

Effects of Weekly Iron and Folic Acid Supplements on Malaria Risk in Nulliparous Women in Burkina Faso: A Periconceptional, Double-Blind, Randomized Controlled Noninferiority Trial

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Background. The safety of iron supplementation for young women is uncertain in malaria-endemic settings.

Methods. This was a double-blind, randomized controlled noninferiority trial in rural Burkina Faso.

Results. A total of 1959 nulliparae were assigned to weekly supplementation (60 mg iron and 2.8 mg folic acid) (n = 980) or 2.8 mg folic acid (n = 979) until first antenatal visit (ANC1), or 18 months if remaining nonpregnant. Three hundred fifteen women attended ANC1, and 916 remained nonpregnant. There was no difference at ANC1 in parasitemia prevalence (iron, 53.4% [95% confidence interval [CI], 45.7%–61.0%]; control, 55.3% [95% CI, 47.3%–62.9%]; prevalence ratio, 0.97 [95% CI, .79–1.18]; *P* = .82), anemia (adjusted effect, 0.96 [95% CI, .83–1.10]; *P* = .52), iron deficiency (adjusted risk ratio [aRR], 0.84 [95% CI, .46–1.54]; *P* = .58), or plasma iron biomarkers. Outcomes in nonpregnant women were parasitemia (iron, 42.9% [95% CI, 38.3%–47.5%]; control, 39.2% [95% CI, 34.9%–43.7%]; prevalence ratio, 1.09 [95% CI, .93–1.28]; *P* = .282); anemia (aRR, 0.90 [95% CI, .78–1.05]; *P* = .17), and iron deficiency (aRR, 0.99 [95% CI, .77–1.28]; *P* = .96), with no iron biomarker differences.

Conclusions. Weekly iron supplementation did not increase malaria risk, improve iron status, or reduce anemia in young, mostly adolescent menstruating women, nor in early pregnancy. World Health Organization Guidelines for universal supplementation for young nulliparous women may need reassessment.

Clinical Trials Registration. NCT01210040.

Keywords. iron; malaria; adolescents; nonpregnant; pregnant.

Iron deficiency debilitates millions of young women in malaria-endemic areas [1]. Iron supplementation is recommended where anemia is prevalent, but its safety and efficacy in malarious areas is uncertain. Reviews of daily or intermittent iron supplementation of nonpregnant women include few studies from malaria-endemic settings [2–4], and in iron supplementation

trials in pregnancy, malaria outcomes are mostly not reported [5]. Two trials showed no increased malaria risk at delivery with daily antenatal iron supplementation, but malaria exposure was very low in one trial [6]. The other recruited mainly multigravidae [7], although in sub-Saharan Africa the highest prevalence of *Plasmodium falciparum* malaria occurs during early pregnancy in primigravidae [8]. World Health Organization (WHO) guidelines on iron supplementation are applicable to both all-age pregnant women and all-age menstruating women. Safety of iron–folic acid supplementation in young nulliparous women before and during their first pregnancy in malaria-endemic areas has not been evaluated [9], and primigravidae are a higher malaria risk group than multigravidae.

As there had been no periconceptional trials of the safety and efficacy of iron supplementation in young nulliparous women living in malaria-endemic areas, we carried out a double-blind, randomized controlled noninferiority trial of malaria risk prior to and during early pregnancy in nulliparous women receiving

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weekly iron and folic acid supplementation over a period of up to 18 months. The primary outcome was malaria parasitemia at first antenatal visit (ANC1). Secondary outcomes assessed malaria in women who remained nonpregnant, and supplement efficacy in reducing iron deficiency. Malaria endemicity in this study area is typical of many settings in sub-Saharan Africa with perennial malaria transmission.

METHODS

This periconceptional trial compared 2 cohorts supplemented with iron and folic acid vs folic acid alone: nonpregnant women and women who experienced pregnancy (additional details in [Supplementary Data 1](#)). The study received ethical approval in Burkina Faso (Comité d’Ethique pour la Recherche en Santé), the United Kingdom (Liverpool School of Tropical Medicine), and Belgium (University Hospital, Antwerp). All women in the trial provided written informed consent.

The study was conducted (April 2011–January 2014) in the Nanoro Health and Demographic Surveillance System area (DHSS) where malaria is hyperendemic, with highest transmission between June and December [10]. The iron regimen used followed WHO guidelines updated in 2016 [3, 11] and satisfied recommended nutrient intakes for nonpregnant women assuming 10%–25% iron absorption, with potential for good compliance and reduced side effects compared to daily use.

Sample

Eligible participants were nulliparous, nonpregnant residents in 30 villages within the DHSS area. Potential participants aged 15–24 years identified through the DHSS database were approached by female field assistants (FFAs) for prescreening (additional details on informed consent procedures are shown in [Supplementary Data 1](#)). We excluded women with possible or confirmed pregnancy, chronic disease (eg, sickle cell), or illness requiring hospitalization. In this area, human immunodeficiency virus prevalence was 1.2% among women 15–49 years of age and 0.76% among pregnant women [12]. At enrollment demographic data, history of illness, obstetric history, last menstrual period, age at menarche, and sexual activity were recorded. A study clinician performed a general clinical examination, duplicate anthropometric measurements, and axillary temperature. We collected venous blood for plasma ferritin, serum transferrin receptor (sTfR), hepcidin, and C-reactive protein (CRP) measurements. Plasma was stored at -80°C (additional laboratory details are shown in [Supplementary Data 1](#)). All participants received a long-lasting insecticidal net (LLIN) and single doses of albendazole (400 mg) and praziquantel.

Recruitment continued until the target number was reached. We anticipated 34% malaria prevalence in controls at ANC1 [13]. Three hundred ninety women in the intervention and control groups would provide 90% power to detect a noninferiority margin of 10% of malaria prevalence between trial arms.

The number of nonpregnant nulliparous women aged 15–24 years necessary to reach the required number of pregnant women was estimated from the proportion of nulliparous women (0.5) of childbearing age (0.23), aged 15–24 years (0.33), which provides 1973 potentially eligible women for a population of 52 000 in the DHSS area ([Figure 1](#)).

