

8-30-2023

Prevention of Maternal and Neonatal Death/Infections with a Single Oral Dose of Azithromycin in Women in Labour in Low-Income and Middle-Income Countries (A-PLUS): A Study Protocol for a Multinational, Randomised Placebo-Controlled Clinical Trial

Follow this and additional works at: <https://jdc.jefferson.edu/gha>

Jennifer Hemingway-Foday

 Part of the [Diseases Commons](#)

Alan Tita

[Let us know how access to this document benefits you](#)

Elwyn Chomba

Recommended Citation

Musaku Mwenechanya

Hemingway-Foday, Jennifer; Tita, Alan; Chomba, Elwyn; Mwenechanya, Musaku; Mweemba,

Treacius; Mweemba, Treacius; Lokangaka, Adrien; Tshefu Kitoto, Antoinette; Lomendje, Gustave;

Hibberd, Patricia L.; Patel, Archana; Das, Prabir Kumar; Kurhe, Kunal; Goudar, Shivaprasad S;

Kavi, Avinash; Metoud, Mrityunjay; Saleem, Sarah; Tikmani, Shiyam S.; Esamai, Fabian;

Nyongesa, Paul; Sagwe, Amos; Figueroa, Lester; Mazariegos, Manolo; Billah, Sk Masum; Haque,

Rashidul; Shahjahan Siraj, Md; Goldenberg, Robert L.; Bauserman, Melissa; Bose, Carl; Liechty,

Edward A.; Ekhuagere, Osayame A.; Krebs, Nancy F.; Derman, Richard; Petri, William A.; Koso-

Thomas, Marion; McClure, Elizabeth; and Carlo, Waldemar A., "Prevention of Maternal and

Neonatal Death/Infections with a Single Oral Dose of Azithromycin in Women in Labour in Low-Income and Middle-Income Countries (A-PLUS): A Study Protocol for a Multinational,

Randomised Placebo-Controlled Clinical Trial" (2023). *Global Health Articles*. Article 31.

<https://jdc.jefferson.edu/gha/31>

This Article is brought to you for free and open access by the Jefferson Digital Commons. The Jefferson Digital Commons is a service of Thomas Jefferson University's [Center for Teaching and Learning \(CTL\)](#). The Commons is a showcase for Jefferson books and journals, peer-reviewed scholarly publications, unique historical collections from the University archives, and teaching tools. The Jefferson Digital Commons allows researchers and interested readers anywhere in the world to learn about and keep up to date with Jefferson scholarship. This article has been accepted for inclusion in Global Health Articles by an authorized administrator of the Jefferson Digital Commons. For more information, please contact: JeffersonDigitalCommons@jefferson.edu.

Authors

Jennifer Hemingway-Foday, Alan Tita, Elwyn Chomba, Musaku Mwenechanya, Trecious Mweemba, Tracy Nolen, Adrien Lokangaka, Antoinette Tshetu Kitoto, Gustave Lomendje, Patricia L. Hibberd, Archana Patel, Prabir Kumar Das, Kunal Kurhe, Shivaprasad S Goudar, Avinash Kavi, Mrityunjay Metgud, Sarah Saleem, Shiyam S. Tikmani, Fabian Esamai, Paul Nyongesa, Amos Sagwe, Lester Figueroa, Manolo Mazariegos, Sk Masum Billah, Rashidul Haque, Md Shahjahan Siraj, Robert L. Goldenberg, Melissa Bauserman, Carl Bose, Edward A. Liechty, Osayame A. Ekhaguere, Nancy F. Krebs, Richard Derman, William A. Petri, Marion Koso-Thomas, Elizabeth McClure, and Waldemar A. Carlo

BMJ Open Prevention of maternal and neonatal death/infections with a single oral dose of azithromycin in women in labour in low-income and middle-income countries (A-PLUS): a study protocol for a multinational, randomised placebo-controlled clinical trial

To cite: Hemingway-Foday J, Tita A, Chomba E, *et al*. Prevention of maternal and neonatal death/infections with a single oral dose of azithromycin in women in labour in low-income and middle-income countries (A-PLUS): a study protocol for a multinational, randomised placebo-controlled clinical trial. *BMJ Open* 2023;**13**:e068487. doi:10.1136/bmjopen-2022-068487

► Prepublication history and additional supplemental material for this paper are available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2022-068487>).

Received 19 September 2022
Accepted 15 May 2023



► <http://dx.doi.org/10.1136/bmjopen-2022-067581>










Link or Companion Paper: [bmjopen-2022-067581](https://doi.org/10.1136/bmjopen-2022-067581) Copyright may not be established in the United States for works of government employees (17 U.S.C. § 105). The NIH Public Access Policy requires NIH authors to submit final, peer-reviewed journal manuscripts to the digital archive PubMed Central upon acceptance for publication but no later than 12 months after publication."

For numbered affiliations see end of article.

