CRF	CRF – Eligibility Screener		
No.	Question	Response	
001	Please record today's date:	/ /	
		D D / M M / Y Y Y Y	
002	Did the participant provide written consent for this study prior to conduct of any study procedures?	0 = No End form 1 = Yes	
003	Did the participant consent to off-site visits?	0 = No 1 = Yes	
004	Did the participant consent to photos/video for infant?	0 = No 1 = Yes	
005	Has participant enrolled in MATRIX- 007 for a previous pregnancy?	0 = No 1 = Yes	
006	Enter the PID that was used for the mother previously:	Respond only if [005 = 1]	
Pregn	ancy, HIV, and Syphilis Status		
007	Does the participant report they are currently pregnant?	<pre>0 = No End form 1 = Yes 2 = Not sure</pre>	
800	Has participant received an ultrasound in the last 5 days that confirmed the pregnancy?	0 = No Skip to 011 1 = Yes	
009	Does participant have the results of the ultrasound with them today that includes an intrauterine pregnancy with a documented gestational age measured in weeks and days and/or the estimated date of delivery, and with fetal heart tones detected?	0 = No Skip to 011 1 = Yes (viable pregnancy)	

010	Date of Ultrasound	Respond only if [009 = 1] Date should not exceed 5 days before today's date. / /
011	Has participant received a urine or serum human chorionic gonadotropin (hCG) (pregnancy) test within the last 5 days as part of antenatal care or other health services with the results available?	0 = No Skip to 014 1 = Yes
012	Date of previous test:	Date should not exceed 5 days before today's date. / / D D / M M / Y Y Y Y
013	Result of previous hCG test:	<pre>0 = Negative End form 1 = Positive Skip to 015</pre>
014	Will hCG testing (urine pregnancy test) to confirm the pregnancy status be needed at this MATRIX-007 visit?	0 = No 1 = Yes
015	Has the participant had HIV testing within the last 5 days as part of antenatal care or other health services with available results?	0 = No Skip to 019 1 = Yes
016	Enter date that HIV testing was performed:	Date should not exceed 5 days before today's date. / / D D / M M / Y Y Y Y

017	Enter the HIV testing outcome done within the last 5 days of today's visit:	<pre>0 = HIV negative Skip to 020 1 = HIV positive (seroconversion confirmed) Skip to 018 then end form 2 = Inconclusive End form</pre>
018	Does the participant have ART documentation for seroconversion?	<pre>0 = No 1 = Yes (seroconversion confirmed) END screening after responding, participant is ineligible.</pre>
019	Will rapid HIV testing be needed at this MATRIX-007 visit? Note: If no, but syphilis testing is needed, HIV testing will be included in the combo rapid test.	0 = No End form 1 = Yes
020	Has the participant received a syphilis test within the last 3 months and during this pregnancy from a healthcare provider with the results available?	<pre>0 = No/Not sure Skip to 025 1 = Yes</pre>
021	Enter date that syphilis testing was performed:	Date should not exceed 3 months before today's date
022	Type of previous syphilis testing:	<pre>0 = RPR or VDRL 1 = Treponemal antibody 2 = Rapid (with or without HIV testing)</pre>
023	Was there a positive test result?	<pre>0 = No Skip to 025 1 = Yes Submit a PSRT query and add to Medical Conditions/Events CRF once enrolled.</pre>
024	Has the participant received appropriate treatment?	Respond only if [023 = 1] 0 = No 1 = Yes 2 = Don't know/Uncertain

025	Will rapid syphilis/HIV (combo) testing be needed at this MATRIX-007 visit?	0 = No 1 = Yes	
	HCG, HIV, syphilis Test Results Perform any needed pregnancy, HIV and/ syphilis testing and document results below		
026	Result of urine hCG test performed today by MATRIX-007 study:	Respond only if [014 = 1] 0 = Negative End form 1 = Positive 2 = Not done	
027	Result of rapid HIV test done today by MATRIX-007 study:	Respond only if [019 = 1] 0 = HIV negative 1 = HIV positive End form 2 = Not done	
028	Results of syphilis testing done today by MATRIX-007 study:	Respond only if [025 = 1] 0 = Not reactive 1 = Reactive 2 = Not done	
Gesta	Gestational Age		
029	What is the participant's best estimate of the first day of the last normal menstrual period (LMP)?	/ /	
030	Is this LMP certain or uncertain?	<pre>0 = Uncertain 1 = Certain</pre>	
031	Follow this link to identify important dates in the pregnancy Enter best available input: LMP, EDD, or DoC	https://perinatology.com/calculators/bue-Date.htm	
032	Using the calculator at the perinatology.com website or other study-approved calculator, what is the estimated date of conception?		

033	Using the calculator at the perinatology.com website or other study-approved calculator, what is the estimated due date (EDD)?	
034	Using the calculator at the perinatology.com website or other study-approved calculator, what is the gestational age today? Enter the WEEKS output of the GA	
035	Using the calculator at the perinatology.com website or other study-approved calculator, what is the gestational age today? Enter the DAYS output of the GA	
036	Calculated gestational age (total days):	If [≥34 weeks] then END
Expos	ure Assessment	
037	Has the participant ever been a cohort participant in CATALYST?	0 = No End form 1 = Yes
038	Date of 180 days before estimated date of conception:	/ /
039	Read aloud to participant: Have you received a CAB PrEP injection since this date (insert date above)?	0 = No 1 = Yes
040	Read aloud to participant: In the last 28 days, have you used and had the PrEP ring inserted for at least 21 of those days?	0 = No 1 = Yes

041	Read aloud to participant: In the last 28 days, have you taken oral PrEP/tenofovir-based PrEP for at least 21 of those days?	0 = No End form 1 = Yes
Behav	rioral Assessment	
042	Read aloud to participant: Are you currently receiving or do you plan to receive antenatal care?	0 = No End form 1 = Yes
043	Read aloud to participant: Do you intend to relocate away from any of the study-linked CARE PrEP sites within time frame of the study (now up through 6 months postpartum)?	0 = No 1 = Yes End form
044	Read aloud to participant: Do you intend to travel away from any of the study-linked CARE PrEP sites for three or more months?	0 = No 1 = Yes End form
045	Does the participant have adequate locator information?	0 = No End form 1 = Yes
Scree	ning Outcome	
046	Is there any other reason or condition to deem the participant ineligible for the study? Such as that would preclude informed consent, make study participation unsafe for the participant, complicate interpretation of study outcome data, or otherwise interfere with achieving the study objectives.	0 = No Skip to 047 then end form. 1 = Yes

047	Please describe the reason/conditions why the participant is ineligible.	Respond only if [046 = 1] then END
048	Participant Eligibility:	<pre>0 = "Eligible" 1 = "Not Eligible" If participant is exposed only to DVR or Oral PrEP and GA<4 weeks pregnant, then INELIGIBLE until GA>4 weeks.</pre>
049	Enrollment Status:	<pre>0 = Enrolled 1 = Pending exposure status/form completion 2 = Failed - eligible but not interested 3 = Failed - not eligible 4 = Failed - declined to complete eligibility assessment</pre>
050	Participant must meet all eligibility requirements in order to enroll.	