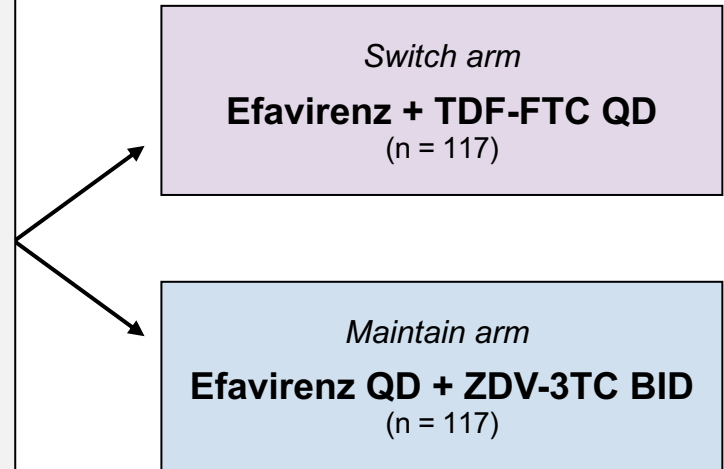


Switch from Efavirenz + ZDV-3TC to Efavirenz + TDF-FTC
SWEET Trial

Switch to Efavirenz + TDF-FTC

SWEET: Study Design

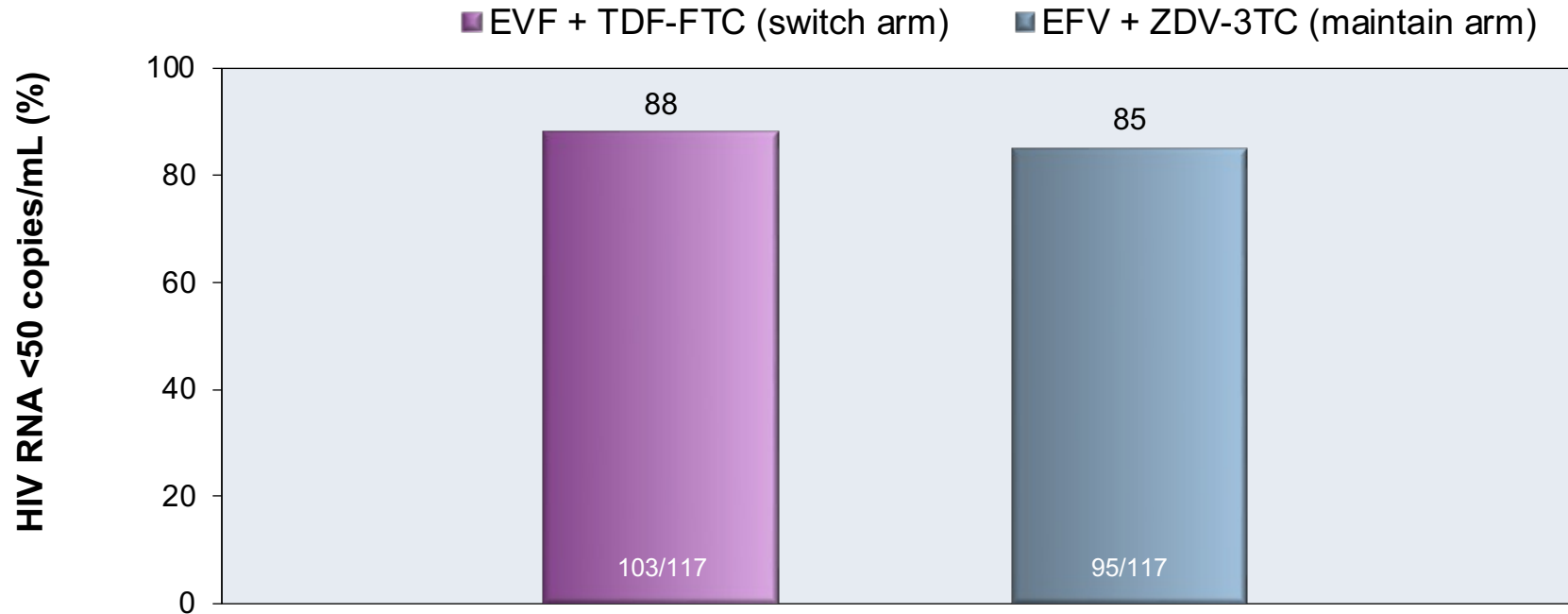
- **Background:** Randomized, controlled, open-label, phase 3 trial evaluating a simplification strategy for patients suppressed on efavirenz-based ART by switching from twice-daily zidovudine-lamivudine to once-daily tenofovir DF-emtricitabine in adults with HIV
- **Inclusion Criteria (n = 234)**
 - Age ≥ 18 years
 - On EFV + ZDV-3TC for >6 months
 - No resistance to study drugs
 - HIV RNA <400 copies/mL for ≥ 3 months and HIV RNA <50 copies/mL on 2 occasions
- **Treatment Arms**
 - Efavirenz + TDF-FTC
 - Efavirenz + ZDV-3TC



