











Prevalence and risk factors of curable sexually transmitted and reproductive tract infections and malaria co-infection among pregnant women at antenatal care booking in Kenya, Malawi and Tanzania: a cross-sectional study of randomised controlled trial data

Georgia R Gore-Langton ¹, Mwayiwawo Madanitsa ^{2,3}, Hellen C Barsosio ⁴, Daniel T R Minja,⁵ Jacklin Moshia,⁶ Reginald A Kavishe,⁶ George Mtove,⁵ Samwel Gesase,⁵ Omari A Msemu,⁵ Simon Kariuki,⁴ Kephias Otieno,⁴ Kamija S Phiri,⁷ John P A Lusingu,⁵ Crispin Mukerebe ⁶, Alphaxard Manjurano,⁶ Pius Ikigo,⁶ Queen Saidi,⁷ Eric D Onyango,⁴ Christentze Schmiegelow,^{8,9} James Dodd,¹⁰ Jenny Hill ¹⁰, Helle Hansson,⁸ Michael Alifrangis,⁸ Julie Gutman ¹¹, Patricia Jean Hunter ¹², Nigel Klein,¹² Ulla Ashorn ¹³, Asma Khalil,¹⁴ Matt Cairns ¹⁵, Feiko O ter Kuile,^{4,10} R Matthew Chico ¹

To cite: Gore-Langton GR, Madanitsa M, Barsosio HC, *et al*. Prevalence and risk factors of curable sexually transmitted and reproductive tract infections and malaria co-infection among pregnant women at antenatal care booking in Kenya, Malawi and Tanzania: a cross-sectional study of randomised controlled trial data. *BMJ Public Health* 2024;**2**:e000501. doi:10.1136/bmjph-2023-000501

► Additional supplemental material is published online only. To view, please visit the journal online (<https://doi.org/10.1136/bmjph-2023-000501>).

OAM died in December 2022
NK died in June 2024

Received 17 August 2023
Accepted 30 July 2024



© Author(s) (or their employer(s)) 2024. Re-use permitted under CC BY. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Dr Georgia R Gore-Langton; georgia.gore-langton@lshtm.ac.uk

ABSTRACT

Objectives Malaria and curable sexually transmitted and reproductive tract infections (STIs/RTIs) are associated with adverse pregnancy outcomes. This study reports the prevalence and risk factors of curable STIs/RTIs, STI/RTI co-infection and STI/RTI and malaria co-infection among HIV-negative pregnant women at their first antenatal care visit in Kenya, Malawi and Tanzania.

Methods HIV-negative pregnant women of all gravidae (n=4680) were screened for syphilis with point-of-care tests and treated if positive. Separately, women provided blood samples (n=4569) for rapid plasma reagin (RPR) testing; positive cases were confirmed by *Treponema pallidum* particle agglutination (TPPA). Women also provided dried blood spots for batch testing of malaria by retrospective polymerase chain reaction (PCR (n=4226) methods. A randomly selected subgroup of women provided vaginal swabs for chlamydia, gonorrhoea and trichomoniasis testing by retrospective PCR batch testing (n=1431), and bacterial vaginosis diagnosis by Nugent scoring (n=1402).

Results Malaria prevalence was 14.6% (95% CI 13.6 to 15.7), 45.9% (43.4 to 48.4) of women were positive for at least one curable STI/RTI and 6.7% (5.5 to 8.1) were co-infected with malaria and a curable STI/RTI. Prevalence of individual STIs/RTIs ranged from 28.5% (26.2 to 30.9) for bacterial vaginosis to 14.5% (12.7 to 16.4) for trichomoniasis, 13.8% (12.1 to 15.7) for chlamydia, 2.7% (1.9 to 3.6) for gonorrhoea and 1.7% (1.4 to 2.2) for RPR/TPPA-confirmed syphilis. The prevalence of STI/RTI co-infection was 10.1% (8.7 to 11.8). Paucigravidae, at highest risk of malaria, were also at greater

WHAT IS ALREADY KNOWN ON THIS TOPIC

- ⇒ Malaria and curable sexually transmitted and reproductive tract infections (STIs/RTIs) during pregnancy are associated with adverse pregnancy outcomes, with the majority of cases asymptomatic.
- ⇒ In areas of moderate-to-high malaria transmission, intermittent preventive treatment with sulfadoxine-pyrimethamine is provided to pregnant women and universal screening and treatment for syphilis is standard; however, other curable STIs/RTIs are most often undetected and untreated.

WHAT THIS STUDY ADDS

- ⇒ This study adds critical information on the prevalence of malaria and STIs/RTIs in three sub-Saharan African countries and highlights the frequency of co-infection.

risk of having chlamydia, gonorrhoea and bacterial vaginosis than multigravidae.

Conclusions Of women infected with malaria, 49.0% also had a curable STI/RTI and one in five women with at least one STI/RTI were co-infected with more than one STI/RTI. Current antenatal interventions that address malaria and curable STIs/RTIs remain suboptimal. New approaches to preventing and managing these infections in pregnancy are urgently needed.

Trial registration number [NCT03208179](https://www.clinicaltrials.gov/ct2/show/study/NCT03208179).

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

- ⇒ Apart from syphilis, the antenatal care visit package lacks interventions that effectively target malaria and curable STIs/RTIs.
- ⇒ In the near-term, this may involve the use of sulfadoxine-pyrimethamine combined with a partner antimalarial therapy such as dihydroartemisinin-piperazine.
- ⇒ Pregnant women also need access to affordable, rapid and accurate point-of-care diagnostics for curable STI/RTIs early in pregnancy and vaccines are needed to protect against these infections in pregnancy.

BACKGROUND

Intermittent preventive treatment in pregnancy (IPTp) with sulfadoxine-pyrimethamine (SP) is one of the interventions recommended by WHO for the prevention of adverse consequences attributable to malaria during pregnancy in areas of moderate to high transmission. Malaria during pregnancy is associated with increased risk of numerous adverse fetal, newborn (including spontaneous abortion, stillbirth, preterm birth, low birth weight, small-for-gestational age) and maternal outcomes (maternal anaemia and death).¹⁻³ In some countries, it is policy to test all women for malaria with rapid diagnostic tests (RDT) at antenatal care (ANC) booking, while in other countries relying on the presentation of symptoms is national policy. Sexually transmitted and reproductive tract infections (STIs/RTIs) are associated with ectopic pregnancies, preterm birth, small-for-gestational age, low birth weight, neonatal death as well as risk of acquiring and transmitting HIV.⁴⁻⁶ The five most common STIs/RTIs are curable: syphilis (*Treponema pallidum*), chlamydia (*Chlamydia trachomatis*), gonorrhoea (*Neisseria gonorrhoea*), trichomoniasis (*Trichomonas vaginalis*) and bacterial vaginosis. Syphilis is particularly pernicious, trebling the risk of stillbirth⁷ and causing 1 in 10 stillbirths in sub-Saharan Africa.⁸ WHO recommends universal syphilis screening at ANC booking and treatment of cases with benzathine penicillin G.⁹ For the diagnosis and treatment of other curable STIs/RTIs, WHO recommends symptom-based management in low-resource settings where laboratory facilities and trained staff may be limited.^{10 11} Syndromic management has proven effective among men who are most often symptomatic; however, women are far more likely to be asymptomatic and, therefore, less likely to be diagnosed and treated.¹²⁻¹⁶

