

HPTN 057

Study-Specific Procedures Manual

A PHASE I OPEN LABEL TRIAL OF THE SAFETY AND PHARMACOKINETICS OF
TENOFVIR DISOPROXIL FUMARATE IN HIV-1 INFECTED
PREGNANT WOMEN AND THEIR INFANTS
PROTOCOL VERSION 2.0, DATED 28 OCTOBER 2009



**Multicenter Study of the International Maternal Pediatric Adolescent AIDS Clinical Trials
Group (IMPAACT)**

Sponsored by:

US National Institute of Allergy and Infectious Diseases (NIAID)
And
The Eunice Kennedy Shriver
US National Institute of Child Health and Human Development (NICHD)

Pharmaceutical Support Provided by:

Gilead Sciences, Inc

XX February 2010
Version 4.0