Switch to Efavirenz + TDF-FTC

SWEET: Results

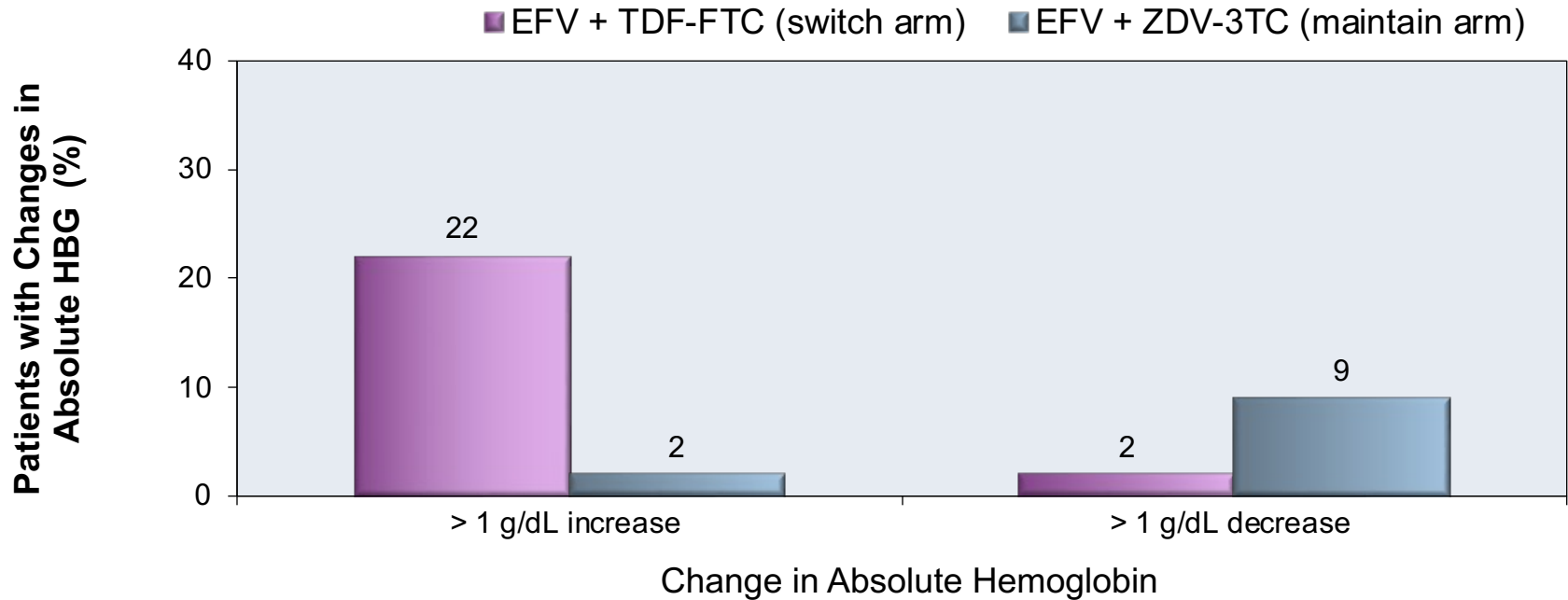
Week 48 Virologic Response (Intention-to-Treat Analysis, Missing=Failure)