Randomization and Blinding

Participants were individually randomized to receive weekly one of identical red colored vegetable cellulose (hypromellose) capsules (disintegration time <30 minutes; mean, 9.5 minutes), containing either ferrous gluconate (60 mg elemental iron, 479 mg gluconate) and folic acid 2.8 mg, or folic acid alone. A block allocation sequence was used with randomly determined block lengths. Four containers (4 × 20 capsules) were assigned the same randomization code and were allocated to each participant. The FFA kept one container per participant, obtaining a replacement as required. Supplements were not kept by participants. Supplement codes, unknown to investigators and maintained independently by the sponsor, were revealed only after database lock and completion of data analysis. Women received a card with a unique study number corresponding to the randomization list. Supplements were stored at 20°C – 25°C and at ambient temperature while with FFAs.

Follow-up

FFAs provided supplements during weekly home visits and ingestion was directly observed. Participants not located for 2 consecutive attempts but then recontacted, were reported as temporarily absent. In cases of fever (temperature $\geq 37.5^{\circ}\text{C}$) or history of fever in the previous 48 hours, the FFA performed a malaria Rapid Diagnostic Test (RDT; Bioline SD, Malaria Antigen P.f) and if positive, collected a blood sample for a thick film. RDT positives were treated with artesunate-amodiaquine following national guidelines. If no menses were reported at 5 consecutive weekly visits, a urine pregnancy test was done for human chorionic gonadotropin, immediately or at the next weekly visit. Supplement safety and tolerability were evaluated by recording and grading adverse events (AEs). There were 2 levels of recording: weekly FFA symptom using a checklist, and AEs through passive follow-up at health facilities. Serious AEs (SAEs), including deaths, were collected by active (weekly) and passive surveillance and reported according to available information from FFAs and health center staff. SAE classification used all available clinical evidence and was not bound by stringent laboratory or diagnostic criteria. If a woman died outside hospital, verbal autopsies were done by the DHSS team following a standardized protocol.

At the end of the first malaria transmission season, finger-prick samples were taken from all nonpregnant participants for an interim safety analysis of malaria risk (microscopy and RDT). Positive RDTs were treated following national guidelines (additional details in procedures, [Supplementary Data 1](#)).

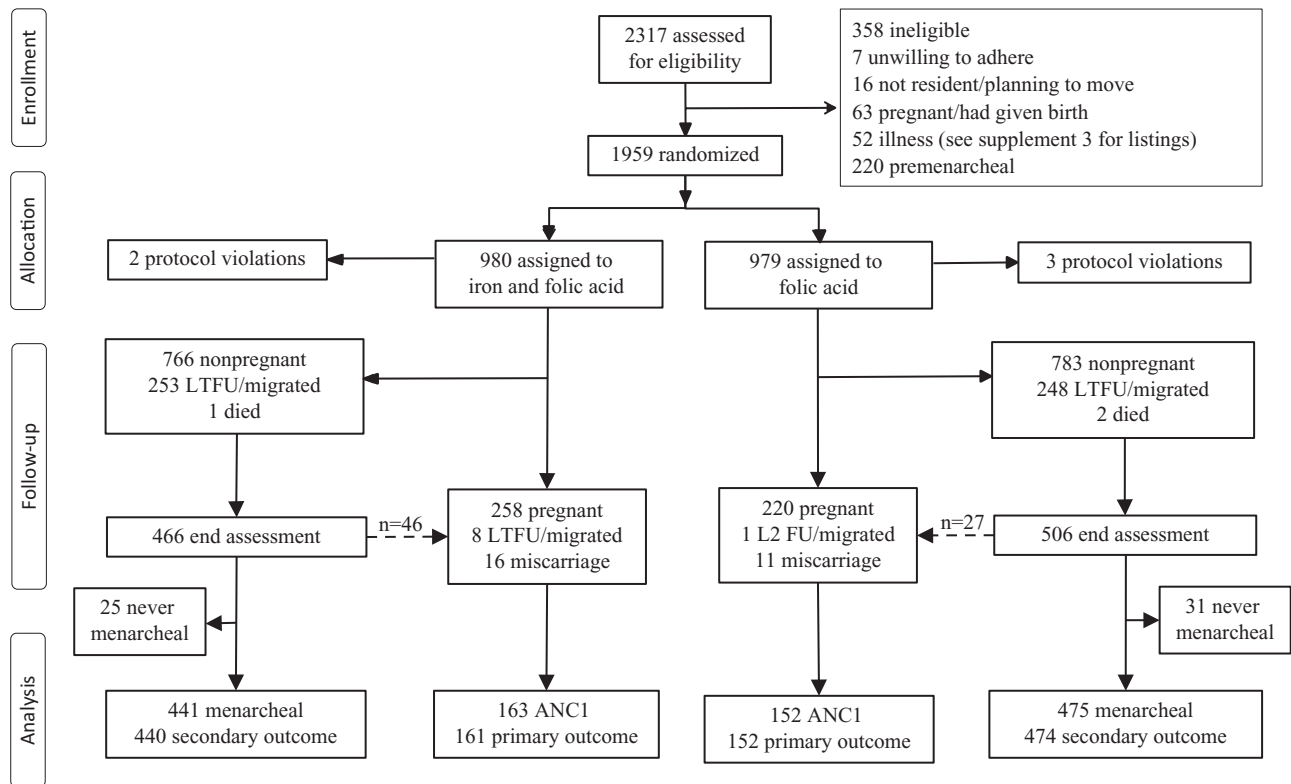


Figure 1. Trial profile. Stippled arrow indicates the number of women identified in early pregnancy at or within 2 months of the end assessment survey. Secondary outcome in nonpregnant women is malaria parasitemia. Abbreviations: ANC1, first antenatal visit; LTFU, lost to follow-up.

Weekly supplements continued for 18 months, when women who remained nonpregnant were referred for end assessment. This included medical history, clinical examination, anthropometry, and axillary temperature. A venous blood sample was collected for malaria smear, hemoglobin (Hemocue AB), plasma ferritin, TfR, hepcidin, and CRP measurements, and for storage on filter paper. If febrile (temperature $\geq 37.5^{\circ}\text{C}$) or experiencing fever in the previous 48 hours, an RDT was performed and women with positive results were treated. Urine pregnancy tests were repeated when indicated. On trial completion, anemic nonpregnant women (hemoglobin [Hb] level < 12 g/dL) received iron and folic acid tablets daily for 1 month and, if severely anemic (Hb < 8 g/dL), were referred to hospital.

