

13. Study Reporting Plan

The purpose of this reporting plan is to describe the reports that the MTN SDMC (SCHARP) plans to generate for MTN-024/IPM 031.

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 follow-up (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-024/IPM 031 SDMC Project Manager in collaboration with other MTN-024/IPM 031 SDMC staff.

MTN-024/IPM 031 Statistical and Data Management Center (SDMC) Staff

Job Role	Name	Email Address
Protocol Statisticians	Elizabeth Brown Jingyang Zhang	erbrown@scharp.org jzhang2@scharp.org
Project Manager	Corey Miller	corey@scharp.org
Statistical Research Associate	Holly Gundacker	hgundack@scharp.org
Operations Programmers	Craig Chin Brad Fischer	craig@scharp.org hfischer@scharp.org
Data Coordinator	Claire Chapdu	cchapdu@scharp.org
Document Specialist	Lori Filipcic	lorif@scharp.org
Lab Programmers	Deb Bassuk Della Wilson	dbassuk@scharp.org della@scharp.org
Clinical Affairs Safety Associate	Jill Zeller	jzeller@scharp.org

13.1 Study Reports

Table 13-1 lists the reports the SDMC will produce and distribute via email. Table 13-2 lists the reports the SDMC will produce and make available via the Atlas website, <https://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 13-1: MTN-024/IPM 031 SDMC Reports Distributed via Email

Report Title	Distribution Frequency	Email Distribution List
Data Quality Control (QC) Report	Monthly, or as needed	<ul style="list-style-type: none"> • SDMC Project Manager • Site Staff as designated by each site
Clinical Data Quality Control (CQC) Report	As needed (as queries are identified)	<ul style="list-style-type: none"> • Site Staff as designated by each site • SDMC Project Manager
Unresolved Adverse Experiences (AE) Listing	Monthly	<ul style="list-style-type: none"> • Site Staff as designated by each site • SDMC Project Manager
Unresolved Product Holds	Monthly	<ul style="list-style-type: none"> • Site Staff as designated by each site • SDMC Project Manager
LDMS Specimen Monitoring Report	Monthly	<ul style="list-style-type: none"> • Site LDMS Lab Staff • Site Staff as designated by each site • Network Lab Representative • SDMC Project Manager

Table 13-2: MTN-024/IPM 031 SDMC Reports Posted on Atlas

Report Title	Update Frequency	Atlas Viewing Area
Screen Out	Daily	Unsecure
Enrollment and Retention Report	Daily	Unsecure
Visit Adherence and Procedure Completion Report	Monthly	Unsecure
Site Data Management Quality Report	Monthly	Unsecure
Data Summary Report	Monthly	Unsecure
Protocol Deviations	Daily	Secure
Protocol Safety Review Team (PSRT) Report	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) Report	As determined by the SMC	<ul style="list-style-type: none"> • MTN-024/IPM 031 SMC members and observers • MTN-024/IPM 031 Protocol Chair • MTN-024/IPM 031 Site Investigators

13.1.1 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

13.1.2 Clinical Data Quality Control (CQC) Report

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

13.1.3 Unresolved Adverse Experiences (AE) Listing

Purpose: To identify those AEs that have been continuing for 90 or more days (per the AE Log CRF) so that AE status updates are made as needed.

Prepared and distributed by: SDMC Clinical Affairs Safety Associate

Components: Listing of AEs that have had a “continuing” status for more than 90 days

13.1.4 Unresolved Product Holds

Purpose: To identify those clinical product holds that have been continuing for 30 or more days (per the PH Log CRF) so that product status updates are made as needed.

Prepared by: SDMC Programmer

Components: Listing of product holds that have been ongoing for 30 or more days.

13.1.5 LDMS Specimen Monitoring Report

Purpose: To monitor specimen storage in LDMS for 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

13.1.6 Screen Out Report

Purpose: To summarize the number of participants screened for the study, the number enrolled and the reasons participants were not enrolled.

Prepared by: SDMC Programmer

Components: Number screened, number enrolled, number screened out per reason listed on the Eligibility Criteria CRF.

13.1.7 Enrollment and Retention Report

Purpose: To monitor participant accrual and retention as reflected by data submitted to the SDMC (via DataFax)

Prepared by: SDMC Programmer

Components:

- Enrollment table includes the number of women enrolled each week and cumulatively

- Retention table includes total number of women 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 women who have completed a visit divided by the total number of women expected for the visit.

13.1.8 Visit Adherence and Procedure Completion Report

Purpose: To provide information on completion of required study procedures during follow-up, and serve as an indication as to the amount of missing data from completed visits.

Prepared by: SDMC Statistical Research Associate

Components: Distribution of visits, including the number of days between target and actual visit dates; listing of number and percentage of completed key required procedures, which may include pelvic exam completion, PK specimen collection, CVL specimen collection, CASI questionnaire completion

13.1.9 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 monthly (previous month)

13.1.10 Data Summary Report

Purpose: To summarize site performance regarding data management quality, enrollment, retention, and selected procedure completion

Prepared by: SDMC Project Manager

Components: Cumulative enrollment and retention data, cumulative procedure completion data for selected study procedures, and monthly and cumulative data management quality data

13.1.11 Protocol Safety Review Team (PSRT) Report

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 Programmer and SDMC Clinical Affairs Safety Associate

Components: Cumulative AE data reported to SCHARP via DataFax

13.1.12 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

13.1.13 Study Monitoring Committee (SMC) Report

Purpose: To monitor study progress at each site

Prepared and distributed by: Prepared by SDMC MTN-024/IPM 031 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 (e.g. procedure completion) as requested by the SMC

13.1.14 Protocol Deviations Report

Purpose: To summarize reported protocol deviations at each site

Prepared and distributed by: Prepared by SDMC Programmer

Components: Listing, by site, of reported protocol deviations as reported on Protocol Deviation Log CRFs received at SCHARP (via DataFax)