Switch to Efavirenz + TDF-FTC

SWEET: Results

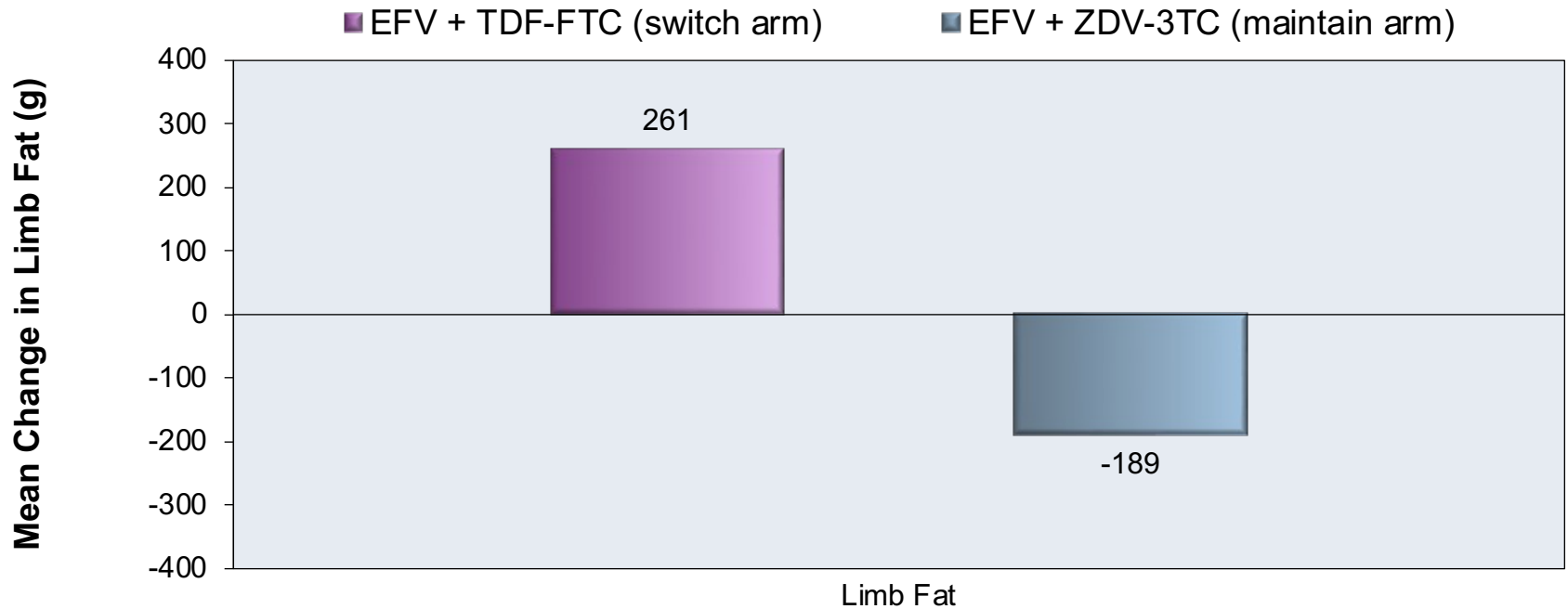
Week 48: Patients with Change in Absolute Hemoglobin from Baseline



