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BMJ Open Cohort profile: effect of malaria in early pregnancy on fetal growth in Benin (RECIPAL preconceptional cohort)

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ABSTRACT

Purpose Retard de Croissance Intra-uterin et PALudisme (RECIPAL) is an original preconceptional cohort designed to assess the consequences of malaria during the first trimester of pregnancy, which is a poorly investigated period in Africa and during which malaria may be detrimental to the fetus.

Participants For this purpose, a total of 1214 women of reproductive age living in Sô-Ava and Akassato districts (south Benin) were followed up monthly from June 2014 to December 2016 until 411 of them became pregnant. A large range of health determinants was collected both before and during pregnancy from the first weeks of gestation to delivery. Five Doppler ultrasound scans were performed for early dating of the pregnancy and longitudinal fetal growth assessment.

Findings to date Pregnant women were identified at a mean of 6.9 weeks of gestation (wg). Preliminary results confirmed the high prevalence of malaria in the first trimester of pregnancy, with more than 25.4% of women presenting at least one microscopic malarial infection during this period. Most infections occurred before six wg. The prevalence of low birth weight, small birth weight for gestational age (according to INTERGROWTH-21st charts) and preterm birth was 9.3%, 18.3% and 12.6%, respectively.

Future plans Retard de Croissance Intra-uterin et PALudisme (RECIPAL) represents at this time a unique resource that will provide information on multiple infectious (including malaria), biological, nutritional and environmental determinants in relation to health outcomes in women of reproductive age, pregnant women and their newborns. It will contribute to better define future recommendations for the prevention of malaria in early pregnancy and maternal malnutrition in Africa. It confirms that it is possible to constitute a preconceptional pregnancy cohort in Africa and provides valuable information for researchers starting cohorts in the future.

INTRODUCTION

Malaria in pregnancy is associated with a wide range of deleterious effects in women and their offspring. In Sub-Saharan African (SSA)

Strengths and limitations of this study

- A unique cohort combining highly detailed, high-quality health-related information on the pregnancies of approximately 400 women recruited in the preconception period, with a substantial related biobank of plasmas, placental and other samples.
- Preconceptional design allowing the early identification and follow-up of pregnant women: screening of women for both microscopic and submicroscopic malaria from the first weeks of pregnancy; accurate estimation of gestational age by using early obstetrical ultrasound scan and collection of some important factors influencing fetal growth such as prepregnancy nutritional status and gestational hypertensive disorders.
- Collection of valuable information for the implementation of future preconceptional cohorts in Africa.
- High attrition in the preconceptional cohort, with a possible impact on the external validity of some findings.
- Reduced sample size at delivery due to a high proportion of spontaneous abortion, with a possible lack of power for analyses on birth outcomes.

countries, preventive strategies are based on long-lasting insecticide-treated bed nets (LLITNs) and intermittent preventive treatment in pregnancy (IPTp). IPTp consists in the administration of sulfadoxine-pyrimethamine (SP) at each antenatal care (ANC) visit from the second trimester of pregnancy onwards.¹ LLITN are distributed at the first ANC visit, which generally occurs around 4–5 months of pregnancy.² Therefore, pregnant women remain unprotected or insufficiently protected during the first months of pregnancy and particularly during the first trimester. However, this period may be a high-risk period for the fetus if pregnancy-associated malarial parasites



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accumulate into the placenta during trophoblast differentiation and vascular remodelling of the uterus.^{3,4} Impaired placentation may then contribute to fetal growth restriction and low birth weight (LBW).

Few studies have investigated malaria in early pregnancy, and while some have shown an adverse effect on birth outcomes, this has not been consistent.⁵ In Burkina Faso and in Benin, malarial infection in the first trimester⁶ or before 4 to 5 months of pregnancy^{7,8} has been associated with a higher risk of LBW or a decrease in birth weight. Conversely, such association was not reported in Uganda and Malawi,^{9,10} but the number of women screened in the first trimester was low in these two studies. One of the problems in interpreting this existing literature relates to the challenges of recruiting women early in pregnancy; most of the studies were not designed to assess the consequences of malaria early in pregnancy and therefore women were generally recruited late in the first trimester, which may lead to misclassification errors and underestimation of exposure. Besides, using LBW as a proxy for fetal growth restriction (FGR) can lead to the overestimation of FGR prevalence and possible bias when estimating the effect

of malaria. Finally, maternal nutrition is one of the main factors influencing fetal growth in developing countries.¹¹ Therefore, adjusting for this factor when assessing the effect of early malaria on FGR is important. As well, a recent study has suggested a higher risk of FGR due to malaria in undernourished women.¹²

The REtard de Croissance Intra-uterin et PALudisme (RECIPAL) study aimed to assess the effect of malaria (both microscopic and submicroscopic) in the first trimester of pregnancy on both the mother and the fetus, with a focus on fetal growth, by following a cohort of pregnant women recruited before conception. Moreover, it also aimed at assessing the influence of the woman's nutritional status in the relationship between malaria in early pregnancy and birth outcomes.

COHORT DESCRIPTION

Study setting

The study cohort is currently conducted in the districts of Sô-Ava and Abomey-Calavi, located in Southern Benin (figure 1). Women's recruitment has been completed by