Women becoming pregnant within the follow-up period were referred to Nanoro hospital for ANC1 with a study nurse/doctor at 13–16 weeks' gestation according to the last menstrual period. The weekly supplement was withdrawn and hematinics provided according to national policy (60 mg iron, 400 μg folic acid daily), although weekly follow-up continued. All women received the first dose of intermittent preventive treatment with sulfadoxine-pyrimethamine (IPTp-SP) if gestational age was > 13 weeks. Women ≤ 13 weeks' gestation, if RDT positive, were treated with oral quinine. At ANC1, procedures included routine antenatal care, venous blood collection for RDT, malaria film, hemoglobin, plasma ferritin, TfR, hepcidin, zinc

protoporphyrin, and CRP measurements. Ultrasound was performed to confirm pregnancy and to determine fetal location, number and viability, and gestational age. Following ANC1, women with moderate anemia (Hb 7.0–9.9 g/dL) received twice the recommended daily supplement during 1 month and, if severely anemic (Hb < 7 g/dL), were referred to Nanoro hospital. Women followed routine ANC visits where they received a second IPTp dose. Routine ANC and unscheduled health center visits were recorded on study questionnaires by study clinicians/nurses who collected data in health centers regularly (additional details in “Methods” in [Supplementary Data 1](#)). Procedures were similar at the second study visit (33–36 weeks' gestation) when women were encouraged to deliver at Nanoro Hospital or the nearest health center, where free obstetric care was provided by the study. Study nurses examined babies for congenital malformations within 24–48 hours of delivery and information on infant survival was obtained by FFAs who visited or telephoned families.

Laboratory Measurements

Blood samples were transported to the laboratory within 3 hours, centrifuged, and aliquoted. Blood films were Giemsa stained and read for malaria and parasite density (additional details on laboratory procedures are shown in [Supplementary Data 1](#)). Hemoglobin was measured (Sysmex automated

analyzer) on fresh whole blood, and zinc protoporphyrin by fluorometry (Aviv Biomedical). Plasma ferritin and sTfR were measured using mean values from duplicate enzyme-linked immunosorbent assay (ELISA) samples (Spectro Ferritin S-22 and TFC 94 TfR, RAMCO Inc). CRP was measured by ELISA (EU59131IBL, GmbH). Intra-assay coefficients of variation were all <10%. Plasma hepcidin was measured by competitive ELISA at an international reference laboratory (see laboratory procedures in [Supplementary Data 1](#)).

Outcomes

The primary outcome was *Plasmodium* parasitemia prevalence at ANC1. Prespecified secondary outcomes at ANC1 were prevalence of the following: RDT positivity with or without fever (temperature $\geq 37.5^{\circ}\text{C}$), clinical malaria (fever or history of fever in previous 48 hours with parasitemia), iron deficiency (definitions in [Supplementary Data 1](#)), anemia, adverse pregnancy outcomes (ie, miscarriage, stillbirth, perinatal or neonatal death), and congenital abnormality. In nonpregnant women at the end assessment, the proportion infected in the first year was assessed, along with the same prespecified outcomes as at ANC1. Adverse events captured incidence of prespecified gastrointestinal events (definitions and MedRA classification, grading and severity; see [Supplementary Data 1](#) and [2](#) and [Supplementary Table 3](#)). Malaria parasitemia prevalence was measured in nonpregnant women during the first rainy season after at least 6 months of supplementation. Adherence to supplementation is reported.

Statistical Analysis

A statistical analysis plan was approved before database lock and release of the data and analyses follow that plan with only minor variation. Analyses followed CONSORT (Consolidated Standards of Reporting Trials) guidelines for noninferiority trials [14]. Investigators were blinded to allocation (coded as A/B) until completion of the primary outcome analysis. Prevalence analyses based on data at specific time points utilized risk-ratio binomial models unadjusted and adjusted for season of transmission (low or high), LLIN use (proportion of weekly visits with reported use prior to visit date), antimalarial use in 4 weeks prior to visit, and whether menarcheal at baseline. Parasite density and iron biomarkers were analyzed using analogous ordinary regression models following logarithmic transformation. Results are expressed as risk ratios (or density/biomarker level ratios) for iron treatment with 95% confidence interval (CI). Analyses of outcomes by iron deficiency utilized the same methodology, but with no menarche covariate. Mean hepcidin levels were compared between malaria parasitemia-positive and -negative women at ANC1 and the end assessment using Mann-Whitney *U* tests. Differences in proportions with elevated hepcidin were estimated using Fisher exact test and based on the 95th percentile value for a healthy Dutch female population of comparable age [15]. Statistical significance was one-sided at $\alpha = .05$.

Malaria incidence was calculated using Poisson regression models for the number of malaria episodes per woman with an offset of the logarithm of the period under observation (ie, until final assessment, ANC1, or loss to follow-up) with adjustment for LLIN use and menarche. Results are presented as incidence ratios for iron treatment with 95% CI. Time to first episode of malaria was analyzed using Cox regression models with participants censored at ANC1, final assessment, or loss to follow-up with adjustment for season (baseline hazard stratified by enrollment month to allow for timing of rainy season), LLIN use, menarche, and iron deficiency at baseline (using adjusted ferritin definition). Results are expressed as hazard ratios for iron treatment with 95% CI. All analyses were performed using R statistical environment version 3.3 [16].

RESULTS

Of 2317 women invited for screening, 1959 were randomly assigned weekly supplements ($n = 980$, iron and folic acid; $n = 979$, folic acid); 1954 (99%) comprised the intention-to-treat population for the primary outcome. Four hundred five pregnancies (21%) occurred during follow-up, with another 73 (4%) early pregnancies identified at or shortly after the end assessment survey (giving 478 pregnancies in total) ([Figure 1](#)). Median weekly supplement adherence at ANC1 was 79% (95% CI, 65%–90%, iron) and 80% (95% CI, 59%–91%, folic acid) and at end assessment for nonpregnant women 83% (95% CI, 72%–91%, iron) and 84% (95% CI, 70%–92%, folic acid) ([Supplementary Table 1](#) and [Supplementary Data 2](#)). Mean plasma ferritin and sTfR/log ferritin ratios were not improved in women with weekly adherence $\geq 80\%$ ([Figure 2](#)). Median weekly LLIN use to ANC1 was 48.8%, and to end assessment was 46.7%.

