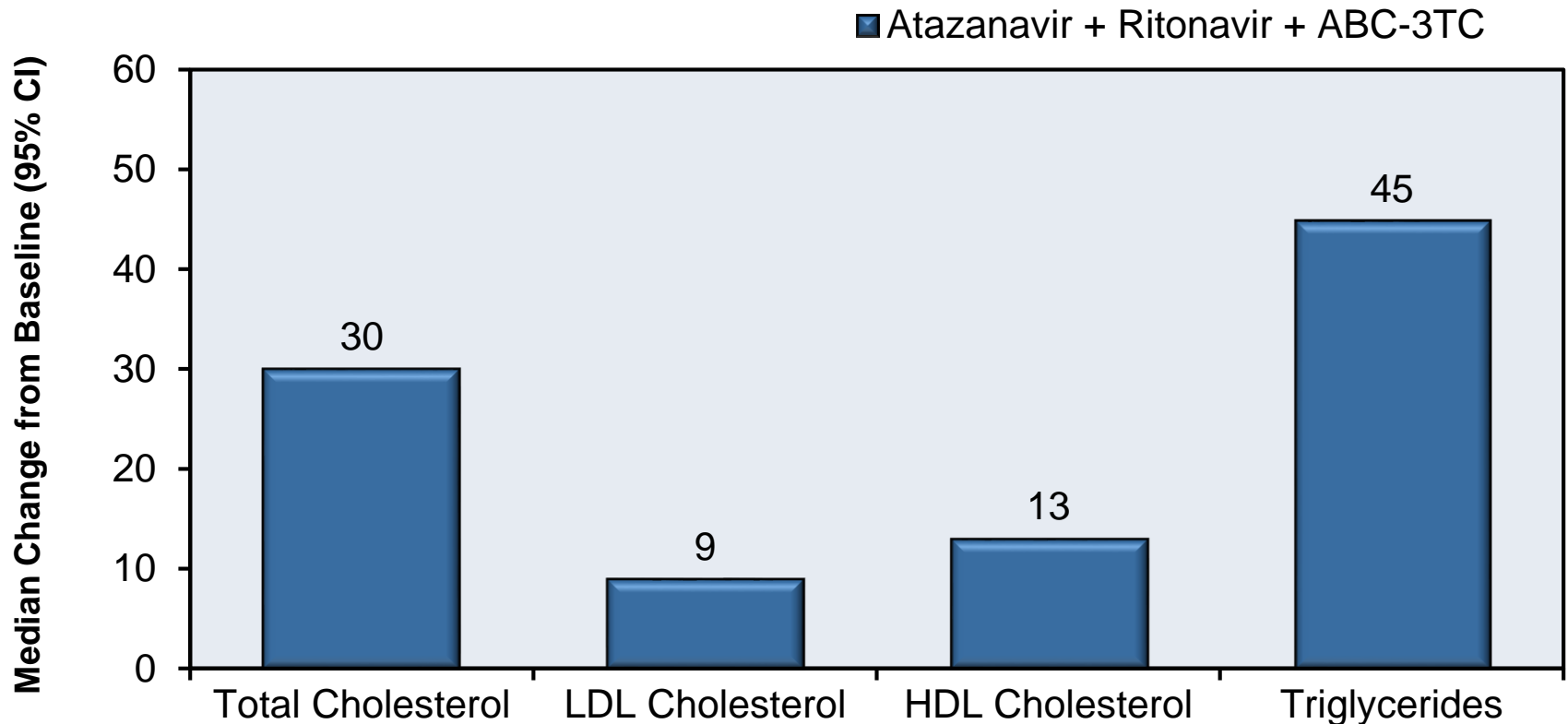
Week 48: Virologic Response, by Baseline HIV RNA (ITT, M=F)



Source: Elion R, et al. HIV Clin Trials. 2008;9:152-63.

Atazanavir + Ritonavir + ABC-3TC in Treatment-Naïve SHARE: Results

Week 48: Change in Fasting Lipids from Baseline



Source: Elion R, et al. HIV Clin Trials. 2008;9:152-63.

Atazanavir/r + ABC-3TC in Treatment-Naïve SHARE: Conclusions

Conclusion: “ABC/3TC and ATV-RTV QD is an effective and well-tolerated regimen in ART-naïve patients through 48 weeks, with a modest impact on fasting lipids.”

Acknowledgment

The **National HIV Curriculum** is an AIDS Education and Training Center (AETC) Program resource funded by the United States Health Resources and Services Administration. The project is led by the University of Washington and the AETC National Coordinating Resource Center.

The content in this slide set does not represent the official views of the U.S. Department of Health and Human Services, Health Resources & Services Administration.