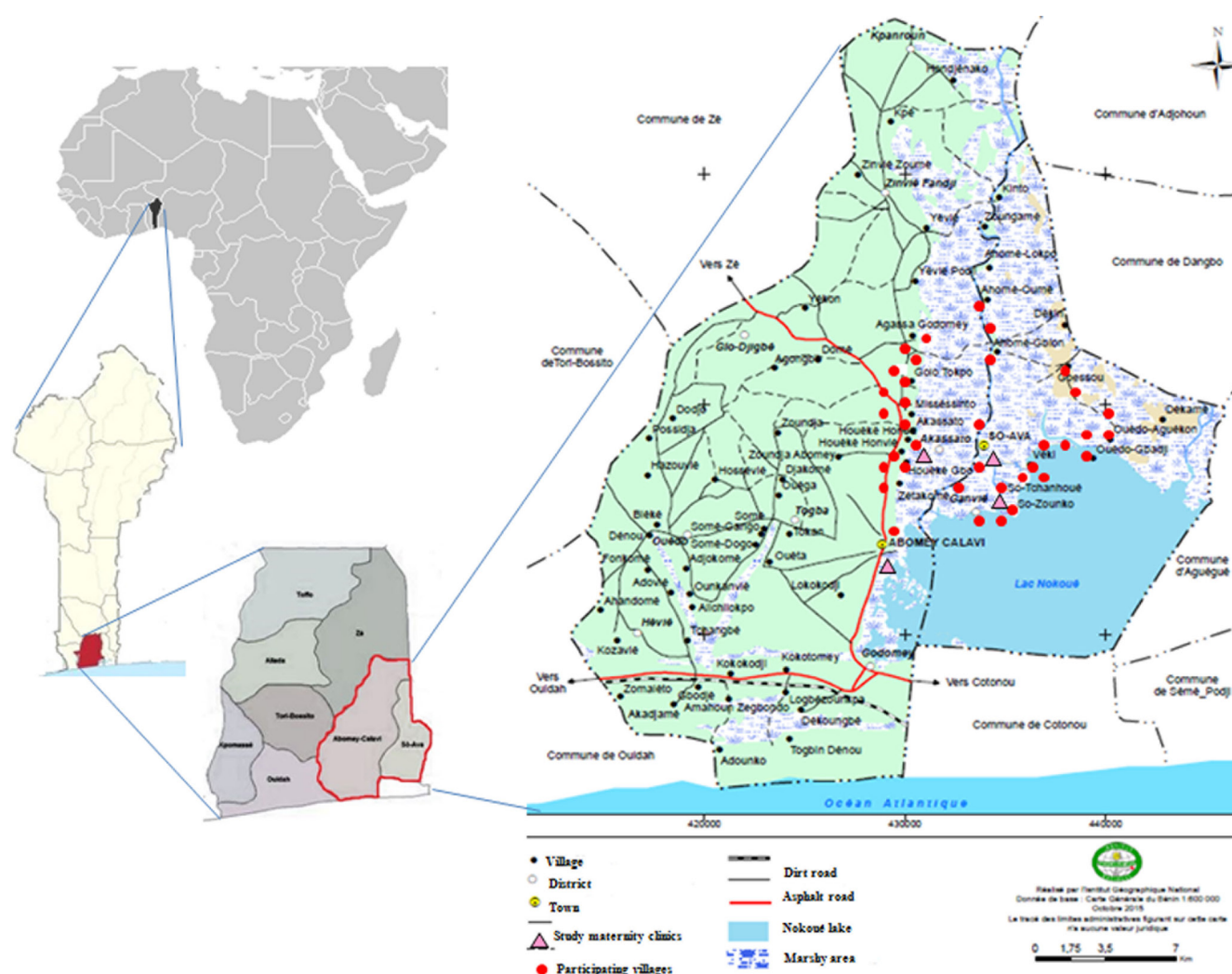


Figure 1 Geographical location of Sô-Ava and Abomey-Calavi districts and of the 35 villages of the RECIPAL study, Southern Benin, 2014–2017. RECIPAL, REtard de Croissance Intra-uterin et PALudisme.

December 2016, and pregnancy follow-up is still ongoing. In the area, malaria is hyperendemic with a mean entomological inoculation rate of 2.1 infected anopheles bites/person/100 nights.¹³ The main mosquito vectors of malaria are *Anopheles gambiae* ss and *Anopheles funestus*.¹⁴ Four subdistricts (Sô-Ava, Vekky and Houedo in Sô-Ava District and Akassato in Abomey-Calavi District) were selected for the study according to the population density and mean number of ANC visits and deliveries per month in the four main public maternity clinics of Sô-Ava and Abomey-Calavi districts. In these four subdistricts, the population size was estimated to 124994 people in 2013, including 17.4% (21 779) women of reproductive age (WRA).¹⁵ Thirty-five out of a total of 36 villages were then selected for the study according to their proximity to the maternity clinics.

Study design, subject identification, recruitment and enrolment procedures

Briefly, WRAs were recruited at community level and followed monthly for a maximum period of 24 months until becoming pregnant. They constituted the RECIPAL initial cohort. The subsample of women who became pregnant was then followed up monthly at the maternity clinic from early pregnancy to delivery. They constituted the RECIPAL final cohort.

A sample size of 466 births was estimated sufficient to evidence a two-fold OR for the association between malaria and poor birth outcomes,⁴ assuming a prevalence of 25% of women with microscopic malaria in the first trimester of pregnancy and 30% of poor birth outcomes (LBW, small-for-gestational age (SGA), preterm birth (PTB) or stillbirth)^{8 16 17} with an 80% power, significant level at 0.05. We estimated that following 2000 WRA (initial cohort) would have allowed identifying 510 pregnant women (final cohort) during a planned 18-month period for recruitment in the initial cohort (based on a total fertility rate of 200/1000/year in Benin (<http://esa.un.org/unpd/wpp/index.htm>)) and allowing 15% of loss to follow-up after recruitment in the initial cohort.

After explanation of the study to the local political and administrative authorities, three concomitant procedures were used to recruit WRA. First, repeated sensitisation events were organised in each of the 35 selected villages for presenting the study to all inhabitants. If interested, women were invited to register with the head of the community and were visited at home the day after for their inclusion. That procedure constituted the main way of recruitment throughout the study. Second, leaders of the community, religious authorities and mass media were involved to help us mobilise women during public meetings (masses, women's association meetings, etc). The last way of recruitment consisted of going door to door to identify eligible and interested women with the assistance a network of community health workers (more details are provided in the online supplementary file S1).

To be included, women had met the following criteria: negative urinary pregnancy test, 18 to 45 years old, no current contraception, no previous fecundity issues,

willingness to become pregnant, no planned travel for more than 2 months within the next 18 months, acceptance of RECIPAL protocol and signed written informed consent. Both oral and written communication was used for providing information on the project to the women and their husbands. The consent form was translated into the local language for women who were illiterate.

From June 2014 to December 2016, 1302 WRA were willing to participate in the study. Among them, 1214 (93.2%) met the inclusion criteria and were recruited, 88 (6.8%) were not included mainly because of a positive urinary pregnancy test at inclusion (63.6%) or past infertility issues (14.8%). Based on both the 2013 national census and the 2011–2012 Demographic Health Survey (DHS-IV) in Benin, we estimated that women included in the project represented 6% of the total number of WRA living in the study area. Compared with women included in Dansou *et al* study,¹⁸ which used individual data of a representative sample of WRA included in the Beninese DHS-IV, WRA included in RECIPAL had a higher level of poverty. The other sociodemographic characteristics such as age, education level, household wealth, gravidity and media exposure were, however, similar (online supplementary file S2).

RECIPAL study was approved by the ethics committees of the Institut des Sciences Biomédicales Appliquées (ISBA) in Benin and from the National Research Institute for Sustainable Development in France as well as by the Ministry of Health in Benin.

Cohort follow-up

Preconceptional follow-up

After enrolment, WRAs were visited at home monthly by specifically trained investigators assisted by local community health workers. Follow-up consisted mainly of recording the first day of last menstrual period (LMP) and performing a urinary pregnancy test every month. To minimise the risk of loss of confidentiality during the tracking of women for follow-up, the study team was instructed to speak only to people—including family members—to whom women have granted them permission to speak. After 12 months, women who were still followed received medical advices to help conceive, and they were invited to attend the maternity clinic for a medical examination in case of symptoms of genital infection. After 24 months, the follow-up ended and women who did not conceive were invited to attend a gynaecologist for fertility issues assessment. More details on the follow-up procedures have been provided in the online supplementary file S3.