In Burkina Faso, the prevalence of malaria and curable STI/RTI co-infection at ANC has been reported at 12.9%. This was based on malaria diagnosis by microscopy, point-of-care tests for syphilis and chlamydia and Nugent scoring for bacterial vaginosis.¹⁵ A study in Zambia using rapid plasma reagin (RPR) with *T. pallidum* haemagglutination assay to confirm syphilis, polymerase chain reaction (PCR) to diagnose malaria, chlamydia, gonorrhoea and trichomoniasis, and Nugent scoring for bacterial vaginosis, reported the prevalence of malaria and curable

STI/RTI co-infection as 38.7%¹⁷; >70% of the STI/RTI cases were asymptomatic.¹³

Information on curable STIs/RTIs during pregnancy is scarce in sub-Saharan Africa given the asymptomatic nature of most infections and the absence of screening for many infections at ANC, apart from HIV and syphilis. Using gold standard diagnostics in three East African countries, we investigated curable STI/RTI prevalence and risk factors, including malaria and curable STI/RTI co-infection with the aim of documenting a more complete estimate of disease burden during pregnancy and informing the development of integrated packages of care to address them and reduce their associated adverse consequences for maternal and newborn health.

METHODS

Study design, population and study site

Participants were enrolled in the IMPROVE (IMproving PRegnancy Outcomes with intermittent preVEntative treatment in Africa) trial (NCT03208179), a three-arm malaria chemoprevention randomised controlled trial conducted in areas of perennial malaria transmission in Kenya, Malawi and Tanzania between 29 March 2018 and 5 July 2019; main trial outcomes have been published elsewhere.¹⁸ Eligible women were between 16 and 28 gestational weeks as measured by ultrasound, HIV-negative and had not previously received SP during the current pregnancy. Recruitment stopped when 4680 participants had been randomly assigned, as per sample size calculations detailed in appendix 8 of the main trial paper.¹⁸ Participants were allocated to receive at each scheduled ANC visit IPTp with SP (n=1561), or dihydroartemisinin-piperazine (n=1561) or dihydroartemisinin-piperazine plus the addition of 2 g azithromycin as part of the first visit only (n=1558) (online supplemental table 1). IPTp-SP is a single-day course and was directly observed in all instances. In contrast, dihydroartemisinin-piperazine is a 3-day course and azithromycin was a 2-day course. Women in these two groups had IPTp directly observed, but took the remaining tablets home for self-administration. Self-reported adherence was high, exceeding 97.5%, a value validated through random and unannounced home visits over the course of the trial, along with telephone calls to all participants on the second and third days of each course.

Malaria and STI/RTI diagnosis

At enrolment, participants provided sociodemographic and maternal health information. Clinical staff collected a finger-prick blood sample from participants for retrospective batch analysis of malaria by ultrasensitive quantitative real-time-PCR.^{19 20} In Kenya and Tanzania, in keeping with national policy, all women were screened for malaria by RDT at enrolment and given first-line malaria treatment if positive. All women were screened for syphilis with a rapid point-of-care test (SD Bioline

V.3.0). Positive cases were treated with 2.4 million units of benzathine penicillin G by intramuscular injection. As part of routine care, clinic staff queried women about whether they had experienced any symptoms associated with STIs/RTIs. Any suspected cases were assessed and treated by the clinic staff based on syndromic management guidelines.

Apart from routine care, all women provided venous blood for RPR (Plasmatec RPR Test Kit) analysis and syphilis confirmation by TPPA (Serodia). Additionally, clinic staff collected cervicovaginal swabs from a randomly selected subset of one-third of participants at each study site. Samples were shipped to a regional reference laboratory at the National Institute for Medical Research, Mwanza, Tanzania, at the end of the trial for retrospective batch analysis and, therefore, the test results were not used for clinical management. Samples were tested for *C. trachomatis* and *N. gonorrhoea* by RT-PCR (Artus CT/NG QS-RGQ Kit), trichomoniasis with SACASETTM Real-TM Kit and bacterial vaginosis using the Nugent scoring (a case defined as a Nugent score of 7–10). Hereafter, when referring to curable STIs/RTIs, we specifically mean: syphilis, chlamydia, gonorrhoea, trichomoniasis and bacterial vaginosis.

Data management and statistical analyses

Binary positive/negative outcome variables were generated for the five STIs/RTIs and malaria. When calculating the prevalence of at least one curable STI/RTI, analysis was restricted to the subset of participants who provided vaginal swabs to diagnose curable STIs/RTIs. We tested associations between maternal age (dichotomised around the median age; <25 years vs ≥25 years) and being positive for each individual STI/RTI using modified Poisson regression models to obtain prevalence ratios (PRs). We pooled data across the three countries, controlling for country and study site. Each potential explanatory variable: gravidity (paucigravidae (primigravidae and secundigravidae combined), multigravidae), maternal education (none/primary, secondary), marital status (single, married), residency (urban, rural), socio-economic status (SES) (low, medium, high tercile) and maternal anaemia (<11 g/dL), was added one-by-one to the model including maternal age, country and site. Being positive for bacterial vaginosis was also investigated as a potential risk factor for each of the four STIs. An explanatory variable which changed the PR of the STI/RTI by maternal age, country and site by >10% was included in the final multivariable model for that specific STI/RTI. Analyses were done in Stata (V.16, College Station, Texas, USA) and R (V.4.03).

We generated prevalence estimates of co-infection (positive for more than one STI/RTI or for malaria and at least one STI/RTI) and summarised co-infection combinations. Risk factors for the most common STI/RTI co-infection combinations were tested using multinomial logistic regression models to estimate the effect of explanatory variables on the probability that the outcome

is in a particular category. For these models, the outcome measure was a mutually exclusive, multicategorical infection status variable, for example: (1) negative for bacterial vaginosis and trichomoniasis, (2) mono-infection with bacterial vaginosis, (3) mono-infection with trichomoniasis or (4) co-infection with bacterial vaginosis and trichomoniasis.

