Pregnant Woman Eligibility Note: Before enrolling a woman, make sure you have enough supplies (study kits, blood and urine collection materials, paper forms, etc.)

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Name of Person Completing the Form:		
Cedula of Pregnant Woman:		
Today's date:/(DD/MMM/YYYY)		
IPS Information		
Clinic name:		
<u>_</u>		
City: □ Barranquilla □ Bucaramanga □ Tuluá		
Pregnant Woman Information		
Last names:		
First name(s):		
Pregnant woman date of birthI(DD/MMM/YYYY) If under 18, date of pregnant woman 18th birthday/	_/(DD	/MMM/YYYY)
Expected delivery date (EDD):/(DD/MMM/YYYY)	Unsure	
How determined: by LMP by ultrasound		
If expected delivery date is unknown:		
Date of LMP:I (DD/MMM/YYYY)	nsure	
Gestational age at eligibility: weeks days		
Inclusion Criteria		
le the consequence of the confirmed division and aliminate of the confirmed division and the confirmed	☐ Yes	□ No
Is the woman pregnant? (by confirmed clinic pregnancy test or ultrasound)	⊔ res	□ INO
Is the woman in the first trimester of pregnancy (≤14 ⁺⁶)? (by LMP or ultrasound)	☐ Yes	□ No
Is the woman planning to have prenatal care at a participating clinic?	\square Yes	\square No
Is the woman 16 years of age or older?	\square Yes	\square No
Does this woman speak Spanish?	☐ Yes	□ No

CDC estimates the average public reporting burden for this collection of information as 5 minutes per response, including the time for reviewing instructions, searching existing data/information sources, gathering and maintaining the data/information needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information,

Last updated 05/19/17 including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0920-XXXX). **Exclusion Criteria** ☐ Yes ☐ No Is the pregnancy an ectopic or a molar pregnancy? ☐ Yes □ No Is the woman incarcerated? Is the woman unable to physically or psychologically participate ☐ No ☐ Yes based on clinical judgement? **Eligibility Determination** The woman is eligible for the study. (All answers to inclusion criteria questions are Yes AND all answers to exclusion criteria are No.) ☐ Yes Eligible ☐ No \rightarrow Not Eligible ☐ Unsure \rightarrow If unsure, then fill out Appendix E1 (contact information) and follow-up in one week. Notes about eligibility determination: If eligible, please complete the sections below. Informed Consent Determination Did the woman receive and provide informed consent or assent for participation? ☐ Yes \rightarrow Enroll ☐ Unsure \rightarrow If unsure, then fill out Appendix E1 (contact information) and followup in one week. ☐ No \rightarrow If no, thank the patient for their time and note that they declined participation in the study. Reason(s) for declining (check all that apply): ☐ Not interested ☐ Concerned about study protocol (safety, invasive) ☐ Concerns about time/transportation ☐ Concerns about family member approval (e.g. partner, parents) ☐ Other concern: Zika Study Kit Did you give the woman a Study Kit before she left? ☐ Yes, she took it ☐ Offered it to her, but she didn't want/take it ☐ Did not offer → **STOP**. Do not enroll if no study kit is available. Schedule enrollment visit for another day.

ZEN Participant and Non-Participant Identification Numbers

Appendix B1 ZEN Colombia Pregnant Woman Eligibility

If the woman was eligible and has consented, assign a ZEN Participant ID. If the woman was not eligible or was eligible and did not consent, assign a ZEN Non-Participant ID (see SOP ZEN-2-02).

If not offered, why?: _____