Gestational follow-up

Once pregnant, women were followed up at the maternity clinic for monthly ANC visits. In addition, they were encouraged to attend the maternity clinic any time in case of any symptoms. Both the maternity staff (for usual ANC follow-up and clinical examination) and study investigators (for all specific issues related to RECIPAL) were involved in women's follow-up. Throughout the study, women were offered free transport to the maternity clinic; in Sô-Ava

District, a free river shuttle was set up. At 37 weeks of gestation (wg), women were visited at home weekly and transportation to the study maternity clinic was scheduled for delivery.

According to Beninese national recommendations, a kit including iron and folic acid tablets for 1 month, mebendazole for deworming, the first dose of SP for IPTp and a LLITN was given to each pregnant woman at the first ANC visit. The maternity staff was encouraged to administer at least three doses of IPTp from the second trimester onwards as recommended by the Beninese National Malaria Control Programme, as well as iron and folic acid supplementation throughout the pregnancy. RECIPAL women benefited from free management of any diseases related to the pregnancy—detected either as part of RECIPAL follow-up or during unscheduled visits, free ANC visits and delivery. Women infected with malaria were treated with quinine in the first trimester of pregnancy and artemether–lumefantrine from the second trimester.

Infant follow-up

The infant follow-up was not initially planned as part of the RECIPAL project. It is currently carried out as part of the SEPSIS project (bioMérieux-funding, UMR216/9198 coordination), which aims to assess immune dysfunctions associated with the risk of sepsis in preterm and growth-restricted newborns as well as in those exposed to malaria in utero. All infants born from the RECIPAL women from April 2016 have been included and followed until 3 months of age.

Cohort attrition

Among the 411 pregnant women, 207 (50.4%) have already delivered, 16.5% (68/411) had a spontaneous abortion (n=68) or a non-viable pregnancy (n=2), 62 did not complete the study because of informed consent withdrawal (45/411, 10.9%) or migration (17/411, 4.1%)

and 72 women are still followed (figure 2). Spontaneous abortions, informed consent withdrawals and migrations occurred at 8.9 (± 3.9), 13.2 (± 8.1) and 14.2 (± 8.6) wg in average (SD), respectively. More details on the reasons for cohort attrition are provided in the online supplementary file S4.

Figure 2 presents the flowchart diagram of the study. Among the 1214 WRA enrolled in the initial cohort, 411 (33.9%) became pregnant, 359 (29.6%) completed the follow-up without conceiving and 444 women (36.6%) did not complete the study mainly because of informed consent withdrawal (272/444) or migration outside the study area (138/444) that occurred after 5.2 and 6.3 months of follow-up in average, respectively.

Data collection

Preconceptional data

Demographic, socioeconomic and household characteristics were collected at inclusion (table 1). At this occasion, screening for malaria, urinary schistosomiasis, *Chlamydia trachomatis* and *Neisseria gonorrhoeae* was performed. Haemoglobin (Hb) and markers of inflammation levels were determined. The first day of LMP was recorded and a urinary pregnancy test (One-Step; International Holding, Germany) was performed each month. Anthropometric measurements were collected every 3 months, and dietary intakes were assessed twice during follow-up.

Gestational data

At each ANC visit, clinical, anthropometric and obstetrical data, as well as LLITN use, IPTp administration, alcohol consumption and iron, folic acid and other micronutrient supplementation intake were collected (table 2). In addition, malarial screening was performed, and proteinuria, glycosuria and urinary infection were detected using a urine dipstick test (Combi-screen;

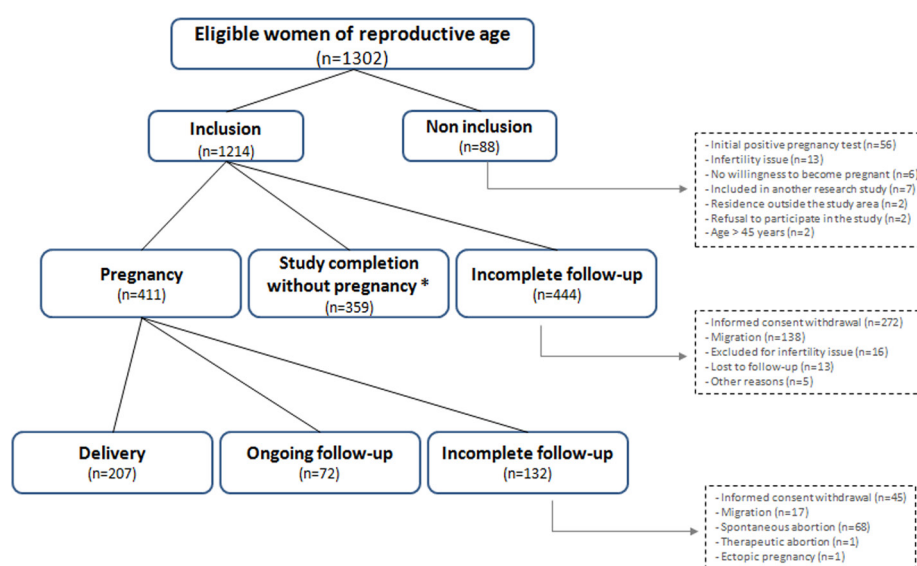


Figure 2 Flow chart diagram of RECIPAL study.* Study completion: follow-up from enrolment until the end of the study (24-month follow-up without pregnancy for women recruited before December 2014 or monthly follow-up without pregnancy for women recruited between December 2014 and December 2016), excluding consent withdrawal, migration and lost to follow-up. RECIPAL, RETard de Croissance Intra-uterin et PALudisme.

Table 1 Clinical, nutritional and biological data collected during the preconceptional follow-up—the RECI PAL study, 2014–2017

	Inclusion	D1	M1	M2, M3	M4	M5, M6	M7	M8, M9	M10	M11, M12	M13	M14, M15	M16	M17, M18	M19	M20, M21	M22
General characteristics																	
Place of visit	*	†	†	*	*	*	*	*	†	*	*	*	*	*	*	*	*
Eligibility criteria	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Informed consent	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
GPS location	†	\$	\$	\$	\$	\$	\$	–	\$	\$	\$	\$	\$	\$	\$	\$	\$
Sociodemographic characteristics	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Housing characteristics	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Economic characteristics	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Clinical and nutritional data																	
Obstetrical history	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Height	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Weight, MUAC, skinfold thickness	\$	†	†	\$	†	\$	†	\$	†	\$	†	\$	†	\$	†	\$	†
Body composition (BIA)	\$	†	†	\$	\$	\$	\$	\$	†	\$	\$	\$	\$	\$	\$	\$	\$
Dietary 24 hours recall	\$	†	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Amenorrhoea/LMP	\$	†	†	†	†	†	†	†	†	†	†	†	†	†	†	†	†
Biological data																	
Urinary pregnancy test	†	\$	†	†	†	†	†	†	†	†	†	†	†	†	†	†	†
Malarial screening (TBS, PCR)	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Hb, ferritin, folic acid levels	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
CRP, AGP, sTfR levels	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
<i>Chlamydia trachomatis</i> †	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
<i>Neisseria gonorrhoeae</i> †	\$	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Urinary schistosomiasis†	\$	†	†	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$	\$
Urine storage**	\$	†	†	\$	\$	\$	†	\$	†	\$	\$	\$	\$	\$	\$	\$	\$

*Household visit.