We present results first for individual infections: prevalence of malaria and then each STI/RTI, followed by risk factors for each STI/RTI. Prevalence of STI/RTI co-infection combinations is then presented followed by risk factors for the most common STI/RTI co-infection combinations. Finally, malaria and curable STI/RTI co-infection prevalence and risk factors are presented.

RESULTS

Study population

In total, 4680 women were recruited, of whom 97.6% (n=4569) were tested for syphilis (RPR and TPPA) and 90.3% (n=4226) were tested for malaria by PCR. Randomly selected participants, 30.6% (n=1431), provided samples for gonorrhoea, chlamydia and trichomoniasis testing by PCR, and 30.0% (n=1402) for bacterial vaginosis testing by Nugent score. In total, 31.8% (n=1488) of women were tested for at least one of chlamydia, gonorrhoea, trichomoniasis or bacterial vaginosis. Maternal demographics did not differ by STI/RTI tested (table 1). See online supplemental figure 1 and online supplemental table 2 for further details on the number and combination of curable STIs/RTIs tested for.

Prevalence: malaria and individual STIs/RTIs

Prevalence of malaria by PCR was 14.6% (95% CI 13.6 to 15.7) at enrolment. In total 469 women, 10.3% (9.4 to 11.2), were RPR positive, 79 of whom were confirmed by TPPA, giving a syphilis prevalence of 1.7% (1.4 to 2.2) (table 2) (see online supplemental table 3 for syphilis titre information). Among the subgroup tested, the prevalence of chlamydia was 13.8% (12.1 to 15.7), gonorrhoea 2.7% (1.9 to 3.6) and trichomoniasis 14.5% (12.7 to 16.4) (table 2, online supplemental figure 2). In Malawi, the prevalence of malaria (19.3% (17.2 to 21.5)) and syphilis (3.2% (2.4 to 4.3)) was higher than in Tanzania and Kenya. In Kenya, the prevalence of chlamydia was particularly high, at 21.0% (17.6 to 24.8) relative to Tanzania, 11.0% (8.7 to 13.8) and Malawi, 8.7% (6.2 to 12.2). In each of the three countries, bacterial vaginosis was the most common STI/RTI, with an overall prevalence of 28.5% (26.2 to 30.9). Among women with bacterial vaginosis, the mean Nugent score was 8.1; a further 9.8% (137/1402) of women had intermediary flora (Nugent score 4–6) (online supplemental table 3), among these women, 79.6% (109/137) were positive for clue cells. Among the subgroup tested, nearly one-half (45.9% (43.4 to 48.4); 683/1488) were positive for at least one STI/RTI.

Table 1 Sociodemographic and health characteristics of participants tested for malaria and STIs/RTIs

Sociodemographic and health characteristics*		Number of participants tested by infection (malaria, STIs/RTIs)			
		Malaria N=4226 (90.3%)	Syphilis N=4569 (97.6%)	Chlamydia, gonorrhoea, trichomoniasis N=1431 (30.6%)	Bacterial vaginosis N=1402 (30.0%)
		N (%)	N (%)	N (%)	N (%)
Maternal age (years)	<25 years	2274 (53.8)	2432 (53.2)	763 (53.4)	744 (53.1)
	≥25 years	1950 (46.2)	2137 (46.8)	668 (46.7)	657 (46.9)
Gravidity	Paucigravidae	2345 (55.7)	2523 (55.4)	783 (55.0)	774 (55.4)
	Multigravidae	1868 (44.3)	2033 (44.6)	642 (45.0)	623 (44.6)
Marital status	Single	507 (12.0)	566 (12.4)	176 (12.3)	174 (12.4)
	Married	3715 (88.0)	4001 (87.6)	1254 (87.7)	1227 (87.6)
Socio-economic status by tercile	Low	1434 (34.0)	1516 (33.2)	466 (32.6)	440 (31.4)
	Medium	1417 (33.6)	1525 (33.4)	485 (33.9)	474 (33.8)
	High	1373 (32.5)	1528 (33.4)	480 (33.5)	488 (34.8)
Maternal education	None/Primary	2739 (65.0)	2963 (65.0)	900 (63.0)	869 (62.1)
	Secondary	1477 (35.0)	1598 (35.0)	528 (37.0)	530 (37.9)
Residency	Rural	3155 (74.7)	3351 (73.4)	1045 (73.1)	1026 (73.2)
	Semi-urban/ Urban	1067 (25.3)	1216 (26.6)	385 (26.9)	375 (26.8)
Anaemia	No	2327 (55.3)	2535 (55.7)	777 (54.4)	766 (54.8)
	Yes	1882 (44.7)	2017 (44.3)	651 (45.6)	633 (45.3)

*The number of missing values was <5 for all sociodemographic and health characteristics. RTI, reproductive tract infection; STI, sexually transmitted infection.

Risk factors associated with individual STI/RTIs

Results of the final adjusted models with each individual STI/RTI as the outcome of interest, and the variables listed in the 'Methods' section explored as risk factors are shown in [table 3](#) for syphilis, chlamydia, gonorrhoea and bacterial vaginosis. In the final syphilis multivariable model, being positive for bacterial vaginosis was associated with an increased risk of infection; the prevalence of syphilis was 3.6% among women with bacterial vaginosis and 0.9% among women without bacterial vaginosis (PR: 3.7; 95% CI 1.6 to 8.4), $p=0.002$ ([table 3](#), online supplemental table 4). Women of low or medium SES were almost nine times more likely to have syphilis than women of high SES ([table 3](#)). Neither gravidity nor maternal age were associated with syphilis. Prevalence was slightly higher among multigravidae, 2.3% than paucigravidae, 1.3%.

Compared with multigravidae, paucigravidae were more likely to have chlamydia, gonorrhoea and bacterial vaginosis. When controlling for maternal age, chlamydia prevalence was 70% higher (PR 1.7; 95% CI 1.2 to 2.4, $p=0.002$) among paucigravidae than among multigravidae, and bacterial vaginosis was 30% higher (PR 1.3; 95% CI 1.1 to 1.6, $p=0.018$) ([table 3](#), online supplemental tables 5 and 6). When controlling for maternal age and education level, the prevalence of gonorrhoea among paucigravidae women was more than double

that of multigravidae (PR 2.5; 95% CI 1.2 to 5.4, $p=0.018$) ([table 3](#), online supplemental table 7).

There were no subgroups identified as being at elevated risk of having trichomoniasis in either the univariate or the multivariate analyses (online supplemental table 8). While maternal age and gravidity were risk factors in univariate analyses for the composite outcome of at least one of chlamydia, gonorrhoea, trichomoniasis or bacterial vaginosis, these were not associated in the final multivariable model (online supplemental table 9).