Baseline

Characteristics were similar between groups ([Table 1](#)). Adolescents (<20 years) comprised 93%. Women who remained nonpregnant were younger with lower body mass index. Women not attending were more illiterate and sexually active ([Supplementary Table 2](#) and [Supplementary Data 2](#)). Iron deficiency prevalence at baseline did not differ in women lost to follow-up before ANC1 (9 of 478 [2%]), or for nonpregnant women lost to follow-up before end assessment (501 of 1549 [32%]) compared to those followed successfully. Iron deficiency prevalence was almost 2-fold higher using the sTfR/log ferritin ratio (22%) as biomarker than adjusted ferritin (12%) ([Table 1](#)). Elevated hepcidin concentration (>10.5 nmol/L) occurred in a quarter of women.

Prevalence of *P. falciparum* parasitemia in nonpregnant women during the first rainy season was 36.4% (95% CI, 33.7%–39.3%; $n = 1167$), with no difference by trial arm.

ANC1 Outcomes

Mean gestational age was 18.5 (standard deviation [SD], 5.5) weeks (iron and folic acid) and 18.0 (SD, 6.0) weeks

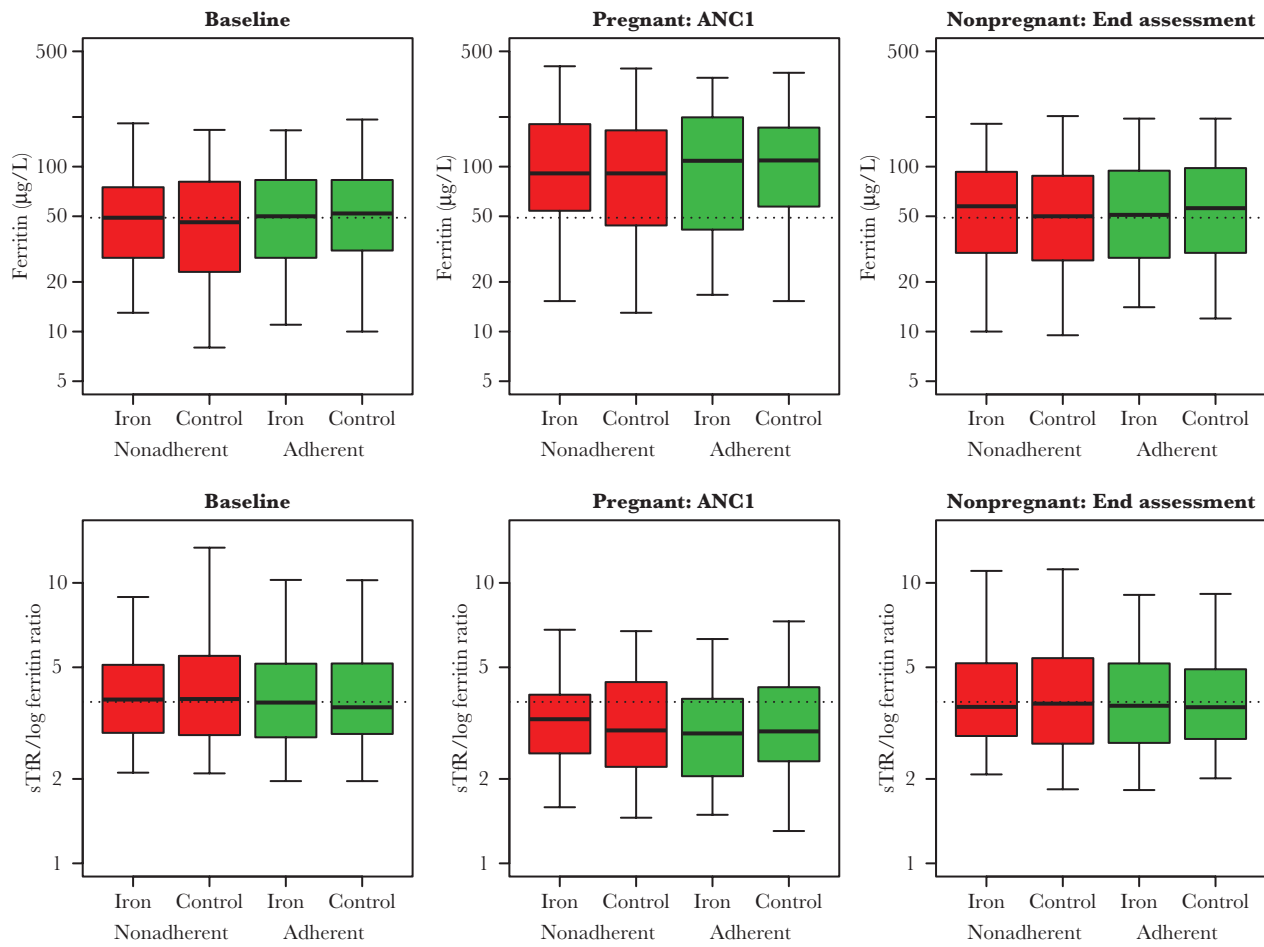


Figure 2. Mean ferritin concentration and serum transferrin receptor (sTfR)/log ferritin ratio in nonadherent and adherent women, by trial arm. Boxplots (median, interquartile range, and 90% range) for iron biomarkers in controls and iron-supplemented women at the 3 assessment points, subdivided by adherence to treatment. Adherence defined as receiving $\geq 80\%$ weekly supplement intake up to first antenatal visit (ANC1), or end assessment in nonpregnant women. Broken reference lines are median baseline levels.

(folic acid). *Plasmodium* parasitemia prevalence was 54.3%. This did not differ by trial arm (adjusted ratio, 1.0 [95% CI, .97–1.03]), nor when associated with fever. Incidence and time to the first symptomatic episode (fever plus parasitemia or RDT positivity) were not different between arms (Table 2).

There was no difference between study arms in anemia or iron deficiency prevalence, or mean concentrations of iron biomarkers (Table 3). Median hepcidin concentration was 3.9 nmol/L (interquartile range [IQR], 1.7–9.0) with, and 2.9 nmol/L (IQR, 0.7–7.0) without, malaria parasitemia ($P = .005$). Elevated hepcidin was more frequent in parasitemic (22%) compared to nonparasitemic women (11%) ($P = .015$).

End Assessment Outcomes

Plasmodium parasitemia prevalence was 49.0% and did not differ by trial arm (adjusted ratio, 1.1 [95% CI, .95–1.28]), or if associated with fever. There was no difference between study arms in prevalence of anemia or iron deficiency, or mean concentrations of iron biomarkers (Table 3). Median hepcidin concentrations in women with and without malaria parasitemia were

3.4 (95% CI 1.7–6.5) nmol/L and 2.9 (95% CI 1.1–6.2) nmol/L, respectively ($P = .011$). Elevated hepcidin was similar in parasitemic (15%) compared with nonparasitemic women (16%) ($P = .85$).