†Visit performed at the study maternity clinic.

‡Information collected/assessed at each specific time point.

§Information not collected/assessed at each specific time point.

¶*Chlamydia trachomatis* and *Neisseria gonorrhoeae* screening (using PCR and serology), as well as schistosomiasis screening (Nytel filter) started 9 and 12 months after study initiation, respectively.

**Urine storage for further analysis.

AGP, alpha1-acid glycoprotein; BIA, bioimpedance analysis; CRP, C-reactive protein; D, day; Hb, haemoglobin; GPS, global positioning system; LMP, last menstrual period; M, months; MUAC, mid-upper-arm-circumference; RECI PAL, REtard de Croissance Intra-uterin et PALudisme; sTfR, soluble transferrin receptor; TBS, thick blood smear.

Table 2 Clinical, nutritional and biological data collected during pregnancy—the RECIPAL study, 2014–2017

	ANCv 1	ANCv 2	ANCv 3	ANCv 4	ANCv 5	ANCv 6	ANCv 7	ANCv8	Delivery
Clinical and nutritional data									
Gestational age (weeks), mean (SD)	6.8±2.5	11.6±2.9	16.7±2.9	22.1±3.7	27.6±3.8	32.7±3.5	36.2±2.5	38.4±1.4	39.1±3.0
Medical and obstetrical history*†	‡	§	§	§	§	§	§	§	§
Alcohol consumption in the last 24 hours	‡	‡	‡	‡	‡	‡	‡	‡	§
Iron and folate intake in the last 24 hours	§	‡	‡	‡	‡	‡	‡	‡	§
Use of ITN the night before the visit	‡	‡	‡	‡	‡	‡	‡	‡	§
Axillary temperature, blood pressure	‡	‡	‡	‡	‡	‡	‡	‡	‡
Gestational age (based on LMP or first US)	‡	‡	‡	‡	‡	‡	‡	‡	‡
Anthropometric measurements									
Weight	‡	‡	‡	‡	‡	‡	‡	‡	‡
Mid-upper-arm-circumference	‡	§	§	‡	§	‡	§	§	§
Skinfold thickness (bicipital and tricipital)	‡	§	§	‡	§	‡	§	§	§
Body composition (bioimpedance analysis)	‡	‡	‡	‡	‡	‡	‡	‡	§
Dietary 24 hours recall	§	‡	§	§	‡	§	‡	§	§
Biological data									
Blood and rhesus group	‡	§	§	§	§	§	§	§	§
HIV 1 screening	‡	§	§	§	§	§	§	§	§
Peripheral malaria (TBS, PCR)	‡	‡	‡	‡	‡	‡	‡	‡	‡
Placental malaria (TBS, PCR, histology)	§	§	§	§	§	§	§	§	‡
Cord blood malaria (TBS, PCR)	§	§	§	§	§	§	§	§	‡
Hb level	§	‡	§	§	§	‡	§	§	§
Serum ferritin, sTfR, CRP, AGP, folic acid levels	§	‡	§	§	§	‡	§	§	§
Lead level	§	‡	§	§	§	‡	§	§	§
Urinary infection and proteinuria (dipstick test)	‡	‡	‡	‡	‡	‡	‡	‡	§
Helminthic intestinal infection (PCR)	‡	§	§	§	§	§	§	§	§
Urinary schistosomiasis	§	§	§	§	§	§	‡	§	§
Urine storage¶	§	‡	§	‡	§	‡	§	§	§

*Sickle cell disease, diabetes or other chronic medical affection; history of preterm delivery or hypertensive disorders.

†History of cigarette smoking during pregnancy collected at delivery.

‡Information collected/assessed at each specific time point.

§Information not collected/assessed at each specific time point.

¶Urine storage for further analysis.

AGP, alpha1-acid glycoprotein; ANCv, antenatal care visit; CRP, C-reactive protein; Hb, haemoglobin; ITN, insecticide-treated bed net; LMP, last menstrual period; RECIPAL, REtard de Croissance Intra-uterin et PALudisme; sTfR, soluble transferrin receptor; TBS, thick blood smear; US, ultrasound scan.

Analyticon, Germany). Dietary intakes were assessed once every trimester. Five Doppler ultrasound scans (US) were performed. A sample of venous blood was collected in the first and third trimesters for Hb, markers of inflammation and lead levels determination.

At delivery, newborns were weighed within 1 hour after birth on an electronic digital scale with an accuracy

of 2 g (SECA 757; SECA, Germany). Newborn's length (SECA infantometer 416, Germany) and head circumference (SECA 201, Germany) were recorded to the nearest millimetre. Malarial screening was performed on maternal venous blood, placental and cord blood (table 2). A placental biopsy was performed for malarial histopathology. Follow-up and quality control

Table 3 Biological samples stored before conception, during pregnancy and at birth—the RECIPAL study, 2014–2017

	Prepregnancy	Pregnancy*	Delivery/birth	Planned analyses
Total blood				
Total blood (200 µL)	–	Mother (x2)	–	Erythrocytic folic acid
Total blood (500 µL)	–	Mother (x2)	Mother–placenta cord	Metal level
Blood spot +4°C (50 µL)	Mother (x1)	Mother (x8)†	Mother–placenta cord	<i>Plasmodium</i> detection (PCR)
Blood spot –20°C (100 µL)	Mother (x1)	Mother (x8)†	Mother–placenta cord	Immunological markers
Red cells (Trizol, 100 µL)	Mother (x1)	Mother (x8)	Mother–placenta cord	<i>Plasmodium</i> phenotyping
Buffy coat PAX gene (300 µL)	–	Mother (x2)	Mother–placenta cord	Immunological markers
Plasma/serum				
Plasma –20°C‡	Mother (x1)	Mother (x2)	Mother–placenta cord	Micro-RNAs, cytokines and chemokines
Plasma –80°C‡	Mother (x1)	Mother (x2)	Mother–placenta cord	Micro-RNAs, cytokines and chemokines
Serum (100 µL)	Mother (x1)		–	<i>Chlamydia trachomatis</i> and <i>Neisseria gonorrhoeae</i> detection
Serum (400 µL)	–	Mother (x2)		Vitamin A
Serum (100 µL)‡	Mother (x1)	Mother (x2)	–	Ferritin, CRP, sTfR, AGP
Urine (1.8 mL)	Mother (x2)	Mother (x3)	–	Pollutant/pesticide
Stool (–80°C)	Mother (x1)	–	–	Helminthic infection
Placenta				
Biopsy with formalin	–	–	Placenta	Histopathology for malaria
Biopsy –20°C	–	–	Placenta	Immunological markers
Biopsy with RNA later	–	–	Placenta	Immunological markers

The frequency of sample collection is indicated in parentheses.

