	Participant ID: _____	Pin # _____
	Discovery Site: _____	Clinical Center _____
	CRF Date: ____/____/____	Visit #: _____

Study Stop Point

***For EPS Pt.s:** Research Coordinator completes at Twelve-month in-clinic contact or at final contact if Participant withdraws from the study early.

***For Healthy/Positive Control Pt.s:** Research Coordinator completes at the conclusion of the Baseline visit.

1. Has the EPS participant successfully completed the 12-month phenotyping visit of the Trans-MAPP Epidemiology Phenotyping Study? ₁ Yes ₀ No
-OR-
 Has the Healthy/Positive Control Participant successfully completed the Baseline visit?

If **No**, indicate reason for withdrawal:


- a. No longer willing to follow the protocol/interested in participating ₁ Yes ₀ No
- b. Lost to follow-up ₁ Yes ₀ No
- c. Participant has personal constraints ₁ Yes ₀ No
- d. Medical condition/event ₁ Yes ₀ No
- e. Physician's Discretion ₁ Yes ₀ No
- f. Other ₁ Yes ₀ No
 Specify: _____

Female Participants only:

- g. Female Participant is pregnant ₁ Yes ₀ No ₉₉ NA
- g1. If **Yes**, date of most recent menstrual period: _____
 (MM/DD/YYYY)

2. Number of Participant's last Contact: _____

3. Date that the participant was last seen: _____
 (MM/DD/YYYY)

	Participant ID: _____	Pin # _____
	Discovery Site: _____	Clinical Center _____
	CRF Date: ____/____/____	Visit #: _____

Study Stop Point

*For EPS Pt.s: Research Coordinator completes at Twelve-month in-clinic contact or at final contact if Participant withdraws from the study early.

*For Healthy/Positive Control Pt.s: Research Coordinator completes at the conclusion of the Baseline visit.

The following section is for Study Close-out.

(PRINCIPAL INVESTIGATOR AND RESEARCH COORDINATOR COMPLETE WHEN PARTICIPANT STOPS PARTICIPATION IN THE STUDY.)

4. Physician Comments (optional): _____

SIGNATURES: Please complete the following section regardless of the reason for termination of study participation.

I verify that all information collected on the Trans-MAPP Epidemiology Phenotyping Study data collection forms for this participant is correct to the best of my knowledge and was collected in accordance with the procedures outlined in the Trans-MAPP Epidemiology Phenotyping Study Protocol and Manual of Procedures.

 Principal Investigator's Signature Date: ____/____/____
 (MM/DD/YYYY)

5. Did the PI sign this form? Yes No

 Research Coordinator's Signature Date: ____/____/____
 (MM/DD/YYYY)

6. Did the RC sign this form? Yes No

7. Research Coordinator ID: _____ (4-digit ID)