

Section 17 - Study Reporting Plan

MTN-005 Statistical and Data Management Center (SDMC) Staff

Job Role	Name	Email Address
Protocol Statistician:	Liza Noonan	liza@scharp.org
Project Manager:	Missy Cianciola	missy@scharp.org
Statistical Research Associate:	Sharavi Gandham	sharavi@scharp.org
Protocol Programmer:	Katie Weaver	kweaver@scharp.org
Data Coordinator:	Sue Tracy-Waisanen	stracy@scharp.org
Document Specialist:	Stacie Kentop	stacie@scharp.org
Reporting Programmer	Cathy Kirkwood	ckirkwood@scharp.org
Lab Operations	Deb Bassuk	dbassuk@scharp.org
CASI Programmer	Lynda McVarish	lmcv@scharp.org
Clinical Affairs Safety Associate	Yevgeny Grigoriev	ygrigori@scharp.org

17.1 Purpose of Reporting Plan

The purpose of this reporting plan is to describe the reports that the MTN SDMC (SCHARP) plans to generate for MTN 005.

The specific purposes of this plan are:

- To identify the purpose and content of each report;
- To identify those responsible for the preparation and distribution of each report;
- To identify who should review the reports so that corrective action (if necessary) is taken; and
- To ensure the Protocol Team approves the plan prior to study initiation.

This reporting plan was prepared by the MTN 005 SDMC Project Manager in collaboration with other MTN 005 SDMC staff.

17.2 Study Reports

Table 15-1 lists the reports the SDMC will produce and distribute via email. Table 15-2 lists the reports the SDMC will produce and make available via the Atlas website:

atlas.scharp.org

Following the tables is a description of each report that includes the purpose of the report, who will prepare the report, and specific components of the report.

Table 17-1: MTN-005 SDMC Reports Distributed via Email

Report Title	Distribution Frequency	Email Distribution List
Data Quality Control (QC)	Every two weeks, or as needed	<ul style="list-style-type: none"> • Site Study Coordinators • Site Data Managers • CORE Clinical Research Managers • SDMC Project Manager
Clinical Data Quality Control (CQC) Queries	Weekly, or as needed	<ul style="list-style-type: none"> • Site Study Coordinators • Site Data Managers • CORE Clinical Research Managers • SDMC Project Manager
Unresolved Adverse Experiences (AE) Listing	Monthly	<ul style="list-style-type: none"> • Site Study Coordinators • Site Data Managers • CORE Clinical Research Managers • SDMC Project Manager
Specimen Monitoring Report	Monthly	<ul style="list-style-type: none"> • Site Study Coordinators • Network Lab Representative • SDMC Project Manager

Table 17-2: MTN-005 SDMC Reports Posted on Atlas

Report Title	Update Frequency	Atlas Viewing Area
Enrollment and Retention	Daily	Unsecure
Visit Adherence and Procedure Completion	Monthly	Unsecure
Site Data Management Quality	Monthly	Unsecure
Safety (PSRT) Reports	One week prior to each scheduled PSRT call	Secure
Network Lab Assay Results Report	Monthly, once NL results are received at the SDMC	Unsecure
Study Monitoring Committee (SMC)	As determined by the SMC	<ul style="list-style-type: none"> • MTN-005 SMC members and observers • MTN-005 Protocol Chair • MTN-005 Site Investigators

17.2.1 Data Quality Control (QC) Report

Purpose: To identify and help correct missing and inconsistent data
Prepared and Distributed by: SDMC Data Coordinator
Components: Quality control notes; overdue visit reminders, missing page reminders

17.2.2 Clinical Data Quality Control (CQC) Queries

Purpose: To identify and help correct inconsistencies/questions identified in safety or clinical data
Prepared and Distributed by: SDMC Clinical Affairs Safety Associate
Components: Queries containing clinically-based questions about safety and clinical data.

17.2.3 Unresolved Adverse Experiences (AE) Listing

Purpose: To identify AEs whose resolution/outcome requires updating on the AE Log case report form.
Prepared and Distributed by: SDMC Clinical Affairs Safety Associate
Components: Listing of AEs that have had an “continuing” status for more than 90 days.sa

17.2.4 Specimen Monitoring Report

Purpose: To monitor storage in LDMS of those specimens marked as “stored” on study CRFs
Prepared by: SDMC Laboratory Programmer
Components: Site-specific listing of all discrepancies between the CRF stored specimen data and LDMS data.

17.2.5 Enrollment and Retention Report

Purpose: To monitor participant accrual and retention as reflected by data submitted to the SDMC (via DataFax)
Prepared by: SDMC Protocol Programmer
Components: Enrollment table includes the number of women enrolled each week and cumulatively. Retention includes: total enrolled (broken down by active, inappropriately enrolled, and lost to follow-up); number expected for a given visit; number not expected for a given visit; and total retention by visit calculated as the number of participants who have completed a visit divided by the total number of participants expected for the visit.

17.2.6 Visit Adherence and Procedure Completion Report

Purpose: To summarize site performance regarding study endpoint data collection
Prepared by: SDMC Statistical Research Associate
Components: Distribution of visits, including the number of days between target and actual visit dates, and the number of days between sequential follow-up visits. Listing of number and percentage of completed key required procedures, which may include specimen collection, safety lab testing, pelvic exam completion, and pregnancy test completion.

17.2.7 Site Data Management Quality Report

Purpose: To summarize site performance regarding data management and quality.

Prepared by: SDMC Project Manager

Components: Total number of CRF pages faxed to SCHARP, total number of QCs applied, percentage of QCs resolved, QC rate per 100 CRF pages, and mean days to fax in CRF pages. Reported cumulatively and for the previous month.

17.2.8 Safety (PSRT) Reports

Purpose: To help the Protocol Safety Review Team monitor study participant safety as reflected by adverse experiences reported to the SDMC (via DataFax).

Prepared by: SDMC Reporting Programmer and SDMC Clinical Affairs Safety Associate

Components: Cumulative AE data reported to SCHARP via DataFax.

17.2.9 Network Lab Assay Results Report

Purpose: To monitor the receipt of lab assay results from the Network Lab.

Prepared by: SDMC Laboratory Programmer

Components: For each specimen analyzed by a Network Lab, the number of results expected (per CRF data) along with the number and percentage of results received and processed at SCHARP.

17.2.10 Study Monitoring Committee Report

Purpose: To monitor study progress at each site

Prepared and distributed by: Prepared by SDMC MTN 005 staff and distributed by SDMC Project Manager

Components: Summary by site and overall of study design and history, accrual, retention, demographics, product adherence, safety/adverse events, pregnancy and pregnancy outcomes. Site data management quality and other components as requested by the SMC.