Correspondence to

Jennifer Hemingway-Foday; hemingway@rti.org

Jennifer Hemingway-Foday ¹, Alan Tita,² Elwyn Chomba,³ Musaku Mwenechanya,³ Trecious Mweemba,³ Tracy Nolen,¹ Adrien Lokangaka,^{4,5} Antoinette Tshetu Kitoto,^{4,5} Gustave Lomendje,⁵ Patricia L Hibberd,⁶ Archana Patel,^{7,8} Prabir Kumar Das,⁷ Kunal Kurhe,⁷ Shivaprasad S Goudar,⁹ Avinash Kavi ⁹, Mrityunjay Metgud,⁹ Sarah Saleem ¹⁰, Shiyam S Tikmani,¹⁰ Fabian Esamai,¹¹ Paul Nyongesa ¹¹, Amos Sagwe,¹¹ Lester Figueroa,¹² Manolo Mazariegos,¹² Sk Masum Billah ^{13,14}, Rashidul Haque,¹⁴ Md Shahjahan Siraj ¹⁴, Robert L Goldenberg,¹⁵ Melissa Bauserman,¹⁶ Carl Bose ¹⁶, Edward A Liechty,¹⁷ Osayame A Ekhuagere ¹⁷, Nancy F Krebs,¹⁸ Richard Derman,¹⁹ William A Petri,²⁰ Marion Koso-Thomas,²¹ Elizabeth McClure ¹, Waldemar A Carlo²

ABSTRACT

Introduction Maternal and neonatal infections are among the most frequent causes of maternal and neonatal mortality, and current antibiotic strategies have been ineffective in preventing many of these deaths. A randomised clinical trial conducted in a single site in The Gambia showed that treatment with an oral dose of 2 g azithromycin versus placebo for all women in labour reduced certain maternal and neonatal infections. However, it is unknown if this therapy reduces maternal and neonatal sepsis and mortality. In a large, multinational randomised trial, we will evaluate the impact of azithromycin given in labour to improve maternal and newborn outcomes.

Methods and analysis This randomised, placebo-controlled, multicentre clinical trial includes two primary hypotheses, one maternal and one neonatal. The maternal hypothesis is to test whether a single, prophylactic intrapartum oral dose of 2 g azithromycin given to women in labour will reduce maternal death or sepsis. The neonatal hypothesis will test whether this intervention will reduce intrapartum/neonatal death or sepsis. The intervention is a single, prophylactic intrapartum oral dose of 2 g azithromycin, compared with a single intrapartum oral dose of an identical appearing placebo. A total of 34 000 labouring women from 8 research sites in sub-Saharan Africa, South Asia and Latin America will be randomised with a one-to-one ratio to intervention/

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ This study evaluates the use of routine prophylactic azithromycin in women who undergo labour and attempt a vaginal delivery, including those at high risk of infection due to prolonged labour or membrane rupture, which was identified as a top research priority by the WHO and other stakeholders.
- ⇒ The study evaluates a simple, inexpensive intervention that can be scaled up to reduce the burden of both maternal and neonatal mortality due to infections if successful.
- ⇒ To optimise generalisability, this study will be carried out in a racially, ethnically and geographically diverse population in seven low-and-middle-income countries.
- ⇒ In response to global public health concerns about increasing antimicrobial resistance, this study includes an ancillary study to assess antimicrobial resistance and microbiome changes.
- ⇒ In some locations, azithromycin and other antibiotics are increasingly being used, especially with increasing use of caesarean section, and thus the impact of the outcome may be diminished.

placebo. In addition, we will assess antimicrobial resistance in a sample of women and their newborns.

Ethics and dissemination The study protocol has been reviewed and ethics approval obtained from all the relevant ethical review boards at each research site. The

results will be disseminated via peer-reviewed journals and national and international scientific forums.

Trial registration number NCT03871491 (<https://clinicaltrials.gov/ct2/show/NCT03871491?term=NCT03871491&draw=2&rank=1>).

INTRODUCTION

Background

Maternal infection during pregnancy and the puerperium, accounting for 10% of maternal deaths, is a top cause of maternal mortality worldwide.¹ Maternal infections increase neonatal sepsis, a leading cause of neonatal death in low-income countries (LICs).^{2–5} Neonatal infection accounts for 15% of neonatal mortality worldwide.² According to the WHO, maternal and neonatal deaths from infections have remained unchanged or increased, whereas other causes of death have declined.^{5 6}

Innovative, effective and scalable interventions are needed to reduce infection-related maternal and neonatal mortality. The evidence backing current WHO guidelines to prevent and treat peripartum infections is limited.⁷ Further, current approaches to address neonatal sepsis have had a limited impact and alternative antibiotic regimens suggesting comparable effectiveness have not reduced newborn deaths.^{8–11}

Risk factors for maternal and neonatal infection and sepsis

Caesarean delivery (CD), especially after labour or membrane rupture, is an important risk factor for maternal peripartum infection.¹² Antibiotic prophylaxis for CD is a well-established, effective strategy to reduce infection.^{13–16} However, with a prevalence of <5% in LICs, CDs do not account for most maternal sepsis worldwide.¹⁷ Many peripartum infections in low-and-middle-income countries (LMICs) occur in women who deliver vaginally, with women undergoing prolonged labour or membrane rupture at highest risk. Although most pregnancies are not at high-risk for infection due to prolonged labour or membrane rupture, pregnancies where this does occur account for about 50% of maternal and neonatal infections in LMICs.¹ Therefore, identification and management of prolonged labour and/or membrane rupture are critical to reduce maternal sepsis.⁷ Thus, the WHO has identified evaluation of routine prophylactic antibiotics for women undergoing vaginal birth or with prolonged labour/membrane rupture as a top research priority.¹