Prevalence: STI/RTI co-infection

Among the 678 women positive for an STI/RTI and tested for additional STIs/RTIs, 22.0% ($n=149$) were co-infected with at least one other STI/RTI, giving an overall co-infection prevalence of 10.1% (95% CI 8.7 to 11.8) among all women tested for at least two STIs/RTIs. Of co-infected women, 80.5% ($n=120$) had two STIs/RTIs, 16.8% ($n=25$) had three STIs/RTIs, 2.0% ($n=3$) had four STIs/RTIs and one woman (0.7%) was co-infected with all five curable STIs/RTIs.

The most common unique co-infection combination was bacterial vaginosis and chlamydia ($n=51$) (a further 22 women were co-infected with these two and additional STIs/RTIs), followed by bacterial vaginosis and trichomoniasis ($n=31$) (a further 21 women were co-infected with these two plus additional STIs/RTIs), 18 women

Table 2 Overall prevalence of malaria and STIs/RTIs at first ANC visit

	Total		Malawi		Tanzania		Kenya	
	Positive/Tested	Prevalence % (95% CI)	Positive/Tested	Prevalence % (95% CI)	Positive/Tested	Prevalence % (95% CI)	Positive/Tested	Prevalence % (95% CI)
Malaria	618/4226	14.6 (13.6 to 15.7)	250/1298	19.3 (17.2 to 21.5)	179/1628	11.0 (9.6 to 12.6)	189/1298	14.6 (12.7 to 16.6)
Syphilis	79/4569	1.7 (1.4 to 2.2)	42/1321	3.2 (2.4 to 4.3)	30/1772	1.7 (1.2 to 2.4)	7/1476	0.5 (0.2 to 1.0)
Chlamydia	198/1431	13.8 (12.1 to 15.7)	31/355	8.7 (6.2 to 12.2)	65/590	11.0 (8.7 to 13.8)	102/486	21.0 (17.6 to 24.8)
Gonorrhoea	38/1431	2.7 (1.9 to 3.6)	11/355	3.1 (1.7 to 5.5)	11/590	1.9 (1.0 to 3.3)	16/486	3.3 (2.0 to 5.3)
Trichomoniasis	207/1431	14.5 (12.7 to 16.4)	60/355	16.9 (13.4 to 21.2)	92/590	15.6 (12.9 to 18.8)	55/486	11.3 (8.8 to 14.5)
Bacterial vaginosis	399/1402	28.5 (26.2 to 30.9)	102/327	31.2 (26.4 to 36.4)	171/586	29.2 (25.6 to 33.0)	126/489	25.8 (22.1 to 29.8)
≥1 STI/RTI	683/1488	45.9 (43.4 to 48.4)	163/391	41.7 (36.9 to 46.7)	283/593	47.7 (43.7 to 51.8)	237/504	47.0 (42.7 to 51.4)
≥1 STI/RTI and malaria	94/1401	6.7 (5.5 to 8.1)	33/378	8.7 (6.3 to 12.0)	30/543	5.5 (3.9 to 7.8)	31/480	6.5 (4.6 to 9.0)

Previous reporting of STI/RTI prevalence was restricted to a slightly smaller subgroup than presented here.¹⁸ These estimates account for all STI/RTI diagnostic results. RTI, reproductive tract infection; STI, sexually transmitted infection.

were co-infected with chlamydia and trichomoniasis (a further 16 women were co-infected with these two plus additional STIs/RTIs). The fourth most common co-infection combination was bacterial vaginosis, trichomoniasis and chlamydia (n=12) (figure 1).

For women positive for each STI/RTI, online supplemental table 10 shows the proportion co-infected with the other four curable STIs/RTIs. Notably, among women who had syphilis, 60.8% (14/23) also had bacterial vaginosis. Among women with gonorrhoea, 43.2% (16/37) were co-infected with bacterial vaginosis, 31.6% (12/38) with chlamydia and 21.1% (8/38) with trichomoniasis. Among women positive for chlamydia, 37.8% (73/193) also had bacterial vaginosis and 17.2% (34/198) also had trichomoniasis.

Risk factors for STI/RTI co-infection

For the two most common STI/RTI co-infection combinations, primigravidae were nearly three times more likely than multigravidae to be co-infected with both STIs/RTIs than have neither infection (online supplemental table 11). Specifically, relative risk ratios (RRRs) of having a co-infection rather than neither infection among paucigravidae relative to multigravidae were RRR=2.6 (95% CI 1.2 to 5.7, p=0.020) for bacterial vaginosis and trichomoniasis, and RRR=2.9 (1.4 to 5.7, p=0.003) for bacterial vaginosis and chlamydia.

When controlling for all variables, women who were positive for bacterial vaginosis were 1.5 (95% CI 1.2 to 2.0 p=0.002) times more likely to have chlamydia, and 3.7 (95% CI 1.6 to 8.6, p=0.002) times more likely to have syphilis, relative to women without bacterial vaginosis (online supplemental table 12).

Prevalence: STI/RTIs and malaria co-infection

Among the subgroup of women tested for either chlamydia, gonorrhoea, trichomoniasis or bacterial vaginosis at the first ANC visit, 1401 were also tested for malaria by PCR. Of these, 6.7% (95% CI 5.5 to 8.1, n=94) had malaria plus an STI/RTI co-infection, 39.3% (95% CI 36.8 to 41.9, n=551) had STI/RTI only, 7.0% (95% CI 5.8 to 8.5, n=98) had malaria only and 47.0% (95% CI 44.4 to 49.6, n=658) had neither malaria nor an STI/RTI; see online supplemental table 13 for the prevalence of specific co-infection combinations. Prevalence of malaria and a curable STI/RTI co-infection was similar among paucigravidae (7.1% (95% CI 5.5 to 9.2)) and multigravidae (6.2% (95% CI 4.6 to 8.4)), p=0.508. Among women positive for malaria and tested for chlamydia, gonorrhoea, trichomoniasis or bacterial vaginosis, one-half (49.0%, 95% CI 41.9 to 56.0; n=94/192) also had at least one STI/RTI. The prevalence of malaria co-infection among women with bacterial vaginosis did not differ by Nugent score nor did the prevalence of malaria co-infection among women with syphilis differ by RPR titre (online supplemental figures 3, 4 and online supplemental table 14).

