

Section 14 - Study Reporting Plan

14.	Introduction	14-1
14.1	Purpose of Reporting Plan	14-1
14.2	Study Reports	14-1

14. Introduction

The MTN-038 Statistical and Data Management Center (SDMC) Staff are listed below.

Job Role	Name	Email Address
Protocol Statistician	Barbra Richardson	barbrar@uw.edu
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Clinical Data Manager	Amanda Brown	abrown@scharp.org
Clinical Programmer	Jackie Fitzpatrick	jackie@scharp.org
Clinical Safety Associate	Christine Thompson	cethomps@scharp.org

14.1 Purpose of Reporting Plan

The purpose of this reporting plan is to describe the routine reports that the MTN SDMC (SCHARP) plans to generate for MTN-038.

The specific purposes of this plan are to:

- Identify the purpose and content of each report;
- Identify those responsible for the preparation and distribution of each report;
- Identify who should review the reports so that follow-up (if necessary) is done.

This reporting plan was prepared by the MTN-038 SDMC Clinical Data Manager in collaboration with other MTN-038 SDMC staff.

14.2 Study Reports

The reports listed in Table 14-1 are available within the Medidata web-based environment and can be run by designated site users (based on user permissions) at any time to include the most current data available in the Medidata Rave study database.

Table 14-2 lists the reports the SDMC will produce and make available via the MTN-038 Atlas web page:

<https://atlas.scharp.org/cpas/project/MTN/038/begin.view?>

Table 14-3 lists the reports the SDMC will produce and distribute via e-mail.

Following the tables is a description of each report that includes the purpose and components of the report.

Table 14-1: MTN-038 SDMC Reports Available in Medidata

Report Title	Permissions List
Site-specific Query Details	<ul style="list-style-type: none"> • Site Staff as designated by each site • SDMC Clinical Data Manager
Site-specific AE Listing	<ul style="list-style-type: none"> • Site Staff as designated by each site • SDMC Clinical Data Manager
Site-specific Enrolled PTID Listing	<ul style="list-style-type: none"> • Site Staff as designated by each site • SDMC Clinical Data Manager

Table 14-2: MTN-038 SDMC Reports Posted on Atlas

Report Title	Update Frequency	Atlas Viewing Area
Screen Out	Daily	Unsecure
Enrollment	Daily	Unsecure
Retention	Daily	Unsecure
Procedure Completion	Daily	Unsecure
Data Management Quality	Monthly	Unsecure
Data Summary	Monthly	Unsecure
Missed Visit Listing	Daily	Unsecure
Missed Visit Summary	Monthly	Unsecure
Protocol Deviations Listing	Daily	Secure
Protocol Deviations Summary Table	Monthly	Secure
PSRT (Safety)	One week prior to PSRT call	Secure
AE listings	One week prior to PSRT call	Secure
Study Monitoring Committee (SMC)	Approximately every 4-6 months	Secure

Table 14-3: MTN-038 SDMC Reports Distributed via E-mail

Report Title	Distribution Frequency	E-mail Distribution List
LDMS Specimen Monitoring	Monthly	<ul style="list-style-type: none"> • Site LDMS Laboratory Staff • MTN Laboratory Center Representative(s) • SDMC Clinical Data Managers

1. Query Details

Purpose: To provide detailed information on data queries for a given site

Components: By site, displays the query status, query user, marking group, field, form, folder, subject, site group for each data query

2. AE Listing

Purpose: To provide detailed cumulative listings of all reported Adverse Events per site during the study for clinical and data management

Components: Cumulative listing of Adverse Events (AEs) reported for each site

3. Enrolled PTID Listing

Purpose: To provide listing of PTIDs for all participants who have enrolled into the study by site per the Randomization eCRF completion within the Medidata Rave study database.

Components: Cumulative listing of PTIDs enrolled for each site

4. Screen Out

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

Components: Number screened, number enrolled, number screened out per protocol-specified ineligibility reason listed on the Inclusion/Exclusion Criteria CRF

5. Enrollment

Purpose: To report on participant accrual as reflected by data entered into the study database

Components: By site, activation date, dates of first and last enrollments, duration of accrual, enrollment target, total number screened, total number enrolled, screening to enrollment ratio, average number of enrollments per week, and percentage of site target enrolled for the MTN-038 study

6. Retention

Purpose: To report on participant visit retention as reflected by data entered into the study database

Components: By site and by visit, the number of expected participants who have completed the visit; the number of participants who have not completed the visit; the number of visits missed; the number of participants who missed a visit; the number of participants who have terminated early; the number of participants, excluding early terminators, who have completed the visit; and the number of participants not expected at each visit

7. Procedure Completion

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

Components: Overall and by site, listing of number and percentage of required (“expected”) study procedures that were completed at follow-up visits. Procedures are expected if the visit was completed (i.e., not missed).

8. Data Management Quality Report

Purpose: To provide information on site performance with regard to key data management and data quality metrics.

Components: By site and overall data metrics

9. Data Summary Reports

Purpose: To provide summary information on site performance regarding data management quality, enrollment, retention, and selected procedure completion.

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

10. Missed Visit Listing

Purpose: To identify participants who have missed scheduled study visits to help sites focus retention efforts and prevent participants from becoming chronic defaulters

Components: Site-specific listing of cumulative missed visits per the Missed Visit CRF; includes, for each PTID, the enrollment date, visit name, start and end of visit window

11. Missed Visit Summary

Purpose: To provide a subset of Protocol Team members with a cumulative summary of all missed visits for the study

Components: Overall and by site, the number and percentages of missed visits reported for the study

12. Protocol Deviations Listing

Purpose: To provide a subset of Protocol Team members with a cumulative listing of all protocol deviations reported for the study.

Components: Each of the fields/data items as listed on the Protocol Deviations Log CRF.

13. Protocol Deviations Summary Table

Purpose: To provide a subset of Protocol Team members with a cumulative and past month summary of all protocol deviations for the study

Components: Overall and by site, the number and percentages of protocol deviations reported for the study

14. PSRT (Safety) Reports

Purpose: To help the Protocol Safety Review Team (PSRT) monitor participant safety as reflected by adverse events, pregnancies, product holds, and study product discontinuations reported to the SDMC.

Components: Cumulative AE and study product discontinuations reported to the SDMC on the AE Log CRF and Discontinuation of Study Product CRF.

15. AE Listings

Purpose: To provide the MTN-038 Safety Physicians with a cumulative listing of all adverse events in order to monitor participant safety.

Components: Cumulative listing of all adverse events reports to the SDMC per the Adverse Event Log CRF

16. Study Monitoring Committee (SMC) Reports

Purpose: To provide information on study conduct, ability to answer study objectives, and primary endpoint data to SMC members as required in preparation for scheduled reviews

Components: Summary by site and overall of study design and history, participant accrual, retention, completion of primary and main secondary endpoint assessments, data management quality, protocol deviations, and other components as requested by the SMC

17. LDMS Specimen Monitoring

Purpose: To identify stored specimens whose information in LDMS does not match corresponding information collected per study CRFs

Components: Listing of those specimens whose LDMS PTID, visit code, and/or collection date information does not match the information recorded on CRFs; specimens that are stored per CRF but not present in LDMS; specimens that are present in LDMS but not stored per CRF; specimens in LDMS from PTIDs who did not enroll