Intrapartum azithromycin to prevent maternal and neonatal infections

A novel approach to prevent maternal and neonatal infection is to target organisms that may be frequent pathogens but are not historically treatment targets.^{16 18–21} Azithromycin, which is available as a generic agent, has a bimodal half-life of 70 hours in the non-pregnant population. It is commonly used during pregnancy to treat chlamydia, gonorrhoea and

other infections.²² It provides broad coverage against most common pathogens associated with peripartum infections including gram-positive cocci, genital mycoplasmas and *Ureaplasma* infections, and certain gram-negative Bacilli and anaerobes associated with maternal polymicrobial infections and sepsis.²³ In addition, azithromycin also has activity against Group B *streptococcus*, a major cause of neonatal sepsis in developed countries and possibly in LICs.^{24 25}

Evidence from prior studies

In a US randomised clinical trial (RCT), a 50% reduction in maternal peripartum infection resulted from 500 mg intravenous azithromycin added to the standard prophylactic regimen in high-risk women undergoing CD following labour or membrane rupture >4 hours.¹⁸ These results were observed despite universal antibiotics in both treatment groups. Readmission, unscheduled visits, serious adverse events, postpartum fever or subsequent antibiotic treatment were also reduced in the azithromycin group. A related cost-analysis estimated that adjunctive azithromycin saved US\$360 per use in high-risk CDs and US\$143 per use in scheduled CDs.²⁶

A trial in The Gambia suggested the potential for azithromycin prophylaxis to improve maternal and neonatal outcomes.¹⁹ Among 829 participants randomised to 2 g of azithromycin versus placebo before delivery, maternal infections were lower in the azithromycin group, 3.6% vs 9.2%, respectively, (RR 0.40, 95% CI, 0.22 to 0.71, p=0.002). Among newborns, infections were also lower in the azithromycin group (18.1% vs 23.8%; RR 0.76, 95% CI, 0.58 to 0.99; p=0.052). Maternal and neonatal carriage of infectious organisms was lower in the azithromycin group.

Rationale for a trial

Drawing from these findings, the role of a single oral dose of azithromycin (plus usual care) to prevent maternal death or peripartum sepsis and intrapartum/neonatal death or sepsis appears promising. Of interest is the subpopulation at highest risk for infection due to prolonged labour and/or prolonged membrane rupture. Considering the success with a single 2 g dose in The Gambian trial, which is bioequivalent to the 500 mg intravenous dose used successfully in the US trial, as well as the 40% bioavailability of oral azithromycin, 2 g is considered the most appropriate dose for the proposed intervention. An RCT is the best design to evaluate efficacy and provide evidence for future policy and clinical practice decisions. Such an RCT in a racially, ethnically and geographically diverse population could be accomplished through the established infrastructure of the Global Network for Women's and Children's Health Research (Global Network), a multicountry research network with sites in India, Pakistan, Bangladesh, Guatemala, Kenya, Zambia and

the Democratic Republic of Congo (DRC) and an established data coordinating centre (DCC).²⁷

METHODS AND ANALYSIS

Study design and intervention

This study is a masked, placebo-controlled, multicentre, RCT. The intervention is a single 2 g dose of oral azithromycin or identical placebo, administered as four 500 mg pills following randomisation. Both groups will also receive the local standard of care during labour, delivery and postpartum. The trial adheres to the Standard Protocol Items: Recommendations for Interventional Trials statement.²⁸

Primary hypothesis

The Azithromycin-Prevention in Labor Use Study (A-PLUS) trial has both a maternal and neonatal primary hypothesis:

- ▶ Maternal: a single, prophylactic intrapartum oral dose of 2 g azithromycin given to women in labour will reduce *maternal* death or sepsis.
- ▶ Neonatal: a single, prophylactic intrapartum oral dose of 2 g azithromycin given to women in labour will reduce *intrapartum/neonatal* death or sepsis.

Study population

Pregnant women labouring in health facilities at eight Global Network sites will be screened by research staff per the eligibility criteria (box 1). The Global Network sites are described in online supplemental appendix 1 of the trial protocol. The secondary population is a cohort of 5500 high-risk women, defined as term and preterm (≥ 28 weeks) pregnant women with prolonged labour (≥ 18 hours) or prolonged membrane rupture (≥ 8 hours). A cohort of mother–infant dyads will also be randomly selected for an ancillary study to assess antimicrobial resistance and microbiome changes. Recruitment took place from September 2020 to August 2022, and data collection for the primary outcome was completed in October 2022. Data collection for the ancillary study of antimicrobial resistance is expected to be complete by August 2023.

Patient and public involvement

Patients and/or the public were not involved in study design, conduct or reporting.

Primary outcomes

The primary outcomes are:

- ▶ Maternal: incidence of maternal death or sepsis within 42 days post delivery in the intervention versus placebo group.
- ▶ Neonatal: incidence of intrapartum/neonatal death or sepsis with 28 days post delivery in the intervention versus placebo group.