Switch to Efavirenz + TDF-FTC

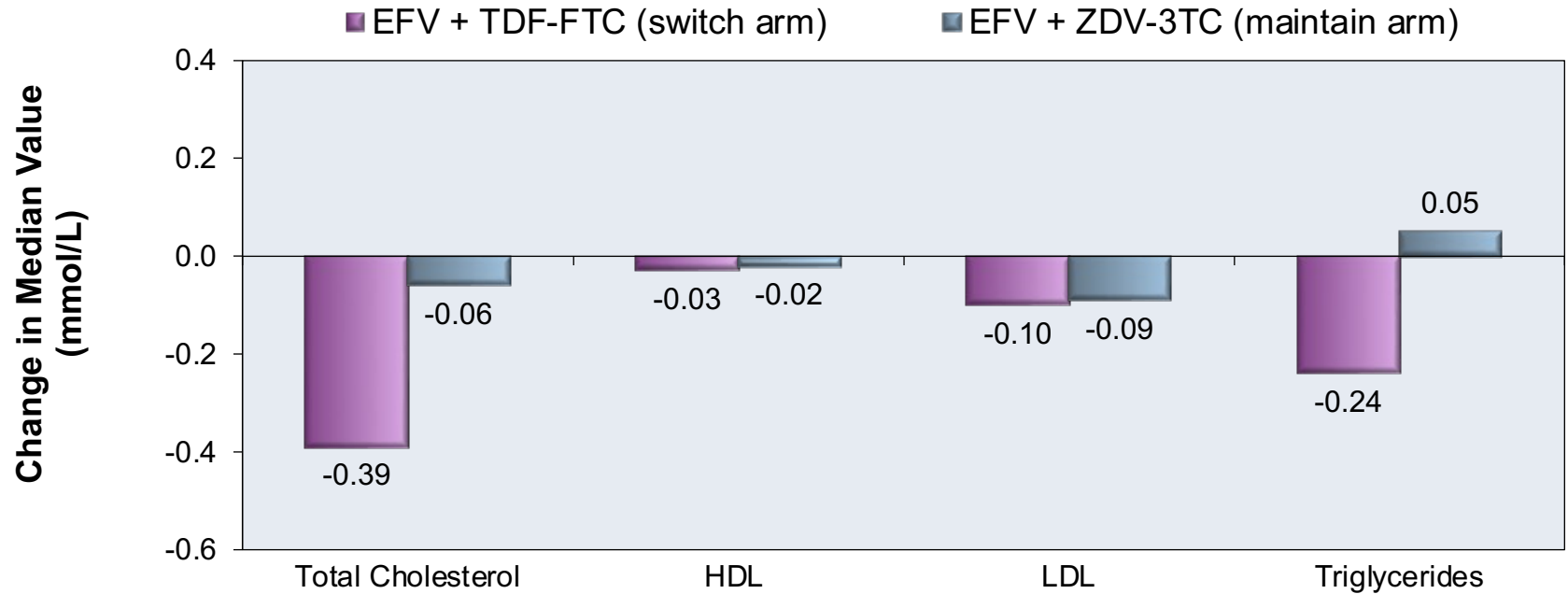
SWEET: Results

Week 24: Change in Limb Fat from Baseline



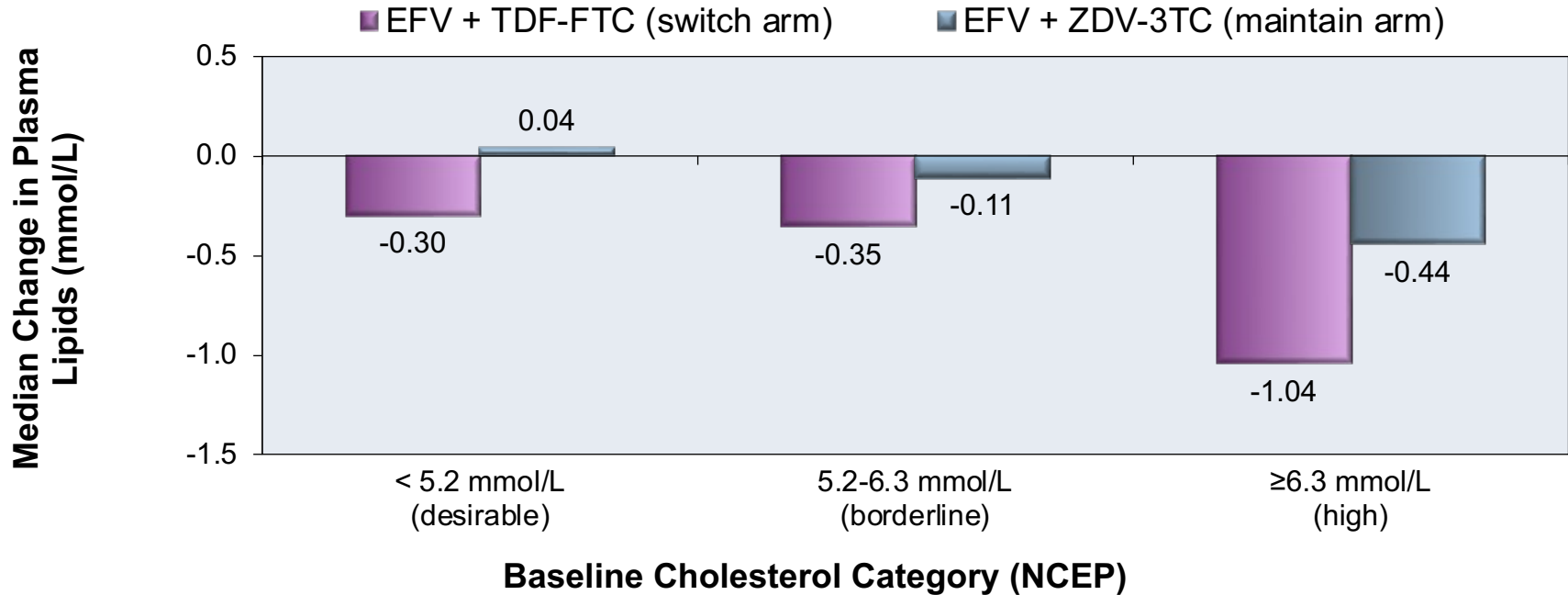
Switch to Efavirenz + TDF-FTC SWEET: Results