Table 3 Risk factors for syphilis, chlamydia, gonorrhoea and bacterial vaginosis during pregnancy

STI/RTI	Risk factor	Positive	Tested	Prevalence % (95% CI)	Prevalence ratio (95% CI)	P value	
Syphilis	Maternal age	<25 years	38	2432	1.6 (1.1 to 2.1)	1.4 (0.4 to 4.9)	0.578
		≥25 years	41	2137	1.9 (1.4 to 2.6)	Ref	
	Gravidity	Paucigravidae	32	2523	1.3 (0.9 to 1.8)	0.5 (0.2 to 1.7)	0.274
		Multigravidae	47	2033	2.3 (1.7 to 3.1)	Ref	
	Socio-economic tercile	Low	44	1516	2.9 (2.2 to 3.9)	8.8 (1.0 to 76.1)	0.048
		Medium	26	1525	1.7 (1.2 to 2.5)	8.8 (1.1 to 68.1)	0.037
		High	9	1528	0.6 (0.3 to 1.1)	Ref	
	Bacterial vaginosis	Negative	9	987	0.9 (0.5,1.7)	Ref	
		Positive	14	391	3.6 (2.1 to 6.0)	3.7 (1.6 to 8.4)	0.002
	Chlamydia	Maternal age	<25 years	124	763	16.3 (13.8 to 19.0)	1.0 (0.7 to 1.4)
≥25 years			74	668	11.1 (8.9 to 13.7)	Ref	
Gravidity		Paucigravidae	137	783	17.5 (15.0 to 20.3)	1.7 (1.2 to 2.4)	0.002
		Multigravidae	61	642	9.5 (7.5 to 12.0)	Ref	
Gonorrhoea	Maternal age	<25 years	28	763	3.7 (2.5 to 5.3)	1.6 (0.7 to 3.3)	0.237
		≥25 years	10	668	1.5 (0.8 to 2.8)	Ref	
	Gravidity	Paucigravidae	29	783	3.7 (2.6 to 5.3)	2.5 (1.2 to 5.4)	0.018
		Multigravidae	9	642	1.4 (0.7 to 2.7)	Ref	
	Maternal education	None/Primary	27	900	3.0 (2.1 to 4.3)	2.4 (1.2 to 5.1)	0.016
		Secondary/ higher	11	528	2.1 (1.2 to 3.7)	Ref	
		Low	14	466	3.0 (1.8 to 5.0)	0.9 (0.3 to 2.6)	0.876
	Socio-economic tercile	Medium	11	485	2.3 (1.3 to 4.1)	0.5 (0.2 to 1.0)	0.054
		High	13	480	2.7 (1.5 to 4.6)	Ref	
	Bacterial vaginosis	Negative	21	962	2.2 (1.4 to 3.3)	Ref	
Positive		16	383	4.2 (2.6 to 6.7)	1.8 (1.0 to 3.4)	0.074	
Bacterial vaginosis	Maternal age	<25 years	222	744	29.8 (26.5 to 33.1)	1.0 (0.8 to 1.2)	0.752
		≥25 years	177	658	26.9 (23.6 to 30.4)	Ref	
	Gravidity	Paucigravidae	241	774	31.1 (27.8 to 34.4)	1.3 (1.1 to 1.6)	0.018
	Multigravidae	156	623	25.0 (21.8 to 28.6)	Ref		

Country and study site controlled for in all models. Statistically significant results are in bold. Ref, reference; RTI, reproductive tract infection; STI, sexually transmitted infection.

Risk factors for malaria and curable STI/RTI co-infection

Risk factor analysis of malaria and curable STI/RTI co-infection identified women who had anaemia as being at greater risk of having a malaria and curable STI/RTI co-infection, rather than neither infection, specifically malaria and chlamydia (PR 7.7 (95% CI 2.5 to 23.7), p<0.001); malaria and trichomoniasis (PR 3.5 (95% CI 1.3 to 9.6), p=0.013); malaria and bacterial vaginosis (PR 2.2 (95% CI 1.2 to 4.1), p=0.011) (online supplemental table 15).

DISCUSSION

Our analysis of malaria and STI/RTI prevalence and risk factors associated with either STI/RTI, STI/RTI

co-infection or STI/RTI-malaria co-infections provides important insight into the epidemiology of infections in pregnancy in three East African countries. We report a very high prevalence of women with at least one curable STI/RTI (45.9%), highlighting the need for improved diagnosis and treatment interventions at ANC, where currently women are rarely screened, unless symptomatic.

The burden of malaria in pregnancy is reduced with the universal provision of IPTp with SP at scheduled ANC visits from the second trimester to delivery, although coverage remains inadequate and many malaria parasites have lost sensitivity to the intervention. Syphilis is managed with universal screening and treatment

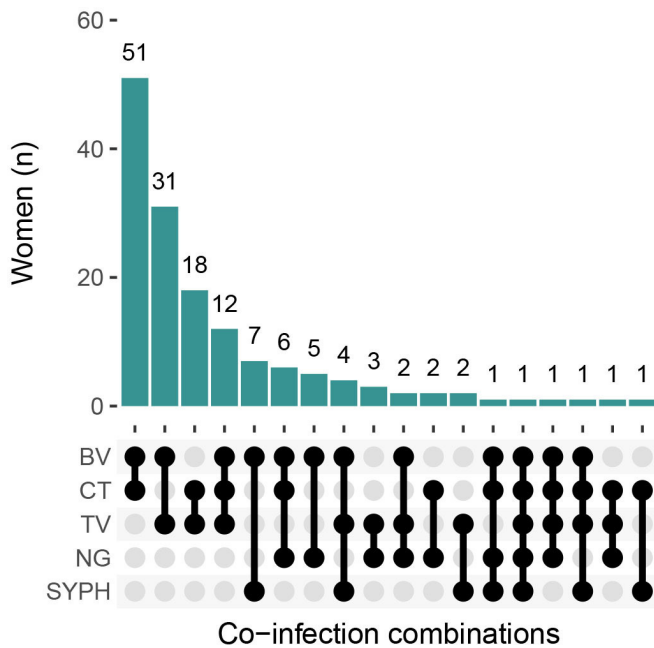


Figure 1 Frequency of unique combinations of sexually transmitted infection and reproductive tract infection co-infections. BV, bacterial vaginosis; CT, chlamydia; NG, gonorrhoea; SYPH, syphilis; TV, trichomoniasis;.

at ANC booking. Other curable STIs/RTIs, however, are relatively neglected. Presumptive treatment of all women for STIs/RTIs alongside IPTp has produced mixed results and raises important issues around antimicrobial resistance.^{21 22} The way forward in the near term for curable STIs/RTIs will likely involve options that range from universal screening for curable STIs/RTIs to risk-based presumptive treatment or risk-based screening-and-treatment.