We use the 2017 WHO definition of sepsis which includes a suspicion of infection and the presence of organ dysfunction based on clinical findings.⁷

Box 1 Eligibility criteria

Inclusion criteria

- ⇒ Pregnant women in labour with gestational age ≥ 28 weeks (by best estimate) who plan to deliver vaginally in a facility.
- ⇒ Admission to health facility with clear plan for spontaneous or induced vaginal delivery.
- ⇒ Presence of one or more live fetus confirmed via a fetal heart rate by Doptone prior to randomisation.
- ⇒ Age ≥ 18 years (minors 14–17 years eligible in countries where married or pregnant minors or their authorised representatives are legally permitted to give consent).
- ⇒ Provision of written informed consent.*

Exclusion criteria

- ⇒ Non-emancipated minors (as per local regulations).
- ⇒ Evidence of chorioamnionitis or other infection requiring antibiotic therapy at time of eligibility (women given single prophylactic antibiotics with no plans to continue after delivery will not be excluded).
- ⇒ Arrhythmia or known history of cardiomyopathy.
- ⇒ Allergy to azithromycin or other macrolides that is self-reported or documented in the medical record.
- ⇒ Any use of azithromycin, erythromycin or other macrolide in the 3 days or less prior to randomisation.
- ⇒ Plan for caesarean delivery prior to enrolment.
- ⇒ Preterm labour undergoing management with no immediate plan to proceed to delivery.
- ⇒ Advanced stage of labour (> 6 cm or 10 cm cervical dilation per local standards) and pushing or too distressed to understand, confirm or give informed consent regardless of cervical dilation.
- ⇒ Not capable of giving consent due to other health problems such as obstetric emergencies (eg, antepartum haemorrhage) or mental disorder.
- ⇒ Any medical condition considered a contraindication per the judgement of site investigators.
- ⇒ Previous randomisation in the trial.

* A model written informed consent form is available as online supplemental appendix 2 of the trial protocol.

Maternal sepsis is defined as organ dysfunction resulting from suspected or confirmed infection that occurs post randomisation during labour or the postpartum period. This WHO definition is operationalised as suspected or confirmed infection based on the presence of fever or hypothermia plus one or more signs of mild-to-moderate organ dysfunction including tachycardia, low blood pressure, tachypnea, altered mental status/confusion, reduced urinary output, jaundice or renal failure.^{7 29–31} Components of peripartum infection considered in diagnosing suspected or confirmed infection include clinical chorioamnionitis, endometritis, wound infections (perineal or caesarean), abdominal or pelvic abscess, mastitis/breast abscess or infection, pyelonephritis, pneumonia, and other bacterial infections (table 1).

Neonatal sepsis is defined as proven or possible serious bacterial infection (PSBI) or pneumonia, or meningitis. PSBI will be determined using WHO criteria of severe chest in-drawing, fever, hypothermia, no movement at all or movement only on stimulation, feeding poorly or not feeding at all and/or convulsions.³² Clinical and

**Table 1** Specified infections considered for maternal sepsis diagnosis

Type of infection	Azithromycin-Prevention in Labor Use Study (A-PLUS) trial definition
Chorioamnionitis	Fever (>100.4°F/38°C) in addition to one or more of the following: fetal tachycardia ≥160 bpm, maternal tachycardia >100 bpm, tender uterus between contractions or purulent/foul smelling discharge from the uterus prior to delivery.
Endometritis	Fever (>100.4°F/38°C) in addition to one or more of maternal tachycardia >100 bpm, tender uterine fundus or purulent/foul smelling discharge from the uterus after delivery.
Wound infection	Purulent infection (superficial or deep infection including necrotising fasciitis) of a perineal or caesarean wound with or without fever. In the absence of purulence, a wound infection requires presence of fever (>100.4°F/38°C) and at least one of the following signs of local infection: pain or tenderness, swelling, heat or redness around the incision/laceration.
Abdominopelvic abscess	Evidence of pus in the abdomen or pelvis noted during open surgery, interventional aspiration or imaging.
Pneumonia	Fever (>100.4°F/38°C) and clinical symptoms suggestive of lung infection including cough and/or tachypnea (>24 breaths/min) or radiological confirmation.
Pyelonephritis	Fever (>100.4°F/38°C) and one or more of the following: urinalysis/dip suggestive of infection, costovertebral angle tenderness or confirmatory urine culture.
Mastitis/breast abscess or infection	Fever (>100.4°F/38°C) and one or more of the following: breast pain, swelling, warmth, redness or purulent drainage.

laboratory signs of infection will also be considered for diagnosis.

Centralised adjudication of key infection outcomes will be implemented to standardise the results. Reported antibiotic treatment and culture status will also be considered for outcome adjudication.

The individual components of these primary outcomes will be analysed.

Secondary maternal outcomes

- ▶ The primary maternal outcome in a high-risk for infection population.
- ▶ Incidence of specific maternal infections (table 1).
- ▶ Subsequent maternal antibiotic therapy after randomisation to 42 days postpartum for any reason.
- ▶ Time from drug administration until initial discharge after delivery.
- ▶ Maternal readmissions and admissions to special care units within 42 days postpartum.
- ▶ Maternal unscheduled visit for care.
- ▶ Maternal gastrointestinal symptoms (nausea, vomiting, diarrhoea) or other reported side effects.
- ▶ Maternal death due to sepsis using the Global Network algorithm for cause of death.³³

Secondary neonatal outcomes

- ▶ Neonatal deaths due to sepsis and all-cause neonatal death in all participants and in labouring women at high risk of infection.
- ▶ Other neonatal infections (eg, eye infection, skin infection, omphalitis, urinary tract infection, respiratory rate ≥60 breaths/minute).
- ▶ Neonatal readmissions and admissions to special care units within 42 days postpartum.
- ▶ Neonatal initial hospital length of stay, defined as time of delivery until initial discharge.

- ▶ Neonatal unscheduled visit for care.
- ▶ Neonatal death due to sepsis using the Global Network algorithm for cause of death.³⁴
- ▶ Incidence of pyloric stenosis within 42 days of delivery.