Week 24: Change in Plasma Lipids from Baseline



Switch to Efavirenz + TDF-FTC SWEET: Results

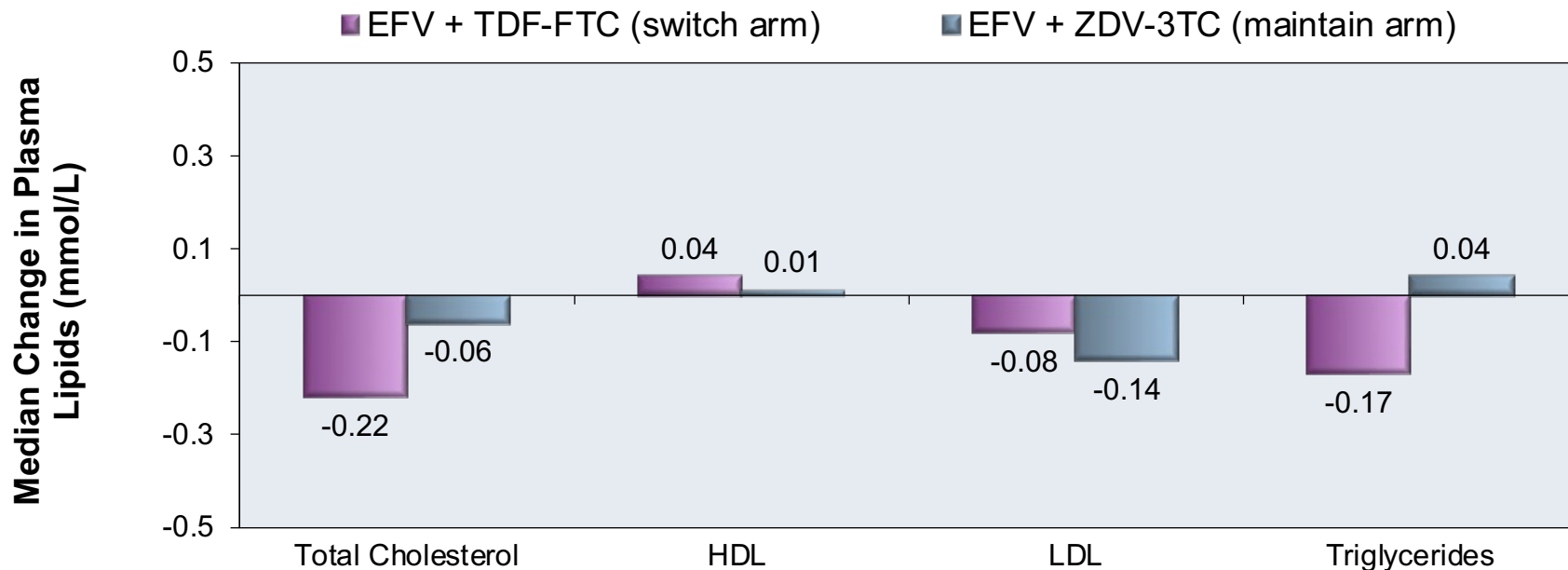
Week 24: Change in Plasma Lipids, by Baseline Cholesterol



Switch to Efavirenz + TDF-FTC

SWEET: Results

Week 48: Change in Plasma Lipids from Baseline



Switch to Efavirenz + TDF-FTC

SWEET: Results

Interpretation: “Switching from zidovudine/lamivudine to tenofovir disoproxil fumarate/emtricitabine in persons on efavirenz therapy maintains virological control, establishes a once-daily regimen, results in improvements in hemoglobin and key lipid parameters, and preserves and restores limb fat relative to continuation of zidovudine/lamivudine.”

Acknowledgments

The **National HIV Curriculum** is supported by the Health Resources and Services Administration (HRSA) of the U.S. Department of Health and Human Services (HHS) as part of a financial assistance award totaling \$1,021,448 with 0% financed with non-governmental sources. The contents are those of the author(s) and do not necessarily represent the official views of, nor an endorsement, by HRSA, HHS, or the U.S. Government. For more information, please visit HRSA.gov. This project is led by the University of Washington's Infectious Diseases Education and Assessment (IDEA) Program.