Individual estimates of trichomoniasis and bacterial vaginosis prevalence were lower than recent estimates among pregnant women in East and Southern Africa.^{17 23} We found that 3 in 10 women had bacterial vaginosis, a consequential burden given its association with adverse reproductive and obstetric sequelae, and increased risk of STI and HIV acquisition and transmission.^{24 25} While prior reports of the prevalence of STI/RTI indicated a prevalence of chlamydia of 5.0% among the general population of women in the WHO African region²⁶ and 5.2% among ANC attendees in Eastern and Southern Africa,²³ we found a prevalence nearly three times higher at 13.8%. The prevalence was particularly high in Kenya, where one in five women overall, and one in four paucigravidae, had chlamydia. Our findings are in line with recent results among women 15–49 years of age in Zimbabwe²⁷ and South Africa,²⁸ potentially suggesting that the prevalence of chlamydia has increased in the subregion over the past decade. Antenatal services should aim to treat chlamydia infection early in pregnancy in order to prevent the associated adverse pregnancy outcomes.²⁹ The importance of this cannot be overstated; a secondary analysis of data from our trial found that fetal growth was restricted in women who had a curable STI/RTI in the

second trimester but was not restricted among women with an STI/RTI occurring only in the third trimester.³⁰ Advancements in the development of rapid, simple and well-performing point-of-care tests for chlamydia and gonorrhoea are much needed.³¹

We previously reported that the prevalence of at least one STI/RTI was 16.9%.¹⁸ This reflected results from the entire study population, for whom two-thirds of women were only tested for syphilis. In this current analysis, we restricted data to the randomly selected subsample of women tested for chlamydia, gonorrhoea, trichomoniasis and bacterial vaginosis. The prevalence of having at least one STI/RTI among this subgroup who underwent comprehensive STI/RTI testing was three times higher, 45.9%.

Paucigravidae were at greatest risk of individual STI/RTI and STI/RTI co-infection, regardless of their age. Many reasons may contribute to this difference. Paucigravidae may have comparatively lower acquired immunity than multigravidae due to fewer lifetime exposures, and may thus harbour STIs/RTIs for longer. Given the association between STIs/RTIs and pelvic inflammatory disease and infertility,³² it is possible that women attending ANC as multigravidae (women able to conceive numerous times) would be those with a lower prevalence of STIs/RTIs.

More women had at least one curable STI/RTI in this population than had malaria,¹⁸ and yet no systematic interventions or screening strategies beyond syndromic management are in place, with the exception of syphilis screening (the coverage of which can also be suboptimal).^{33 34} Results of ongoing trials investigating the integration of curable STI/RTI screening and treatment interventions at ANC will provide much-needed data.^{35 36}

The British HIV Association recommend screening HIV-infected pregnant women for bacterial vaginosis.³⁷ Similarly, given that 60.9% of women with syphilis were positive for bacterial vaginosis, presumptive bacterial vaginosis treatment or a screen-and-treat approach could be considered among all women with a positive syphilis test result. However, the low syphilis prevalence (0.6%) in this setting would have meant that just 3.6% of bacterial vaginosis cases would have received enhanced care under this approach. The low syphilis prevalence may also explain the reason that known risk factors such as maternal age and gravidity were not found to be statistically associated here. Presumptive treatment or a screen-and-treat approach for bacterial vaginosis based on STI presence could be extended to all women with any STI since among those with chlamydia, gonorrhoea and trichomoniasis, 37.8%, 43.2% and 26.4%, respectively, also had bacterial vaginosis. However, this would first require introduction of point-of-care testing for these infections.

Understanding the epidemiology of curable STIs/RTIs associated with adverse pregnancy outcomes is especially important in areas of malaria transmission, as almost half of the women with malaria in our study were co-infected

with a curable STI/RTI. A cohort study nested within the IMPROVE trial showed that infection during pregnancy with malaria-only, STI/RTI-only or co-infection with both, was associated with reduced fetal growth when compared with women without malaria or STI/RTIs. Malaria and STI/RTI co-infection had the most detrimental effect on fetal growth compared with malaria mono-infection or STIs/RTIs mono-infection.³⁰ As clinical trials continue to investigate alternatives to SP for IPTp, collecting data on malaria in pregnancy and on curable STIs/RTIs and co-infection are needed should the non-malarial effects conferred by different IPTp regimens be fully appreciated and the benefits of IPTp on improving birth outcomes maximised.³⁸

Strengths of this study include the large sample size that included sites in three East African countries, and the use of gold standard diagnostics for the five curable STIs/RTIs and malaria. HIV-infected women were excluded from the trial because WHO does not recommend providing IPTp with SP to women who are already receiving cotrimoxazole. SP and cotrimoxazole both contain sulfa and there is a theoretical risk of severe adverse reactions if taken concomitantly. Because the prevalence of malaria and STIs/RTIs among women living with HIV is higher than among HIV-negative women,^{39 40} our prevalence estimates likely understate the true burden among all pregnant women attending ANC.

In addition, 1041 women chose not to participate in the trial or their partner/spouse/another family member discouraged them from joining. We do not know if their non-participation affected our estimates. Finally, our prevalence and risk estimates cannot be assumed to be representative of all pregnant women; ANC attendees, and particularly those women who attended early enough to be eligible for inclusion in the trial, are not necessarily representative of the general pregnant population. Lack of ANC attendance and the opportunities it provides for interventions has been reported as a risk factor for curable STIs/RTIs.⁴¹ Our findings may not reflect the true burden of disease at the community level. Community delivery of IPTp is now a WHO recommendation to complement ANC IPTp administration. Further research into the burden of curable STIs/RTIs among non-ANC attending pregnant women is needed to inform interventions among this group.

In summary, malaria infection and curable STIs/RTIs were highly prevalent among women attending their first ANC visit, with paucigravidae at particular risk. Our results underscore the need for novel approaches to reduce the burden of malaria and curable STIs/RTIs in pregnancy and the need for further research into the burden of infection and co-infection, particularly given that improving pregnancy outcomes may be more difficult to achieve in the presence of co-infection without combination interventions. This will necessitate holistic approaches to ANC. In malaria-endemic East and Southern Africa, this may involve use of SP in combination with a more potent antimalarial therapy such as

dihydroartemisinin-piperazine alongside affordable, rapid and accurate point-of-care diagnostics for curable STIs/RTIs as early as possible in pregnancy.