Study procedures

All sites will train research staff in standardised study procedures to administer the intervention and assess outcomes. The sequence of study activities is described (figure 1). Study procedures are also provided (table 2). A more detailed schedule of study procedures is available as online supplemental appendix 3 of the trial protocol.

Site preparation

Prior to the trial, sites conducted an observational pilot study using the RCT's planned infrastructure to characterise current practices and optimise identification of suspected infection. The pilot study data was used to validate estimates of intrapartum deaths, maternal sepsis and neonatal sepsis for the RCT's sample size. Each site also met with local health authorities and conducted community sensitisation to ensure that study procedures were appropriate for the local context and to encourage facility and community-level engagement.

Screening

Pregnant women admitted for delivery at predefined health facilities will be screened for eligibility by research staff, per harmonised criteria (box 1). If a contraindication to participation is found, the woman will be excluded.

Consent

Research staff at each site are responsible for obtaining informed consent prior to complete cervical dilation or dilation limit approved by local authorities. Participants who are randomly selected to participate in the ancillary

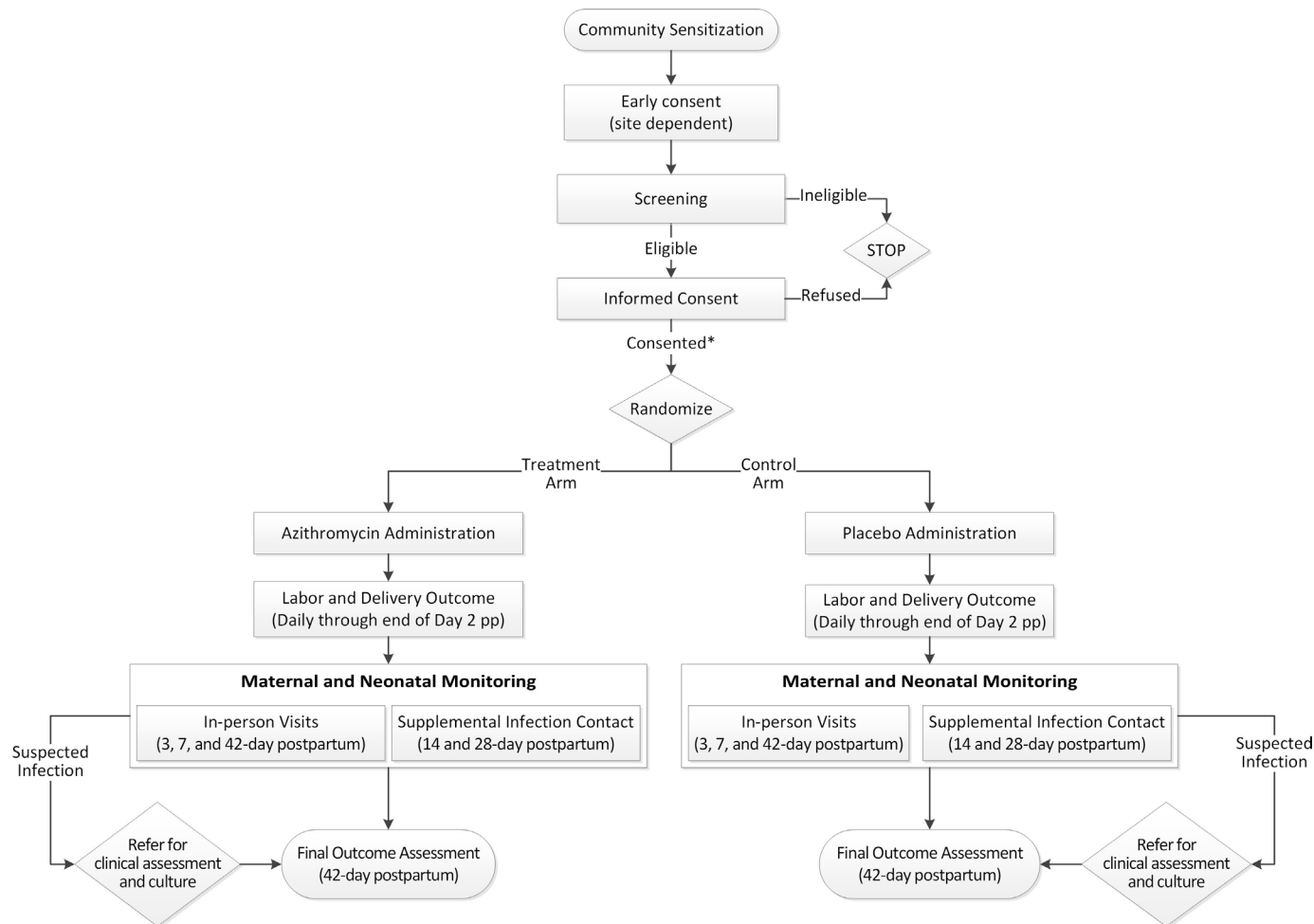


Figure 1 Study Flowchart. *If early consent, confirmation of eligibility and reconfirmation of consent required.

study on antimicrobial resistance will also be consented at this time. To minimise burden on the labouring women, sites may obtain initial consent from potential participants during the antenatal period and reconfirm consent at screening. As literacy levels vary, the consent form will be reviewed verbally. All research staff responsible for consent procedures will be trained and certified in the protection of human subjects and the study-specific consent procedures.