*During pregnancy, samples were collected twice (during the first and third trimester of pregnancy), three times (during each trimester) or eight times (during each ANC visit).

†In case of unscheduled visit, an additional blood spot was collected.

‡Quantity of plasma stored before, during pregnancy and at delivery was 200, 600–800, 600 µL, respectively.

AGP, alpha1-acid glycoprotein; ANC, antenatal care; CRP, C reactive protein; RECIPAL, RETard de Croissance Intra-uterin et PALudisme; sTfR, soluble transferrin receptor.

procedures are provided in the online supplementary file S3.

The types and origins of the biological samples collected before and during pregnancy are listed in table 3. Total blood, serum, plasma urine samples are stored in –20°C or –80°C freezers with a daily monitoring of the temperature.

Specifics components

Malarial diagnosis

Before conception and during pregnancy, malarial screening was performed using both microscopy (thick blood smear) and PCR. Blood smears were stained with Giemsa and parasitaemia was quantified by the Lambaréné method.¹⁹ A molecular diagnostic approach, using a combination of real-time PCR assays comprising genus-specific primers and probes for the gene encoding the small (18S) *Plasmodium* rRNA and an ultrasensitive *Plasmodium falciparum*-specific

detection system²⁰ were used to screen samples containing *Plasmodium* parasites. In addition, any time during pregnancy, a rapid diagnostic test (Pf+pan rapid test SD Bioline Ag, IDA Foundation, Netherlands; BioSynex, France) was performed in case of symptoms suggestive of malaria for immediate diagnosis and treatment.

Ultrasound follow-up

The first US for dating the pregnancy was performed between 9 and 13 wg based on LMP recorded during the preconceptional follow-up (see online supplementary file S5). The gestational age (GA) was estimated in days based on crown-rump length measurement using the Robinson's chart.²¹ At the end, GA estimation was based either on LMP if the difference between the two measurements (LMP/US) was less than 7 days or on US if the difference is >7 days. Following INTERGROWTH-21st methodology, four additional scans were performed for fetal biometry

assessment and uterine and umbilical blood flow measurements.²² Each parameter was measured twice and then averaged; a third measure was performed in case of discrepancy between the two first measurements. A quality control was performed on 10% of the pictures.

Anthropometric measurements and dietary intake assessment

Anthropometric measurements were collected using standard procedures (tables 1 and 2).²³ During household visits, women's body weight was measured with a 200 g precision with calibrated electronic scales (Tefal, France). At the facility level, women were weighed with a Tanita MC-780 body composition device (Tanita, Tokyo, Japan). Height was measured to the nearest millimetre with a SECA 206 (Hamburg, Germany) gauge. The left mid-upper-arm-circumference was measured with a SECA 201 (Hamburg, Germany) ergonomic circumference measuring tape. Skin-fold thickness measurements were made to the nearest millimetre at the triceps and biceps sites using a Holtain (Crymyeh, UK) skinfold calliper. The set of measurements was repeated twice—by the same field investigator—and then the measurements were averaged. Body composition, assessed by bioelectrical impedance analysis method, was measured by the multifrequency body composition analyser Tanita MC-780 with a four-electrode arrangement paired on hands and feet. A quality control of anthropometric data was performed periodically by a senior research scientist in nutrition.

Dietary intakes were assessed by 24-hour recalls using the standard multiple-pass methodology.²⁴ A comprehensive list of the most common food items and recipes was developed from a preliminary study of the diet habits of the study area. This list was used to identify foods mentioned by the women as consumed during the interview. Amounts of food consumed were estimated by household measures consisted of precalibrated utensils, food pictures of different portion sizes and monetary value of known food portions sold on the study area markets. A food composition table and recipes database are under construction for conversion of household units to gram and to determine macronutrients and micronutrients intake by the women.

Characteristics of the study populations

At enrolment, WRA were 27 years old in average (table 4). Nulligravidae represented 8.8% of the cohort. More than half of the women were anaemic, one quarter was infected with schistosomiasis and 38% had an abnormal body mass index. The prevalence of microscopic malaria was 5.7%, with two geographical clusters in Sô-Ava and Vekky subdistricts (figure 3). The median duration of follow-up in the initial cohort was 5.9 months (mean=7.1, range: 0.49–23.7). Compared with women included in the initial cohort, women who did not complete the follow-up were significantly different in terms of residence area, ethnicity, household density and schistosomiasis status (table 4).

Pregnant women were identified at a mean of 6.9 wg and benefited from 7.5 scheduled ANC visits in average (table 4). The prevalence of microscopic malarial infection

was 25.4%, 19.4% and 16.1% during the first, second and third trimester of pregnancy, respectively. The prevalence of SGA was 11.5% and 18.3% using Schmiegelow's and INTERGROWTH-21st charts, respectively (table 5). The proportion of PTB and LBW was 12.6% and 9.3%, respectively. Overall, 29.5% (61/207) of newborns presented at least one poor birth outcome defined as LBW, PTB, SGA or stillbirth.

FINDINGS TO DATE

Fertility and time to pregnancy

The median time to pregnancy was 12.3 months (figure 4). The fertility rate was 5.4 pregnancies/100 persons-month (95% CI =4.9 to 5.9). The probability (95% CI) of conceiving after 5, 10, 15 and 20 months of follow-up were 24.5% (21.9–27.4), 42.1% (38.6–45.9), 56.6% (52.1–61.1) and 62.8% (57.6–68), respectively. Using multivariate survival analysis, factors associated with a lower risk of conceiving were maternal age more than 35 years, a low education level, being polygamous, living in Vekky subdistrict, long-term couple (>8 years), infrequent sexual activity, overweight and urinary schistosomiasis infection (E Yovo, personal communication, 2017).

Malaria in the first trimester of pregnancy

The incidence rate of malarial infection during the first trimester was 19.7 cases per 100 persons-month (95% CI 15.8 to 24.5). Using a multilevel logistic regression, we showed that women infected with malaria before conception were more likely to be infected during the first trimester (adjusted OR (aOR): 2.68, 95% CI 1.24 to 5.78). Gestational age was also negatively correlated with malarial infection (aOR: 0.64, 95% CI 0.41 to 0.98) (M Accrombessi, personal communication, 2017).

STRENGTHS AND LIMITATIONS

The preconceptional design—and early identification of pregnant women—allowed (1) the screening of women for malaria from the very beginning of pregnancy; (2) the accurate estimation of GA by using US before 14 wg; (3) the determination of prepregnancy nutritional status²⁵ and hypertensive disorders screening,²⁶ which highly influence fetal growth and (4) the minimisation of selection bias embedded in the design of women recruited at facility level. Besides, PTB and intrauterine growth restriction could be accurately assessed thanks to the precise estimation of GA and longitudinal fetal growth assessment.

