

## Acronyms

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AE	Adverse Event
CFR	Code of Federal Regulations
CM	Clarification Memo
CORE	Coordinating and Operations Center (FHI)
CRF	Case Report Form
CRPMC	Clinical Research Products Management Center
CTA	Clinical Trials Agreement
CV	curriculum vitae
DAIDS	Division of AIDS
DBS	dried blood spot
EAE	Expedited Adverse Event
EC	Ethics Committee
FWA	Federal Wide Assurance
GCP	Good Clinical Practices
GLP	Good Laboratory Practices
IATA	International Air Transport Association
ICH	International Conference on Harmonization
IMPAACT	International Maternal Pediatric Adolescent AIDS Clinical Trials Group
IoR	Investigator of Record
IRB	Institutional Review Board
LDMS	Laboratory Data Management System
LoA	Letter of Amendment
MOP	Manual of Operations (HPTN)
MTCT	Mother to Child Transmission
NA	not applicable
ND	not done
NIH	National Institutes of Health
OHRP	Office of Human Resource Protections
PAB	Pharmaceutical Affairs Branch
PIBA	Press-in Bottle Adapters
pk	pharmacokinetics
pMTCT	preventing Mother to Child Transmission
PRO	Protocol Registration Office
PSRT	Protocol Safety Review Team
QC/QA	Quality Control/Quality Assurance
RCC	Regulatory Compliance Center
ROS	Review of Symptoms
SDMC	Statistical and Data Management Center (SCHARP)
SOAP	subjective-objective-assessment-plan
SOP	Standard Operating Procedure
SSP	Study-Specific Procedures Manual
TDF	tenofovir disoproxil fumarate
VCT	Voluntary Counseling and Testing
VST	visit code
WHO	World Health Organization