Masking

Both the azithromycin and placebo will be procured from the same manufacturer (Idifarma, Spain). The packaging will be standardised across sites and labelled as: 'azithromycin 2g or placebo' with the expiration date and a unique identifier. Clinical and research staff, and participants, will be masked to treatment status unless a serious adverse event potentially related to the treatment requires unmasking. Each site's study pharmacist will monitor randomisation, drug supply and safety.

Randomisation

A computer-generated randomisation scheme, created and known only by the DCC, will use a randomly permuted block design with randomly varied block to achieve a 1:1

randomisation, stratified by site. Each site will receive study drug directly from the manufacturer, which is sequentially prepared per the randomisation scheme in identical packaging. The study drug will be distributed to participating health facilities and dispensed sequentially by study staff.

Monitoring before discharge (baseline assessment)

Routine postdelivery care will be provided by clinical providers, trained to contact research staff in case of suspected or confirmed infection. Research staff will collect demographic characteristics, key clinical measures and outcomes from randomisation until discharge. Data will be abstracted from medical records or collected directly from participants, as relevant. All data will be directly entered into a secure electronic data capture platform, Research Electronic Data Capture (REDCap). Deidentified data will then be transmitted to the DCC. In addition, participants will be educated on signs and symptoms of infection and encouraged to call research staff with any concerns.

Monitoring after discharge (postpartum follow-up assessments)

After discharge, research staff will contact participants at the following timepoints:

**Table 2** Schedule of study procedures

	Antenatal Care visits	During labour/pre delivery	Post delivery/pre discharge	3-day postpartum (pp)	7-day pp	14-day pp	28-day pp	42-day pp
Community sensitisation	X							
Screening								
Eligibility confirmation		X						
Clinical assessment		X						
Consent		X						
Randomisation		X						
Drug administration		X						
Baseline data collection								
Sociodemographic information		X	X					
Medical history		X	X					
Labour and delivery (L&D) information			X					
Monitoring								
Drug side effects*		X	X	X	X			X
Maternal events during L&D		X	X					
Neonatal events during L&D			X					
Maternal infection/sepsis*			X	X	X			X
Neonatal infection/sepsis*			X	X	X			X
Maternal death*			X	X	X			X
Stillbirth or neonatal death<28 days of birth*			X	X	X	X	X	X
Infant mortality≥28 days of birth*								X
Pyloric stenosis*			X	X	X			X
Other maternal outcomes*			X	X	X			X
Other neonatal outcomes*			X	X	X			X
Unintended medical visits*			X	X	X			X
Serious adverse events*			X	X	X			X

*Events that may be reported by participant between scheduled study visits.

- ▶ In-person visits at 3-day, 7-day and 42-day postpartum to identify maternal or infant infection, unexpected medical visits, side effects and other study outcomes. If suspected infection, research staff will collect and send relevant specimens to the local microbiology lab for analysis.
- ▶ Supplemental phone contacts at 14-day and 28-day postpartum to review maternal and neonatal signs of infection (WHO criteria). If signs are identified, participants will visit a study facility for further assessment. These supplemental contacts will reinforce the participants' ability to self-assess maternal and neonatal infection and improve identification between the 7-day and 42-day visits.

If indicated, records of non-study visits to health providers during the follow-up period will be reviewed to ascertain study outcomes. Providers may be called if clarification is needed. Readmissions and related diagnoses identified during follow-up visits will be validated through medical record review. All data will be entered into REDCap, and deidentified data will be transmitted to the DCC.

Safety monitoring

Surveillance of maternal side effects including nausea, vomiting and diarrhoea/loose stools, abdominal pain, vaginitis and dizziness potentially associated with azithromycin will be conducted during labour and postpartum. For infants, findings suggestive of pyloric stenosis will be assessed during the follow-up visits. Maternal and neonatal surveillance will also include assessment of unintended medical visits, maternal deaths, stillbirths, neonatal death<28 days of birth and infant death>28 days. Additional maternal and neonatal risks associated with azithromycin use include anaphylaxis, allergic reactions (rash), liver failure and arrhythmias. Although rare, these side effects will be monitored and reported as serious adverse events. All safety outcomes will be reviewed by the data monitoring committee (DMC) appointed by the National Institute of Child Health and Human Development (NICHD).

Analytical plan

Baseline demographic characteristics and key clinical measures will be compared between the treatment arms

using contingency table approaches for categorical variables and analysis of variance models' continuous variables.

P values presented will be based on two-sided tests unless otherwise specified, adjusted for site. For most analyses, the interaction between treatment and site will be assessed and if significant, also be presented by site. For continuous outcomes, distributional properties will be evaluated and if required, transformations or non-parametric tests employed. Additional details for potential covariate adjustments in secondary analyses or handling violations of analytic method assumptions will be detailed in the statistical analysis plan.

Three key populations are of interest for study analyses:

1. The intention-to-treat (ITT) population will be the primary analysis population and will include all women randomised and their infants. Analyses of this cohort will be conducted based on randomised treatment.
2. The high risk for infection subgroup will include all women in the ITT and their infants' meeting criteria for being high risk (ie, prolonged labour and/or rupture of membranes) at randomisation. Analyses of this cohort will be conducted based on randomised treatment.
3. The as-treated population will include all randomised participants that receive any study drug during the study and their infants. Analyses of this cohort will be conducted based on treatment received.