In addition to malaria, RECIPAL will provide valuable information on WRA in Africa in terms of fertility and nutrition, which are seldom evaluated and may help to define new preconceptional interventions for improving maternal and child health.²⁷ Also, it yields logistical and ethical information for the implementation of future preconceptional cohorts.

Table 4 Characteristics of women included in the initial and final RECIPAL cohorts, Southern Benin, 2014–2017

Characteristics		Initial cohort (n=1214 WRA)	WRA with incomplete follow-up (n=444)	P value (1214 vs 444)*	WRA with complete follow-up but no pregnancy (n=359)	Final cohort Pregnant women (n=411)	P value (411 vs 359)*
General characteristics							
Subdistrict, n (%)	Sô-Ava	279 (23.0)	102 (23.0)		45 (12.5)	132 (32.1)	
	Vekky	423 (34.8)	228 (51.4)	<0.001	57 (15.9)	138 (33.8)	<0.001
	Houedo	186 (15.3)	33 (7.4)		120 (33.4)	33 (7.8)	
	Akassato	326 (26.9)	81 (18.2)		137 (38.2)	108 (26.3)	
Ethnic group, n (%)	Toffin	864 (71.2)	346 (77.9)		218 (60.7)	300 (72.9)	
	Aizo	195 (16.1)	56 (12.6)	0.001	80 (22.3)	59 (14.4)	0.002
	Fon	86 (7.1)	23 (5.2)		31 (8.6)	32 (7.8)	
	Others	69 (5.6)	19 (4.3)		30 (8.4)	20 (4.9)	
Age (years)	Mean (\pm SD)	27.8 (\pm 5.5)	28.0 (\pm 5.6)	0.31	28.8 (\pm 5.6)	26.8 (\pm 5.0)	<0.001
Marital status, n (%)	Polygamy	445 (36.7)	173 (38.9)		154 (42.9)	118 (28.7)	
	Monogamy	729 (60.0)	257 (57.9)	0.45	204 (56.8)	268 (65.2)	<0.001
	Cohabitation	40 (3.3)	14 (3.1)		1 (0.3)	25 (6.1)	
Education, n (%)	Illiterate	889 (73.2)	336 (75.7)	0.14	263 (73.3)	290 (70.6)	0.41
Household density	Mean (\pm SD)	5.7 (\pm 3.0)	6.0 (\pm 3.4)	0.009	5.2 (\pm 2.3)	5.9 (\pm 3.0)	0.001
Gravidity, n (%)	Nulligravidae	107 (8.8)	48 (10.8)		26 (7.2)	–	
	Primigravidae	167 (13.8)	58 (13.1)	0.17	43 (12.0)	33 (8.0)	0.22
	Multigravidae	940 (77.4)	338 (76.1)		290 (80.8)	378 (92.0)	
Number of live birth(s)	Mean (\pm SD)	2.8 (\pm 2.0)	2.6 (\pm 2.0)	0.07	2.9 (\pm 2.1)	2.7 (\pm 1.9)	0.16
Short stature, n (%)	Maternal length <150 cm	92 (8.0)	27 (6.1)	0.46	32 (8.9)	33 (8.0)	0.65
Mid-upper-arm-circumference, n (%)	<23 cm	76 (6.3)	23 (5.2)	0.62	27 (7.5)	26 (6.3)	0.51
Preconceptional characteristics							
BMI, n (%)	<18.5 kg/m ²	95 (8.3)	26 (6.9)		29 (8.1)	40 (9.7)	
	18.5–24 kg/m ²	706 (61.7)	229 (61.1)	0.36	212 (59.2)	265 (64.5)	0.10
	\geq 25 kg/m ²	343 (30.0)	120 (32.0)		117 (32.7)	106 (25.8)	
Schistosomiasis infection, n (%)†	Yes	177 (24.7)	50 (31.2)	0.03	63 (19.6)	64 (27.0)	0.04
Anaemia, n (%)‡	Yes	596 (49.1)	210 (47.3)	0.07	165 (46.0)	221 (53.8)	0.03
Microscopic malaria, n (%)§	Yes	64 (5.7)	28 (7.5)	0.06	12 (3.4)	24 (5.9)	0.10
Gestational characteristics							
Median time to pregnancy (months)	Median	–	–	–	–	12.3	–
Fertility incidence rate (95% CI)		–	–	–	–	5.4 (4.9–5.9)	–
Gestational age at first ANC visit (weeks)	Mean (\pm SD)	–	–	–	–	6.9 (\pm 2.5)	–

Continued

Table 4 Continued

Characteristics		Initial cohort (n=1214 WRA)	WRA with incomplete follow-up (n=444)	P value (1214 vs 444)*	WRA with complete follow-up but no pregnancy (n=359)	Final cohort Pregnant women (n=411)	P value (411 vs 359)*
Gravidity, n (%)	Primigravidae	–	–	–	–	33 (8.0)	–
	Multigravidae	–	–	–	–	378 (92.0)	–
Number of IPTp doses, n (%)**	≥2	–	–	–	–	136 (65.7)	–
Inadequate gestational weight gain†, n (%)**	Yes	–	–	–	–	120 (57.9)	–
Number of ANC visits (scheduled)**	Mean (±SD)	–	–	–	–	7.5 (±1.1)	–
Number of ANC visits (unscheduled)**	Mean (±SD)	–	–	–	–	2.1 (±1.3)	–
HIV status, n (%)**	Positive	–	–	–	–	4 (1.9)	–
Gestational hypertension, n (%)**	Yes	–	–	–	–	7 (3.4)	–
Anaemia, n (%)‡**	During pregnancy‡‡	–	–	–	–	142 (68.6)	–
	In the first trimester	–	–	–	–	98 (48.5)	–
	In the third trimester	–	–	–	–	118 (57.0)	–
Microscopic malaria, n (%)**	During pregnancy‡‡	–	–	–	–	93 (44.9)	–
	In the first trimester ‡‡	–	–	–	–	51 (25.4)	–
	In the second trimester‡‡	–	–	–	–	40 (19.4)	–
	In the third trimester‡‡	–	–	–	–	33 (16.1)	–
Placental malarial infection, n (%)**	Yes	–	–	–	–	13 (7.8)	–

*Student's t-test and χ^2 test were used for comparing continuous and categorical variables, respectively.

†Schistosomiasis infection status before conception has been assessed in 718 WRA (initial cohort): 160 with incomplete preconceptional follow-up, 321 with complete preconceptional follow-up without pregnancy and 237 included in the final cohort.

‡According to WHO thresholds (12 g/dL for non-pregnant women and 11 g/dL for pregnant).

§Before conception, 78 missing and 5 missing values in the initial and final cohort, respectively. During pregnancy, 6, 1 and 2 missing values at first, second and third trimester of pregnancy, respectively; 40 missing values for placental malaria.

¶Incidence rate defined as the number of pregnancies/persons-month at risk.