Author affiliations

¹Department of Disease Control, London School of Hygiene & Tropical Medicine, London, UK

²School of Global and Public Health, Kamuzu University of Health Sciences, Blantyre, Malawi

³Department of Clinical Sciences, Academy of Medical Sciences, Malawi University of Science and Technology, Thyolo, Malawi

⁴Kenya Medical Research Institute, Center for Global Health Research, Kisumu, Kenya

⁵National Institute for Medical Research, Tanga Centre, Tanga, Tanzania

⁶National Institute for Medical Research, Mwanza Centre, Mwanza, Tanzania

⁷Kilimanjaro Clinical Research Institute and Kilimanjaro Christian Medical University College, Moshi, Tanzania

⁸Centre for Medical Parasitology, Department of Immunology and Microbiology, University of Copenhagen and Department of Infectious Diseases, Copenhagen University Hospital, Copenhagen, Denmark

⁹Department of Gynaecology and Obstetrics, Copenhagen University Hospital—North Zealand, Hilleroed, Denmark

¹⁰Department of Clinical Sciences, Liverpool School of Tropical Medicine, Liverpool, UK

¹¹Malaria Branch, Division of Parasitic Diseases and Malaria, Centre for Global Health, Centers for Disease Control and Prevention, Atlanta, Georgia, USA

¹²Great Ormond Street Institute of Child Health, University College London, London, UK

¹³Centre for Child Adolescent and Maternal Health Research, Faculty of Medicine and Health Technology, Tampere University, Tampere, Finland

¹⁴Fetal Medicine Unit, Saint George's Hospital, London, UK

¹⁵Department of Infectious Disease Epidemiology, Faculty of Epidemiology and Population Health, London School of Hygiene & Tropical Medicine, London, UK

X Hellen C Barsosio @drbarsosio

Contributors GRG-L, RMC, MM, Ftk conceived and designed the study. MM, HCCB, DTRM, GM, RAK, SK, KO, JD, UA, JH, JG, CM, SG, OAM, KSP, JPAL, AM, PI, QS, EDO, JM, MA, PJH, NK, CH, Ftk, RMC contributed to data acquisition. QS, CM, HH, RAK, SK, MA coordinated the laboratory component. GRG-L conducted the statistical analysis and wrote the first draft of the manuscript. GRG-L and RMC are the guarantors. All authors contributed to the data interpretation and critical revision for important intellectual content. All authors approved the final version submitted.

Funding The IMPROVE study was supported by the EDCTP2 programme (TRIA.2015-1076) under Horizon 2020; the UK Department of Health and Social Care, the UK Foreign Commonwealth and Development Office, the UK Medical Research Council and Wellcome Trust, through the Joint Global Health Trials scheme (MR/P006922/1); and the Swedish International Development Cooperation Agency. GRG-L was supported by an MRC PhD studentship (MR/N013638/1). CS was supported by Danish Independent Research Fund: Clinician Scientist Positions (1030-00371B). MC was supported by a Sir Henry Dale Fellowship jointly funded by the Wellcome Trust and the Royal Society (220658/Z/20/Z).

Disclaimer Dr. Julie Gutman is staff at the US Centers for Disease Control and Prevention (CDC). The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the CDC.

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval The trial was approved by the ethics committees of the Kenya Medical Research Institute, Nairobi, Kenya (SERU 75-3421); the College of Medicine, Blantyre, Malawi (P.02/17/2110); the Medical Research Coordinating Committee, Dar Es Salaam, Tanzania (NIMR/HQ/R.8a/Vol.1X/2533); the Liverpool School of Tropical Medicine (trial sponsor), Liverpool, UK and the London School of Hygiene & Tropical Medicine, London, UK. This activity was reviewed by Centres for Disease Control (CDC) and was conducted consistent with applicable US federal law and CDC policy. All study participants provided written informed consent at enrolment.

This consent included the long-term storage of samples for future research as well as the shipping of samples to external laboratories for relevant analyses. Personal and medical information relating to research participants was treated as confidential and only accessed by research staff. The risk of disclosure was minimised by secure storage of documents and by replacing personal identifiers with a unique study code to conceal the identity of the participant. Adverse events, particularly those associated with the study medication, were monitored throughout the trial and not found to differ significantly between treatment groups among pregnant women or their infants.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available on reasonable request. Individual participant data will be available from the Worldwide Antimalarial Resistance Network data repository

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: <https://creativecommons.org/licenses/by/4.0/>.