The final determination of analysis population membership will be via a masked data review prior to final study analyses to address any potential anomalous cases that may arise in this large study population (eg, randomisation/treatment of a woman who is discharged prior to delivery due to false labour or unresponsiveness to induction).

Primary analysis

Incidence of maternal death or sepsis and intrapartum/neonatal death or sepsis will be compared separately between the treatment arms using generalised linear models fit with each binary outcome separately as the outcome measure. Estimates of RR and associated 95% CIs will be reported. The model will include terms for treatment and site. As randomisation occurs at the pregnancy level and approximately 1%–2% of pregnancies are anticipated to be multiple gestations, models for neonatal outcomes will account for correlation among multiples assuming an exchangeable covariance structure. For the two primary outcomes, analyses will be conducted using the ITT population and the p values associated with the treatment term will be used to formally test each of the two primary hypotheses at the $\alpha=0.05$ level.

As secondary analyses of the primary outcomes, assuming an overall treatment effect is observed, the models will be run including region (Africa, Latin America or Asia) and a treatment by region interaction

term. If the interaction term has a $p<0.1$, effects will be reported by region with treatment effect within region tested at the 0.025 level.

Additional exploratory models will also be run including individually: (1) a treatment-by-site interaction term, (2) any other antibiotic use during labour (yes or no) and its interaction with treatment and (3) mode of delivery (caesarean or vaginal) and its interaction with treatment. If the interaction term for any of these models has a $p<0.1$, then effects will also be reported by the relevant subgroups.

From each final model, estimates of relative risk associated with treatment will be obtained including unadjusted estimates of risk from the primary model as well as estimates of risk adjusted for potential confounders from the secondary analyses.

Secondary analyses: women at high risk for infection cohort

The major secondary aim is assessing the two primary outcomes (ie, incidence of maternal death or sepsis and incidence of intrapartum/neonatal death or sepsis) in the high-risk cohort. These analyses will also assess if the treatment effect differs between the high-risk subgroup versus non-high-risk women, defined as all women and their infants in the ITT population that delivered prior to meeting high-risk criteria. Specifically, the maternal and neonatal primary outcomes will be modelled including a treatment by risk status interaction term and excluding data from individuals meeting high-risk criteria after randomisation.

Secondary analyses: other secondary outcomes

Other maternal and neonatal binary outcomes will be analysed using the approaches described for the primary analysis for the ITT population and approaches detailed for the secondary analysis for the high-risk (HR) cohort. A similar process with generalised linear models employing an appropriate link function will be used to analyse the outcomes of maternal and neonatal initial hospital length of stay. Binary safety outcomes will also be analysed using the approaches detailed for the primary analysis. These analyses will be conducted using the as-treated population.

Sample size for primary outcome

Sample size estimates were generated to evaluate the potential benefits of peripartum prophylactic azithromycin in two population cohorts of women (table 3). The first population comprises all women delivering in facilities.

Power calculations for the overall study population were generated for the primary maternal and neonatal outcome measures. For each, estimates of the required sample size needed to detect a risk reduction of 20%, 25% and 30% were generated for power of 0.8, 0.85 and 0.9. The risk of sepsis or maternal death was assumed to be 3%, based on data collected from 2010 to 2018 through the Global

**Table 3** (A) Sample size for the overall population (alpha=0.05). (B) Sample size for high-risk group (alpha=0.05). (C) Sample size by region (alpha=0.025)

Baseline risk	Risk reduction	Evaluable sample size per arm		
		Power=0.80	Power=0.85	Power=0.90
(A)				
3%	20%	11 455	13 103	15 334
3%	25%	7133	8159	9548
3%	30%	4815	5508	6446
8%	20%	4096	4686	5483
8%	25%	2554	2921	3419
8%	30%	1727	1975	2311
14%	20%	2204	2521	2950
14%	25%	1377	1575	1842
14%	30%	932	1066	1247
(B)				
6%	20%	5568	6369	7453
6%	25%	3470	3969	4644
6%	30%	2344	2681	3138
(C)				
8%	20%	4961	5607	6477
8%	25%	3093	3496	4038
8%	30%	2091	2363	2729
14%	20%	2669	3017	3485
14%	25%	1677	1884	2176
14%	30%	1129	1276	1474

Network's Maternal Newborn Health Registry, a prospective, population-based registry of pregnant women and neonates receiving care in defined Global Network catchment areas,²⁷ augmented with active surveillance. For the neonatal outcome, the underlying risk of the combined outcome was estimated to be between 8% and 14% based on Global Network data.³⁵ We assumed that the risk of sepsis not resulting in death is approximately 4%–10%. We will also test the neonatal risk separately in south Asia and sub-Saharan Africa. Approximately 37.5% of randomised mothers will be from sub-Saharan Africa and 50% will be from Asia, reflecting the rates observed in the Global Network.^{35 36}

For the primary neonatal outcome of interest of intrapartum/neonatal sepsis or death, assuming that the loss to follow-up will be in the 2%–3% range, this sample size of 34 000 will provide 90% power to detect a 25% reduction in neonatal mortality and sepsis in the sub-Saharan African region and will provide 90% power to detect a 20% reduction in Asia assuming the baseline risk is at least 8%. For the primary maternal outcome of maternal death or sepsis, the sample size will provide 90% power to detect a 20% reduction from 3% in the population aggregated across all study sites. Each site will aim to recruit an equal number of participants (n=4250 per site) but site-specific sample

sizes may be adjusted if targets are not met. No site will be permitted to recruit more than 20% of the overall study sample site.