**Results based on the 207 women who have already delivered.

††A gestational weight gain was considered inadequate when the total weight gain during pregnancy was below to 12.5 kg, 11.5 kg, 7 kg and 5 kg in the underweight women (prepregnancy BMI < 18.5 kg/m²), normal weighted women (prepregnancy BMI between 18.5 and 24.9 kg/m²), overweight women (prepregnancy BMI between 25.0 and 29.9 kg/m²) and obese women (prepregnancy BMI ≥30 kg/m²) before conception, respectively.

‡‡ At least one episode.

ANC, antenatal care; BMI, body mass index; IPTp, intermittent preventive treatment in pregnancy; RECIPAL, REtard de Croissance Intra-uterin et PALudisme; WRA, women of reproductive age.

Only 107 (8%) nulligravidae were included in the study. The main reason is that RECIPAL was implemented shortly after a study, which has recruited nulligravid WRAs living

in the same area. This study aimed at assessing maternal immune responses in early pregnancy to help designing a phase 1 vaccine trial against malaria. It included 278

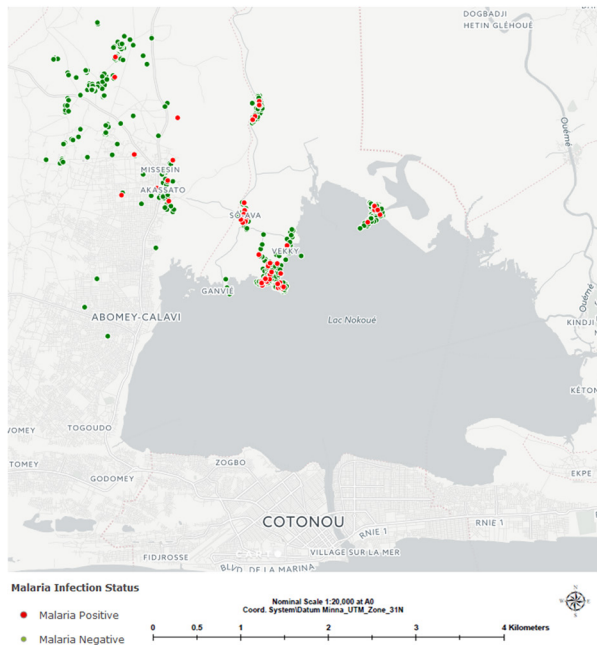


Figure 3 Malarial status (infected vs non-infected) in women of reproductive age at inclusion in RECIPAL (initial cohort), Southern Benin, 2014–2017. For online interactive map, check this link: Malaria Infection Status of Women at Inclusion of Cohort. RECIPAL, REtard de Croissance Intra-uterin et PALudisme.

nulligravid WRAs, of whom 60 became pregnant.²⁸ Both studies were conducted concomitantly for several months and women already included in the first study could not be eligible for RECIPAL. Because primigravidae were under-represented in the final RECIPAL cohort, it is likely that the prevalence of maternal conditions related to primigravidity were underestimated. Although we cannot completely exclude selection bias, it should not have strongly influenced the association between malaria in early pregnancy and FGR.

Finally, because of constraints related to the study design (long duration of the preconceptional follow-up and weariness of the women) and African sociocultural realities (rumour about blood collection, suspicion regarding medical care provided free of charge, etc), a noticeable proportion (22%) of women withdrew their informed consent before the end of the 24-month follow-up. Overall, the initial cohort attrition was 37%. Since some baseline characteristics in women who completed the preconceptional follow-up and those who did not were different, this might impact the external validity of some of results. During pregnancy, there was a high proportion (16.5%) of spontaneous abortions, which contributed to the final cohort attrition. At the end, birth outcomes

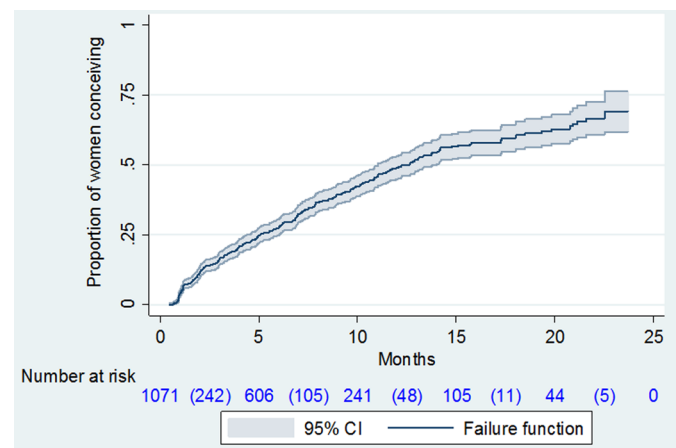


Figure 4 Kaplan-Meier failure estimate of the probability of conceiving; 1214 women of reproductive age included in the initial RECIPAL cohort, Southern Benin, 2014–2017. Probability of conceiving (solid line) and its 95% CI (grey lines). Number of pregnancy events are in parentheses. Number of censored women at 5, 10, 15, 20 and 24 months were 223, 260, 88, 50 and 30, respectively. RECIPAL, REtard de Croissance Intra-uterin et PALudisme.

Table 5 Preliminary data description of newborns at birth (subsample of the cohort)—the RECIPAL study (2014–2017)

Newborn characteristics*		n	Mean (\pm SD) or proportion (%)
Gender	Male	205	55.2
Stillbirth	Per 1000 live births	206	14.8
Preterm birth (<37 weeks)	Yes	207	12.6
Small birth weight for gestational age †‡§	Yes	191	11.5
Small birth weight for gestational age ‡§	Yes	191	18.3
Birth weight (g)‡		193	3043.9 (\pm 432.8)
	<2500	193	9.3
Poor birth outcome‡¶	Yes	207	29.5

*Data based on the first 207 deliveries.

†Small birth weight for gestational age: <10th percentile of birth weight for gestational age using Schmiegelow's charts.

‡Twins excluded.

§Small birth weight for gestational age: <10th percentile of birth weight for gestational age using INTERGROWTH-21st charts.

¶Stillbirth, preterm birth, small birth weight for gestational age (using INTERGROWTH-21st charts) or low birth weight.

RECIPAL, REtard de Croissance Intra-uterin et PALudisme.

could be evaluated in only 68% of women, with a possible lack of power for some upcoming analyses.

CONCLUSION

If the deleterious effects of malaria in the first trimester of pregnancy are confirmed, our results may argue in favour of starting preventive measures from the very beginning of pregnancy or even before pregnancy. Artemisinin-based combination therapies have been shown to be safe during the first trimester of pregnancy and, therefore, may be a good option for the replacement of SP.²⁹ Besides, a vaccine against pregnancy-associated malarial parasites, which could elicit protective immunity prior to pregnancy to best protect pregnant women during early pregnancy may be proposed as a complementary strategy. Such a vaccine is currently under evaluation.³⁰

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Contributors Project management: MA, NF, AM, MC and VB (principal investigator). Field, epidemiology and data collection: MA, EY, GC, GA, PA, NF and VB. Nutritional data collection: YM-P, AG, NF-F, DD, MA and EY. Biology and molecular analyses: NF, NT-N and SH. Obstetrical data collection: MA, EY, NJ and VB. Statistical analysis: MA, GC, MC, JZ, FB-L, YM-P, AG, NF-F, DD and VB. Manuscript writing: MA, GA, GC, AG, YM-P, NFF, DD, JZ, NTM, FB-L, SH, MC, NF and VB.