Adverse Events

Supplements were well tolerated but with more frequent gastrointestinal events with iron supplementation (Relative Risk, 1.29 [95% CI, .93–1.79]; $P = .12$). Adverse events for all categories did not differ by trial arm (Supplementary Table 3 and Supplementary Data 2). We recorded 106 adult and 81 infant SAEs with almost equal frequency between groups (Table 4). Six adult deaths occurred (3 at delivery), unrelated to the intervention or malaria. All congenital abnormalities (1.4%) occurred in controls. There were 17 infant/perinatal deaths, 12 to mothers who received iron.

DISCUSSION

We found that weekly iron and folic acid supplementation for up to 18 months did not affect significantly prevalence of

Table 1. Baseline Characteristics of the Intention-to-Treat Dataset

Characteristic	All Iron (n = 978)	All Control (n = 976)	Nonpregnant Iron (n = 766)	Nonpregnant Control (n = 783)	Pregnant Iron (n = 258)	Pregnant Control (n = 220)
Sociodemographics						
Mean age, y (SD)	16.8 (1.7)	16.8 (1.8)	16.7 (1.7)	16.7 (1.8)	17.1 (1.7)	17.1 (1.7)
Age <20 y	914/978 (93.5)	910/976 (93.2)	719/766 (93.9)	737/783 (94.1)	239/258 (92.6)	199/220 (90.5)
Ethnicity						
Mossi	936/978 (95.7)	946/975 (97.0)	730/766 (95.3)	757/782 (96.8)	250/258 (96.9)	214/220 (97.3)
Missing	0/978 (0)	1/976 (0.1)	0/766 (0)	1/783 (0.1)	0/258 (0)	0/220 (0)
Religion						
Catholic	414/978 (42.3)	405/975 (41.5)	317/766 (41.4)	327/782 (41.8)	118/258 (45.7)	86/219 (39.3)
Protestant	126/978 (12.9)	120/975 (12.3)	113/766 (14.8)	106/782 (13.6)	18/258 (7.0)	19/219 (8.7)
Muslim	251/978 (25.7)	250/975 (25.6)	187/766 (24.4)	203/782 (26)	75/258 (29.1)	52/219 (23.7)
Traditional	187/978 (19.1)	200/975 (20.5)	149/766 (19.4)	146/782 (18.7)	47/258 (18.2)	62/219 (28.3)
Missing	0/978 (0)	1/976 (0.1)	0/766 (0)	1/783 (0.1)	0/258 (0)	1/220 (0.5)
Education						
None	595/975 (61.0)	578/974 (59.3)	447/763 (58.6)	450/781 (57.6)	176/257 (68.5)	142/220 (64.5)
Primary	212/975 (21.7)	201/974 (20.6)	168/763 (22.0)	169/781 (21.6)	54/257 (21)	38/220 (17.3)
Secondary	168/975 (17.2)	195/974 (20.0)	148/763 (19.4)	162/781 (20.7)	27/257 (10.5)	40/220 (18.2)
Missing	3/978 (0.3)	2/976 (0.2)	3/766 (0.4)	2/783 (0.3)	1/258 (0.4)	0/220 (0)
Literacy						
Literate	329/966 (34.1)	347/966 (35.9)	278/756 (36.8)	297/774 (38.4)	68/255 (26.7)	62/218 (28.4)
Missing	12/978 (1.2)	10/976 (1.0)	10/766 (1.3)	9/783 (1.1)	3/258 (1.2)	2/220 (0.9)
Occupation ^a						
Student	300/978 (30.7)	317/976 (32.5)	255/766 (33.3)	274/783 (35)	60/258 (23.3)	51/220 (23.2)
Trading	32/978 (3.3)	31/976 (3.2)	24/766 (3.1)	21/783 (2.7)	10/258 (3.9)	10/220 (4.5)
Domestic	534/978 (54.6)	515/976 (52.8)	401/766 (52.3)	407/783 (52)	154/258 (59.7)	120/220 (54.5)
Farmer	375/978 (38.3)	380/976 (38.9)	269/766 (35.1)	290/783 (37)	130/258 (50.4)	101/220 (45.9)
Other	5/978 (0.5)	4/976 (0.4)	3/766 (0.4)	2/783 (0.3)	3/258 (1.2)	3/220 (1.4)
Clinical characteristics						
Menarcheal	844/978 (86.3)	829/976 (84.9)	639/766 (83.4)	645/783 (82.4)	241/258 (93.4)	203/220 (92.3)
Sexually active	249/978 (25.5)	241/975 (24.7)	159/766 (20.8)	164/782 (21.0)	98/258 (38)	82/220 (37.3)
Height, cm (SD)	159.0 (6.0)	159.2 (6.0)	158.6 (6.0)	159.1 (6.1)	160.0 (5.7)	159.4 (5.6)
Weight, kg (SD)	50.2 (6.8)	50.2 (7.2)	49.6 (6.9)	49.7 (7.2)	51.6 (6.0)	51.6 (6.7)
BMI, kg/m ² (SD)	19.8 (2.1)	19.7 (2.2)	19.7 (2.1)	19.6 (2.2)	20.1 (1.8)	20.3 (2.1)
BMI <18.5 kg/m ²	257/978 (26.3)	277/976 (28.4)	222/766 (29.0)	246/783 (31.4)	46/258 (17.8)	39/220 (17.7)
MUAC, cm (SD)	23.7 (2.1)	23.7 (2.2)	23.6 (2.1)	23.6 (2.2)	24.1 (1.8)	24.3 (2.2)
Serum iron biomarkers						
Median CRP, mg/L (IQR)	0.59 (0.23–1.47)	0.51 (0.20–1.35)	0.58 (0.23–1.43)	0.50 (0.18–1.29)	0.72 (0.27–1.64)	0.70 (0.22–1.66)
Missing	13/978 (1.3)	13/976 (1.3)	6/766 (0.8)	12/783 (1.5)	7/258 (2.7)	2/220 (0.9)
CRP ≥10 mg/L	38/965 (3.9)	41/963 (4.3)	30/760 (3.9)	31/771 (4.0)	9/251 (3.6)	10/218 (4.6)
Median ferritin, µg/L (IQR)	49.0 (28.0–78.0)	49.0 (26.0–82.0)	49.0 (28.0–79.8)	50.0 (27.0–81.0)	48.0 (28.0–74.5)	44.0 (26.0–81.0)
Missing	12/978 (1.2)	14/976 (1.4)	8/766 (1.0)	12/783 (1.5)	4/258 (1.6)	3/220 (1.4)
Median sTfR, mg/L (IQR)	6.32 (5.08–7.87)	6.28 (5.14–7.90)	6.32 (5.10–7.95)	6.26 (5.09–7.92)	6.20 (5.02–7.67)	6.34 (5.31–7.80)
Missing	11/978 (1.1)	18/976 (1.8)	7/766 (0.9)	16/783 (2.0)	4/258 (1.6)	3/220 (1.4)
Median sTfR/log ₁₀ ferritin ratio (IQR)	3.80 (2.88–5.13)	3.74 (2.88–5.36)	3.80 (2.88–5.13)	3.69 (2.86–5.25)	3.77 (2.88–5.13)	3.87 (2.94–5.61)
Missing	14/978 (1.4)	18/976 (1.8)	9/766 (1.2)	16/783 (2.0)	5/258 (1.9)	3/220 (1.4)
Median hepcidin, nmol/L (IQR)	4.80 (2.00–10.75)	4.30 (1.90–10.10)	4.80 (1.90–10.75)	4.40 (1.90–10.00)	4.90 (2.32–10.67)	4.10 (1.80–10.70)
Missing	11/978 (1.1)	16/976 (1.6)	7/766 (0.9)	14/783 (1.8)	4/258 (1.6)	3/220 (1.4)
Elevated hepcidin >10.5 nM/L ^b	251/967 (26.0)	222/960 (23.1)	196/759 (25.8)	172/769 (22.4)	65/254 (25.6)	55/217 (25.3)
Missing	11/978 (1.1)	16/976 (1.6)	7/766 (0.9)	14/783 (1.8)	4/258 (1.6)	3/220 (1.4)
Iron deficiency ^c (adjusted ferritin)	105/962 (10.9)	125/961 (13)	84/758 (11.1)	100/770 (13)	25/250 (10)	26/217 (12)
Missing	16/978 (1.6)	15/976 (1.5)	8/766 (1.0)	13/783 (1.7)	8/258 (3.1)	3/220 (1.4)
Iron deficiency ^d (sTfR/log ferritin ratio >5.6)	205/964 (21.3)	218/958 (22.8)	162/757 (21.4)	168/767 (21.9)	52/253 (20.6)	55/217 (25.3)
Missing	14/978 (1.4)	18/976 (1.8)	9/766 (1.2)	16/783 (2.0)	5/258 (1.9)	3/220 (1.4)

