## **BIRTH COHORT-3 SCREENING CASE RECORD FORM**

Study ID	<u>  B   3    0                               </u>	Patient Initials:	Date of Screening:   _	_ /  /	<u> </u>
		Last First	day	month	vear

Study Selection Criteria						
Date of Birth:   _ /     Age in years: day month year	When any box in the "Exclude" column is ticked stop & proceed to the last question. If not excluded, proceed to the next section.					
Selection criteria	Include	Exclude				
Age ≥16 years	□ Yes	□ No				
Residence within Busia District, Uganda	□ Yes	□ No				
Agreement to come to the study clinic for any febrile episode or other illness	□ Yes	□ No				
Agreement to avoid medications given outside the study protocol	□ Yes	□ No				
Plan to deliver in the hospital	□ Yes	□ No				
History of adverse event to SP or DP	□ No	□ Yes				
Active medical problem requiring inpatient evaluation at the time of screening	□ No	□ Yes				
Intention of moving outside of Busia District, Uganda	□ No	□ Yes				
Chronic medical condition requiring frequent medical attention	□ No	□ Yes				
Prior SP preventive therapy or any other antimalarial therapy during pregnancy	□ No	□ Yes				
Confirmed HIV-Status by rapid test (Attach copy of source document)	□ Negative	□ Positive				
Confirmed viable intrauterine pregnancy by ultrasound	□ Yes	□ No				
Estimated gestation age between 12-20 weeks:  LMP known □ Yes □ No  LMP:     /      (EDD by LMP = LMP + 280 days)  EDD by LMP:     /    /    EDD by US:     /    /     # days difference between LMP and US:  First trimester (6-12w): Use US EDD if more than 7 days difference or LMP unknown  Second trimester (13-24w): Use US EDD if more than 14 days difference or LMP unknown  Third trimester (>24w): Use US EDD if more than 21 days difference or LMP unknown  Final EDD:     /    /    GA:     w     d	□ 12-20wks	□ <12wks □ >20wks				
Early or active labor (documented by cervical change with uterine contractions)	□ No	□ Yes				
Mother's willingness to provide informed consent for herself and her unborn child	□ Yes	□ No				
Mother provided informed consent for use of biological specimens for her and her child*? □ Yes □ No						
Is father of the unborn child readily available to provide consent?    Yes   No						
If yes, is father of the unborn child willing to provide consent?	□ Yes	□ No				
(Mother can still participate in the study if father is not readily available to provide consent)						
All criteria for study inclusion met? □ Yes □ No (If yes, proceed to the enrolment form)						
Initials:						

9 March 2016

Version 1.0

Entered\_\_\_\_\_\_ Date\_\_\_/\_\_/\_\_
Verified\_\_\_\_\_ Date\_\_\_/\_\_/\_\_