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Competing interests None declared.

Patient consent Obtained.

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Data sharing statement The researchers associated with this study are open for the sharing and reuse of the data, but an end-user data use agreement will be required for accessing the data. Collaborations are encouraged, although data sets are not currently publicly available. Any researcher interested in exploring RECIPAL data should contact directly the principal investigator, Valerie Briand (email: valerie.briand@ird.fr).

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REFERENCES

1. WHO Malaria Policy Advisory Committee and Secretariat. Malaria Policy Advisory Committee to the WHO: conclusions and recommendations of eighth biannual meeting (September 2015). *Malar J* 2016;15:117.
2. Crawley J, Hill J, Yartey J, et al. From evidence to action? Challenges to policy change and programme delivery for malaria in pregnancy. *Lancet Infect Dis* 2007;7:145–55.
3. Umbers AJ, Aitken EH, Rogerson SJ. Malaria in pregnancy: small babies, big problem. *Trends Parasitol* 2011;27:168–75.
4. Griffin JB, Lokomba V, Landis SH, et al. Plasmodium falciparum parasitaemia in the first half of pregnancy, uterine and umbilical artery blood flow, and foetal growth: a longitudinal Doppler ultrasound study. *Malar J* 2012;11:319.
5. Huynh BT, Cottrell G, Cot M, et al. Burden of malaria in early pregnancy: a neglected problem? *Clin Infect Dis* 2015;60:598–604.
6. Valea I, Tinto H, Drabo MK, et al. An analysis of timing and frequency of malaria infection during pregnancy in relation to the risk of low birth weight, anaemia and perinatal mortality in Burkina Faso. *Malar J* 2012;11:71.
7. Cottrell G, Mary JY, Barro D, et al. The importance of the period of malarial infection during pregnancy on birth weight in tropical Africa. *Am J Trop Med Hyg* 2007;76:849–54.
8. Huynh BT, Fievet N, Gbaguidi G, et al. Influence of the timing of malaria infection during pregnancy on birth weight and on maternal anaemia in Benin. *Am J Trop Med Hyg* 2011;85:214–20.
9. De Beaudrap P, Turyakira E, White LJ, et al. Impact of malaria during pregnancy on pregnancy outcomes in a Ugandan prospective cohort with intensive malaria screening and prompt treatment. *Malar J* 2013;12:139.
10. Kalilani-Phiri L, Thesing PC, Nyirenda OM, et al. Timing of malaria infection during pregnancy has characteristic maternal, infant and placental outcomes. *PLoS One* 2013;8:e74643.
11. Osrin D, de L Costello AM. Maternal nutrition and fetal growth: practical issues in international health. *Semin Neonatol* 2000;5:209–19.
12. Cates JE, Unger HW, Briand V, et al. Malaria, malnutrition, and birthweight: a meta-analysis using individual participant data. *PLoS Med* 2017;14:e1002373.
13. Djénontin A, Bio-Bangana S, Moiroux N, et al. Culicidae diversity, malaria transmission and insecticide resistance alleles in malaria vectors in Ouidah-Kpomasse-Tori district from Benin (West Africa): A pre-intervention study. *Parasit Vectors* 2010;3:83.
14. Corbel V, N'Guessan R, Brengues C, et al. Multiple insecticide resistance mechanisms in *Anopheles gambiae* and *Culex quinquefasciatus* from Benin, West Africa. *Acta Trop* 2007;101:207–16.
15. Institut National de la Statistique et de l'Analyse Economique. Fourth (2013) general census of population and habitation in Benin. 2015.
16. Briand V, Saal J, Ghafari C, et al. Fetal growth restriction is associated with malaria in pregnancy: a prospective longitudinal study in Benin. *J Infect Dis* 2016;214:417–25.
17. Unger HW, Orme-Kaius M, Karl S, et al. Factors associated with ultrasound-aided detection of suboptimal fetal growth in a malaria-endemic area in Papua New Guinea. *BMC Pregnancy Childbirth* 2015;15:83.

18. Dansou J, Adekunle AO, Arowojolu AO. Factors associated with antenatal care services utilisation patterns amongst reproductive age women in Benin Republic: an analysis of 2011/2012 benin republic's demographic and health survey data. *Niger Postgrad Med J* 2017;24:67–74.
19. Planche T, Krishna S, Kombila M, *et al.* Comparison of methods for the rapid laboratory assessment of children with malaria. *Am J Trop Med Hyg* 2001;65:599–602.
20. Hofmann N, Mwingira F, Shekalaghe S, *et al.* Ultra-sensitive detection of *Plasmodium falciparum* by amplification of multi-copy subtelomeric targets. *PLoS Med* 2015;12:e1001788.
21. Robinson HP, Fleming JE. A critical evaluation of sonar "crown-rump length" measurements. *Br J Obstet Gynaecol* 1975;82:702–10.
22. Papageorgiou AT, Sarris I, Ioannou C, *et al.* Ultrasound methodology used to construct the fetal growth standards in the INTERGROWTH-21st Project. *Int J Gynaecol Obstet* 2013;120(Suppl 2):27–32.
23. Lohman T, Roche A, Martorell R. *Anthropometric standardisation reference manual*. Champaign, IL: Human Kinetics, 1998.
24. Gibson R, Gibson RS. *Principles of nutritional assessment (second edition)*. Oxford, UK: Oxford University Press, 2005.
25. Kramer MS. The epidemiology of adverse pregnancy outcomes: an overview. *J Nutr* 2003;133:1592S–6.
26. Ndao CT, Dumont A, Fievet N, *et al.* Placental malarial infection as a risk factor for hypertensive disorders during pregnancy in Africa: a case-control study in an urban area of Senegal, West Africa. *Am J Epidemiol* 2009;170:847–53.
27. Persson LÅ, Arifeen S, Ekström EC, *et al.* Effects of prenatal micronutrient and early food supplementation on maternal hemoglobin, birth weight, and infant mortality among children in Bangladesh: the MINIMat randomized trial. *JAMA* 2012;307:2050–9.
28. Gbédandé K, Fievet N, Viwami F, *et al.* Clinical development of a VAR2CSA-based placental malaria vaccine PAMVAC: quantifying vaccine antigen-specific memory B & T cell activity in Beninese primigravidae. *Vaccine* 2017;35:3474–81.
29. Moore KA, Simpson JA, Paw MK, *et al.* Safety of artemisinins in first trimester of prospectively followed pregnancies: an observational study. *Lancet Infect Dis* 2016;16:576–83.
30. Tuikue-Ndam N, Deloron P. Developing vaccines to prevent malaria in pregnant women. *Expert Opin Biol Ther* 2015;15:1173–82.