Data are presented as No. (%) unless otherwise indicated.

Abbreviations: BMI, body mass index; CRP, C-reactive protein; IQR, interquartile range; MUAC, mid-upper arm circumference; SD, standard deviation; sTfR, serum transferrin receptor.

^aMultiple answers possible for subsistence activities (ie, trading, farming, and domestic work).

^bHepcidin 95% range for women aged 18–24 years from reference Dutch population was median 2.6 nM; 2.5th percentile 0.7 nM; and 97.5th percentile 10.5 nM [15].

^cFerritin <15 µg/L if CRP <10 mg/L or ferritin <70 µg/L if CRP ≥10 mg/L.

^dRatio: sTfR (µg/L) to log₁₀ ferritin (µg/L) >5.6.

Table 2. Malaria at First Antenatal Visit and at End Assessment in Nonpregnant Nulliparae

Endpoint	Iron	Control	Ratio	P Value	Adjusted Ratio	Adjusted P Value ^a
Prevalence: ANC1						
No.	163	152				
RDT positive, No. (%)	95/161 (59.0) [51.3–66.3]	89/152 (58.6) [50.6–66.1]	1.01 [.84–1.21]	1.000	1.00 [.98–1.03]	.718
RDT positive and fever, No. (%)	15/163 (9.2) [5.7–14.6]	6/152 (3.9) [1.8–8.3]	2.33 [.93–5.85]	.072	2.30 [.91–5.79]	.075
Microscopy positive ^b , No. (%)	86/161 (53.4) [45.7–61.0]	84/152 (55.3) [47.3–62.9]	0.97 [.79–1.18]	.820	1.00 [.97–1.03]	.975
Clinical malaria ^c , No. (%)	9/161 (5.6) [3.0–10.3]	4/152 (2.6) [1.0–6.6]	2.12 [.67–6.75]	.259	2.14 [.68–6.76]	.194
Parasite density, parasites/mm ³ , GM	2000 [1483–2696]	2244 [1674–3008]	0.89 [.59–1.35]	0.585	0.89 [.58–1.36]	.584
Prevalence: end assessment ^a						
No.	441	475				
RDT positive & fever, No. (%)	19/432 (4.4) [2.8–6.8]	30/464 (6.5) [4.6–9.1]	0.68 [.39–1.19]	.188	0.69 [.39–1.21]	.193
Microscopy positive ^b , No. (%)	189/441 (42.9) [38.3–47.5]	186/474 (39.2) [34.9–43.7]	1.09 [.93–1.28]	.282	1.10 [.95–1.28]	.180
Clinical malaria ^c , No. (%)	13/440 (3.0) [1.7–5.0]	22/474 (4.6) [3.1–6.9]	0.64 [.32–1.25]	.228	0.67 [.34–1.31]	.237
Parasite density, parasites/mm ³ , GM	302 [248–368]	316 [259–386]	0.96 [.72–1.27]	.754	0.96 [.73–1.27]	.793
Incidence to end assessment in nonpregnant ^d						
Person-years of follow-up (No.)	1114.9 (936)	1121.1 (942)				
RDT positive (no./No.)	0.40 (445/1115)	0.40 (444/1121)	1.01 [.88–1.15]	.908	1.01 [.89–1.15]	.867
RDT positive and fever (no./No.)	0.16 (174/1115)	0.17 (188/1121)	0.93 [.76–1.14]	.494	0.94 [.76–1.15]	.525
Microscopy positive (no./No.)	0.25 (282/1115)	0.26 (286/1121)	0.99 [.84–1.17]	.919	1.00 [.85–1.18]	.971
Clinical malaria ^c (no./No.)	0.07 (79/1115)	0.08 (87/1121)	0.91 [.67–1.24]	.558	0.92 [.68–1.25]	.582
Proportion infected in first year ^e						
No. at risk	978	976				
No. censored at 1 y (%)	294 (30.1)	294 (30.1)				
RDT positive and fever, %	13 [11–15]	14 [12–16]	0.89 [.72–1.11]	.295	0.91 [.73–1.13]	.386
Clinical malaria ^c , %	6 [5–8]	7 [5–8]	0.92 [.67–1.25]	.588	0.92 [.68–1.26]	.617

Values in square brackets indicate the 95% confidence interval.