ORCID iDs

Georgia R Gore-Langton <http://orcid.org/0000-0002-7392-3127>

Mwayiwawo Madanitsa <http://orcid.org/0000-0002-4743-9382>

Hellen C Barsosio <http://orcid.org/0000-0002-0261-9814>

Crispin Mukerebe <http://orcid.org/0009-0001-6128-8840>

Jenny Hill <http://orcid.org/0000-0003-1588-485X>

Julie Gutman <http://orcid.org/0000-0001-6769-7322>

Patricia Jean Hunter <http://orcid.org/0000-0002-6224-0654>

Ulla Ashorn <http://orcid.org/0000-0002-1720-868X>

Matt Cairns <http://orcid.org/0000-0003-1068-9713>

R Matthew Chico <http://orcid.org/0000-0003-1795-7100>

REFERENCES

- Rogerson SJ, Desai M, Mayor A, *et al*. Burden, pathology, and costs of malaria in pregnancy: new developments for an old problem. *Lancet Infect Dis* 2018;18:e107–18.
- Moore KA, Simpson JA, Scoullar MJL, *et al*. Quantification of the association between malaria in pregnancy and stillbirth: a systematic review and meta-analysis. *Lancet Glob Health* 2017;5:e1101–12.
- Fried M, Duffy PE. Malaria during Pregnancy. *Cold Spring Harb Perspect Med* 2017;7:a025551.
- Adachi K, Nielsen-Saines K, Klausner JD. Chlamydia trachomatis infection in pregnancy: The global challenge of preventing adverse pregnancy and infant outcomes in Sub-Saharan Africa and Asia. *Biomed Res Int* 2016;2016:9315757.
- Juliana NCA, Suiters MJM, Al-Nasiry S, *et al*. The association between vaginal microbiota dysbiosis, bacterial vaginosis, and aerobic vaginitis, and adverse pregnancy outcomes of women living in Sub-Saharan Africa: a systematic review. *Front Public Health* 2020;8:567885.
- Gao R, Liu B, Yang W, *et al*. Association of maternal sexually transmitted infections with risk of preterm birth in the United States. *JAMA Netw Open* 2021;4:e2133413.
- Watson-Jones D, Changalucha J, Gumodoka B, *et al*. Syphilis in pregnancy in Tanzania. I. Impact of maternal syphilis on outcome of pregnancy. *J Infect Dis* 2002;186:940–7.
- Lawn JE, Blencowe H, Waiswa P, *et al*. Stillbirths: rates, risk factors, and acceleration towards 2030. *The Lancet* 2016;387:587–603.
- World Health Organisation. I. Impact of maternal syphilis on criteria and processes for validation: elimination of mother-to-child transmission of HIV and syphilis (Second Edition). 2017.
- World Health Organisation. *Management of patients with sexually transmitted diseases. Report of a WHO Study Group, vol. 810*. Geneva: World Health Organization, 1991.
- World Health Organisation. *Guidelines for the management of symptomatic sexually transmitted infections; Licence: CC BY-NC-SA 3.0 IGO*. 2021.
- Romoren M, Velauthapillai M, Rahman M, *et al*. Trichomoniasis and bacterial vaginosis in pregnancy: inadequately managed with the syndromic approach. *Bull World Health Organ* 2007;85:297–304.
- Chaponda EB, Bruce J, Michelo C, *et al*. Assessment of syndromic management of curable sexually transmitted and reproductive tract infections among pregnant women: an observational cross-sectional study. *BMC Pregnancy Childbirth* 2021;21:98.
- Thomas T, Choudhri S, Kariuki C, *et al*. Identifying cervical infection among pregnant women in Nairobi, Kenya: limitations of risk assessment and symptom-based approaches. *Sex Transm Infect* 1996;72:334–8.
- Lingani M, Zango SH, Valéa I, *et al*. Malaria and curable sexually transmitted and reproductive tract infection among pregnant women in rural Burkina Faso. *Trop Med Health* 2021;49:90.
- Zango SH, Lingani M, Valea I, *et al*. Malaria and curable sexually transmitted infections in pregnant women: A two-years observational study in rural Burkina Faso. *PLoS ONE* 2020;15:e0242368.
- Chaponda EB, Chico RM, Bruce J, *et al*. Malarial Infection and Curable Sexually Transmitted and Reproductive Tract Infections Among Pregnant Women in a Rural District of Zambia. *Am J Trop Med Hyg* 2016;95:1069–76.
- Madanitsa M, Barsosio HC, Minja DTR, *et al*. Effect of monthly intermittent preventive treatment with dihydroartemisinin-piperazine with and without azithromycin versus monthly sulfadoxine-pyrimethamine on adverse pregnancy outcomes in Africa: a double-blind randomised, partly placebo-controlled trial. *Lancet* 2023;401:1020–36.
- 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.
- Saidi Q, Minja D, Njau J, *et al*. Ultrasensitive qPCR-based detection of plasmodium falciparum in pregnant women using dried blood or whole blood pellet samples processed through different DNA extraction methods. *Am J Trop Med Hyg* 2021;106:846–9.
- Kimani J, Phiri K, Kamiza S, *et al*. Efficacy and safety of azithromycin-chloroquine versus sulfadoxine-pyrimethamine for intermittent preventive treatment of plasmodium falciparum Malaria infection in pregnant women in Africa: an open-label, randomized trial. *PLoS ONE* 2016;11:e0157045.
- Hallamaa L, Ashorn P, Cheung YB, *et al*. The Impact of Antenatal Azithromycin and Monthly Sulfadoxine-Pyrimethamine on Maternal Malaria during Pregnancy and Fetal Growth: A Randomized Controlled Trial. *Am J Trop Med Hyg* 2023;108:768–76.
- Chico RM, Mayaud P, Ariti C, *et al*. Prevalence of malaria and sexually transmitted and reproductive tract infections in pregnancy in sub-Saharan Africa: a systematic review. *JAMA* 2012;307:2079–86.
- Atashili J, Poole C, Ndumbe PM, *et al*. Bacterial vaginosis and HIV acquisition: a meta-analysis of published studies. *AIDS* 2008;22:1493–501.
- Cohen CR, Lingappa JR, Baeten JM, *et al*. Bacterial vaginosis associated with increased risk of female-to-male HIV-1 transmission: a prospective cohort analysis among African couples. *PLoS Med* 2012;9:e1001251.
- Rowley J, Vander Hoorn S, Korenromp E, *et al*. Chlamydia, gonorrhoea, trichomoniasis and syphilis: global prevalence and incidence estimates, 2016. *Bull World Health Organ* 2019;97:548–562P.
- Kranzer K, Simms V, Dauya E, *et al*. Identifying youth at high risk for sexually transmitted infections in community-based settings using a risk prediction tool: a validation study. *BMC Infect Dis* 2021;21:1234.
- Garrett NJ, Osman F, Maharaj B, *et al*. Beyond syndromic management: Opportunities for diagnosis-based treatment of sexually transmitted infections in low- and middle-income countries. *PLoS ONE* 2018;13:e0196209.
- Chen X-S. Adverse pregnancy outcomes due to Chlamydia trachomatis. *Lancet Infect Dis* 2018;18:499.
- Mtoto G, Chico RM, Madanitsa M, *et al*. Fetal growth and birth weight are independently reduced by malaria infection and curable sexually transmitted and reproductive tract infections in Kenya, Tanzania, and Malawi: A pregnancy cohort study. *Int J Infect Dis* 2023;135:28–40.
- Unemo M, Bradshaw CS, Hocking JS, *et al*. Sexually transmitted infections: challenges ahead. *Lancet Infect Dis* 2017;17:e235–79.
- Tsevat DG, Wiesenfeld HC, Parks C, *et al*. Sexually transmitted diseases and infertility. *Am J Obstet Gynecol* 2017;216:1–9.
- Perez F, Mayaud P. One step in the right direction: improving syphilis screening and treatment in pregnant women in Africa. *Lancet Glob Health* 2019;7:e550–1.

- 34 Kanyangarara M, Walker N, Boerma T. Gaps in the implementation of antenatal syphilis detection and treatment in health facilities across sub-Saharan Africa. *PLOS ONE* 2018;13:e0198622.
- 35 Medina-Marino A, Cleary S, Muzny CA, *et al.* Sexually transmitted infection screening to prevent adverse birth and newborn outcomes: study protocol for a randomized-controlled hybrid-effectiveness trial. *Trials* 2022;23:441.
- 36 Martin K, Dziva Chikwari C, Dauya E, *et al.* Investigating point-of-care diagnostics for sexually transmitted infections and antimicrobial resistance in antenatal care in Zimbabwe (IPSAZ): protocol for a mixed-methods study. *BMJ Open* 2023;13:e070889.
- 37 British HIV Association. British hiv association guidelines for the management of hiv in pregnancy and postpartum 2018 (2020 third interim update). 2020.
- 38 Roh ME, Kuile FOT, Rerolle F, *et al.* Overall, anti-malarial, and non-malarial effect of intermittent preventive treatment during pregnancy with sulfadoxine-pyrimethamine on birthweight: a mediation analysis. *Lancet Glob Health* 2020;8:e942–53.
- 39 Medina-Marino A, Mudau M, Kojima N, *et al.* Persistent *Chlamydia trachomatis*, *Neisseria gonorrhoeae* or *Trichomonas vaginalis* positivity after treatment among human immunodeficiency virus-infected pregnant women, South Africa. *Int J STD AIDS* 2020;31:294–302.
- 40 ter Kuile FO, Parise ME, Verhoeff FH, *et al.* The burden of co-infection with human immunodeficiency virus type 1 and malaria in pregnant women in sub-saharan Africa. *Am J Trop Med Hyg* 2004;71:41–54.
- 41 Rours GIJG, Verkooyen RP, Hop WCJ, *et al.* Sexually transmitted infections in pregnant urban South African women: socio-economic characteristics and risk factors. *S Afr J Epidemiol Infect* 2006;21:14–9.