Abbreviations: ANC1, first antenatal visit; GM, geometric mean; RDT, rapid diagnostic test.

^aPrevalence adjusted for season at visit date, bed net use up to visit date, antimalarial use in month prior to visit, and menarcheal status at baseline. Ratio is risk ratio for iron treatment.

^bParasite positive on blood smear.

^c*Plasmodium falciparum* blood smear positive and fever $\geq 37.5^{\circ}\text{C}$.

^dIncidence per person-year adjusted for weekly bed net adherence and menarcheal status at baseline; ratio is incidence ratio.

^eCox regression model of time to first malaria episode, adjusted for baseline iron deficiency (ferritin), bed net use over observation time, visit, and menarcheal status at baseline, and stratified by enrollment month; ratio is hazard ratio.

Plasmodium infection, iron deficiency, or anemia, compared with folic acid supplements alone in either the pregnant or nonpregnant cohort. Women receiving iron experienced non-significantly increased gastrointestinal effects. Asymptomatic *falciparum* malaria infection was highly prevalent despite weekly active surveillance, access to free treatments, and treatment of all RDT positives at the interim safety survey.

Our trial was double blind, tablet consumption was directly observed with balanced adherence covering wet and dry seasons, and active surveillance identified malaria episodes. Inadequate supplementation is unlikely as women absent for a number of weeks were tracked and resumed supplements, allowing them to reestablish their iron levels. Untreated chronic asymptomatic malaria may lead to tissue pathology within the gut [17] as well as elevated hepcidin concentrations [18], which would be expected to reduce iron absorption. In relation to a safety trial, an effect of iron supplementation on malaria risk becomes difficult to establish if there is limited iron absorption. Baseline prevalence of iron deficiency was comparable to estimates in a recent meta-analysis for low-income countries [19]. In the only in vivo

iron absorption study to date in young women, conducted in Benin, markedly reduced iron absorption with asymptomatic malaria infection was demonstrated, and those participants had lower median parasite densities than we report at ANC1 [20]. Increasing iron dosage may be inadequate to overcome this limitation. Because participant diets also influence iron absorption due to high phytate content or other micronutrient deficiencies, effects of iron supplements may also depend on food habits and bioavailability. Better malaria control for adolescents could potentially both lower infection rates and improve food iron absorption, helping to meet growth requirements.

In 2016, WHO revised its guidelines on iron supplementation in menstruating women and adolescents girls, distinguishing settings where anemia prevalence is $\geq 40\%$ from $>20\%$. It recommended daily use (30–60 mg elemental iron) provided for 3 consecutive months per year where prevalence is $\geq 40\%$ [11], and weekly intermittent supplementation in 3 monthly annual cycles where anemia prevalence is 20%–40% [3]. Among nonpregnant controls in our study, anemia prevalence was 45% at the end assessment but $\leq 20\%$ were iron deficient. As we saw

Table 3. Anemia and Iron Biomarkers at First Antenatal Visit and at End Assessment in Nonpregnant Nulliparae

Endpoint	No.	Iron	Control ^a	Effect ^b	P Value	Adjusted Effect ^c	Adjusted P Value
ANC1							
Mean hemoglobin, g/dL	314	10.17 [9.96–10.38] (1 missing)	10.22 [9.99–10.46]	-0.06 [-.36 to .25]	.722	-0.01 [-.32 to .3]	.944
Anemia ^d	314	112/162 (69.1) [6.16–75.7]	107/152 (70.4) [62.7–77.1]	0.98 [85–114]	.902	0.96 [83–110]	.524
Severe anemia ^e	314	2/162 (1.2) [3–4.4]	4/152 (2.6) [1.0–6.6]	0.469 [0.87–2.524]	.435	0.443 [0.82–2.401]	.343
Iron deficiency (%) (adjusted ferritin) ^f	310	11/160 (6.9) [3.9–11.9]	19/150 (12.7) [8.3–18.9]	0.54 [2.7–1.10]	.123	0.53 [2.6–1.09]	.083
Iron deficiency (%) (sTfR/log ferritin ratio) ^g	312	18/162 (11.1%) [7.1–16.9]	19/150 (12.7%) [8.3–18.9]	0.88 [4.8–1.61]	.728	0.84 [4.6–1.54]	.578
Mean ferritin, µg/L	313	91 [78–107] (1 missing)	87 [74–103] (1 missing)	1.05 [84–131]	.694	1.07 [86–133]	.563
Mean sTfR, mg/L	313	6.0 [5.6–6.4] (1 missing)	5.9 [5.6–6.3] (1 missing)	1 [91–1.1]	.949	1 [91–1.1]	.962
Mean ZPP, µmol/mol heme	285	105 [100–111] (14 missing)	106 [101–112] (16 missing)	0.99 [92–1.07]	.841	0.99 [92–1.07]	.857
Mean hepcidin, nmo/L	311	3.01 [2.5–3.7] (3 missing)	2.86 [2.3–3.5] (1 missing)	1.05 [1.79–1.41]	.729	1.07 [80–1.43]	.664
Elevated hepcidin, >10.5 nM/L ^h	311	29/160 (18.1) [12.9–24.8]	25/151 (16.6) [11.5–23.3]	1.09 [6.7–1.78]	.766	1.12 [69–1.83]	.649
Nonpregnant							
Mean hemoglobin, g/dL	913	12.02 [11.91–12.12] (1 missing)	11.93 [11.82–12.04] (2 missing)	0.09 [-.07 to .24]	.261	0.06 [-.09 to .22]	.420
Anemia ^d	913	179/440 (40.7) [36.2–45.3]	217/473 (45.9) [41.4–50.4]	0.89 [76–103]	.124	0.90 [78–105]	.168
Severe anemia ^e	913	0/440 (0.0) [0.0–0.9]	3/473 (0.6) [0.2–1.8]	NA	.250	NA	.996
Iron deficiency (%) (adjusted ferritin) ^f	910	38/439 (8.7) [6.4–11.7]	50/471 (10.6) [8.1–13.7]	0.82 [55–122]	.369	0.82 [55–123]	.336
Iron deficiency (sTfR/log ferritin ratio) ^g	909	89/437 (20.4) [16.9–24.4]	97/472 (20.6) [17.2–24.4]	0.99 [77–1.28]	1.000	0.99 [77–1.28]	.960
Mean ferritin, µg/L	912	49 [46–53] (2 missing)	51 [47–55] (2 missing)	0.97 [87–1.09]	.632	0.97 [86–1.08]	.564
Mean sTfR, mg/L	909	6.4 [6.1–6.6] (4 missing)	6.5 [6.2–6.7] (3 missing)	0.98 [94–1.04]	.558	0.98 [93–1.04]	.534
Mean ZPP, µmol/mol heme	914	101 [97–104] (1 missing)	101 [98–105] (1 missing)	0.99 [94–1.04]	.783	0.99 [94–1.04]	.784
Mean hepcidin, nmo/L	909	2.85 [2.5–3.2] (3 missing)	2.83 [2.5–3.2] (4 missing)	1.01 [86–1.17]	.939	1.0 [86–1.17]	.960
Elevated hepcidin, >10.5 nM/L ^h	909	70/438 (16.0) [12.8–19.7]	72/471 (15.3) [12.3–18.8]	1.05 [77–1.41]	.785	1.05 [78–1.43]	.736

Values in square brackets indicate the 95% confidence interval (CI).

Abbreviations: ANC1, first antenatal visit; NA, not applicable; sTfR, serum transferrin receptor; ZPP, zinc protoporphyrin.

^aData are presented as No. (%) for binary variables, mean [95% CI] for hemoglobin; geometric mean [95% CI] for iron biomarkers.

^bRisk ratio for binary outcomes; difference between arms for hemoglobin; ratio between arms for iron biomarker levels.

^cAnemia measures adjusted for baseline menarche, season at assessment, and use of antimalarials in the previous month. Iron measures adjusted for baseline menarche.

^dHemoglobin <11 g/dL.

^eHemoglobin <8 g/dL.

^fFerritin <15 µg/L if C-reactive protein (CRP) <10 mg/L or ferritin <70 µg/L if CRP ≥10 mg/L.

^gRatio: sTfR µg/L to log₁₀ ferritin >5.6.

^hHepcidin 95% reference range for women aged 18–24 years from reference Dutch population was median 2.6 nM; 2.5th percentile 0.7 nM; and 97.5th percentile 10.5 nM [15].

ⁱHemoglobin <12 g/dL.

^jHemoglobin <7 g/dL.

Table 4. Serious Adverse Events

Adverse Event	Iron	Control
Adult SAE	(n = 978)	(n = 976)
Accidental death ^a	0	1
Obstetric death	2	1
Other obstetric SAE ^b	12	7
Severe malaria	27	20
Other death ^c	1	1
Other ^d	12	22
Total SAE	54	52
Infant SAE	(n = 231)	(n = 206)
Miscarriage	16	11
Stillbirth	13	17
Perinatal death*	6	0
Neonatal death	4	3
Infant death	1	1
Death from congenital abnormality ^e	0	1
Other congenital abnormalities ^{f, **}	0	5
Other obstetric SAE ^b	2	0
Other death ^g	1	0
Total SAE	43	38

Abbreviation: SAE, serious adverse event.

^aOne death from drowning.

^bSystems Order Classification = Pregnancy, puerperium and perinatal conditions, excluding deaths.

^cNeoplasms: benign, malignant, and unspecified (liposarcoma and thoracic pain).

^dSee Supplementary Data for summary of adverse events.

^eSpina bifida.

^fClub foot; congenital anomaly; polydactyly (all classified as unlikely or definitely not treatment related). Excludes abnormalities associated with stillbirth/miscarriage (one microcephaly).

^gParalytic ileus.

* $P = .021$ (Fisher exact test).

** $P = .011$ for all abnormalities.

no improved iron status at the higher dosage of 60 mg with up to 18 months of supplementation (maximum of 78 doses), it is questionable whether daily dosage, given over 3 months, would be more beneficial (maximum 90 doses). This population has low iron deficiency prevalence despite high anemia prevalence. Without evidence that iron supplementation is effective in young nulliparous women living in malaria-endemic areas, the basis for the WHO recommendation is weak and possibly unsafe in this group.

Adverse events were evenly balanced across trial arms, including miscarriages, which could result from early pregnancy malaria. Gastrointestinal adverse events were nonsignificantly increased in iron-supplemented women. We have showed a 2-fold increased use of antibiotics for treatment of gastrointestinal infections in these women, with increased use of antifungals for lower genital infection in the nonpregnant iron-supplemented cohort [21]. Iron supplementation is reported to increase diarrhea frequency in menstruating women from non-malaria-endemic areas [2].

Our study has some limitations. Nonattendance was greater than expected, with 25% of nonpregnant women not attending

end assessment and 30% of pregnant women at ANC1, although contact with their families was retained to monitor AEs up to birth. Attrition was equivalent between trial arms. Domestic labor, early sexual activity, pregnancy before marriage, and migration outside the study area at marriage were contributory factors in failure to attend [22], and would be higher than in a nonadolescent population. Six percent of women were identified as nonmenarcheal at end assessment, possibly related to intentional misreporting to gain free treatment [23]. Bias due to different premenarcheal iron requirements would be unlikely as there were equal numbers across study arms, but premenarcheal status reduced the conception rate.

CONCLUSIONS

In a high-malaria-transmission area, weekly iron supplementation did not increase malaria risk, improve iron status, or reduce anemia in young, mostly adolescent menstruating women, nor in early pregnancy. Iron absorption studies are required to clarify whether chronic malaria influences iron uptake. Baseline characteristics were typical for adolescents in low-income, rural sub-Saharan Africa; thus, our results should apply to similar malaria-endemic areas with high rates of asymptomatic adolescent malaria. Studies are warranted to improve malaria prevention and control in adolescent populations and to shape relevant interfacing iron deficiency reduction strategies. Iron supplementation, as routinely given to populations such as this, is not helpful and is potentially harmful. WHO guidelines for universal supplementation in young nulliparous women may need to be reassessed.

Supplementary Data

Supplementary materials are available at *The Journal of Infectious Diseases* online. Consisting of data provided by the authors to benefit the reader, the posted materials are not copyedited and are the sole responsibility of the authors, so questions or comments should be addressed to the corresponding author.

Notes

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