Data monitoring plan and stopping rules

Each site will report data, including adverse events, to the Global Network's DCC. The data will be used to evaluate protocol adherence and site performance (eg, recruitment, loss to follow-up, data quality). The DCC will provide standardised progress reports to NICHD and the site investigators monthly to monitor outcome variables and adverse events.

Trial oversight will be handled by two principal groups: (1) a protocol-focused steering committee (table 4) and (2) a DMC, designated by NICHD to ensure safe and ethical treatment of participants through biannual (at minimum) review of data on participant safety, study progress and futility. One formal interim analysis of efficacy and futility will be conducted by the DMC during the study.

Adverse event monitoring plan

Adverse events will be reported and submitted to the DCC (and IRBs) who will report these cumulative masked data to the DMC in the biannual safety reviews. Safety reports will be reviewed internally by the DCC quarterly and the DMC chair will be notified if any potential safety signals

Table 4 Azithromycin-Prevention in Labor Use Study (A-PLUS) steering committee

A-PLUS central study team	
A-PLUS lead study site (University of Alabama–Birmingham/University Teaching Hospital, Lusaka, Zambia)	
Waldemar A Carlo, MD Principal investigator University of Alabama at Birmingham, USA	Alan Tita, MD, PhD A-PLUS lead investigator University of Alabama at Birmingham, USA
Elwyn Chomba, MBChB, DCH, MRCP Senior foreign investigator University Teaching Hospital, Lusaka, Zambia	Musaku Mwenechanya Country coordinator University Teaching Hospital, Lusaka, Zambia
Trecious Mweemba A-PLUS coordinator University Teaching Hospital, Lusaka, Zambia	
RTI International (Global Network Data Coordinating Center)	
Elizabeth M McClure, PhD Principal investigator RTI International	Tracy Nolen, DrPh Coprincipal investigator and senior statistician RTI International
Jennifer J Hemingway-Foday, MPH, MSW A-PLUS protocol manager RTI International	
Eunice Kennedy Shriver National Institute of Child Health and Human Development	
Marion Koso-Thomas, MD, MPH Medical officer Global Network for Women's and Children's Health	
A-PLUS Research sites	
Democratic Republic of Congo (Kinshasa School of Public Health/University of North Carolina)	
Antoinette Tshetu, MD, PhD, MPH Senior foreign investigator Kinshasa School of Public Health, Kinshasa, Democratic Republic of Congo	Carl L Bose, MD Principal investigator University of North Carolina School of Medicine, USA
Adrien Lokangaka, MD, MPH Country coordinator Kinshasa School of Public Health	Gustave Lomendje A-PLUS coordinator Kinshasa School of Public Health
Guatemala (Instituto de Nutricion de Centro America y Panama (INCAP)/University of Colorado)	
Manolo Mazariegos, MD, MPH Senior foreign investigator INCAP, Guatemala City, Guatemala	Nancy F Krebs, MD Principal investigator University of Colorado Health Science Center
Lester Figueroa Country coordinator INCAP, Guatemala City, Guatemala	
Bangladesh (International Centre for Diarrhoeal Disease Research(ICDDR,b)/University of Virginia)	
Rashidul Haque, MD Senior foreign investigator ICDDR,b	William Petri, MD Principal investigator University of Virginia
Sk Masum Billah Country coordinator ICDDR,b	Md Shahjahan Siraj A-PLUS coordinator ICDDR,b
Belagavi, India (KLE University's JN Medical College/Thomas Jefferson University)	
Shivaprasad S Goudar MD, MHPE Senior foreign investigator KLE University's J N Medical College	Richard Derman, MD, MPH Principal investigator Thomas Jefferson University
Avinash Kavi A-PLUS coordinator KLE University's J N Medical College	Mrityunjay Metgud A-PLUS coordinator KLE University's J N Medical College
Pakistan (Aga Khan University/Columbia University)	
Sarah Saleem, MD Senior foreign investigator Aga Khan University	Robert L Goldenberg, MD Principal investigator Columbia University
Shiyam Sunder Tikmani A-PLUS coordinator Aga Khan University	

Continued

**Table 4** Continued**A-PLUS Research sites**

Nagpur, India (Lata Medical Research Foundation/Boston University)	
Archana Patel, MD, DNB, MSCE Senior foreign investigator Lata Medical Research Foundation	Patricia L Hibberd, MD, PhD Principal investigator Boston University School of Public Health
Prabir Das Country coordinator Lata Medical Research Foundation	Kunal Kurhe A-PLUS coordinator Lata Medical Research Foundation
Kenya (Moi University School of Medicine/University of Indiana School of Medicine)	
Fabian Esamai, MBChB, MMed, PhD Senior foreign investigator Moi University School of Medicine, Eldoret, Kenya	Edward A Liechty, MD Principal investigator Indiana University School of Medicine
Paul Nyongesa Country coordinator Moi University School of Medicine, Eldoret, Kenya	Amos Sagwe A-PLUS coordinator Moi University School of Medicine, Eldoret, Kenya
Osayame Austine Ekhaguere Coinvestigator Indiana University School of Medicine	

are identified to allow for more frequent DMC monitoring if needed.

ETHICS AND DISSEMINATION**Ethics approval and consent to participate**

The A-PLUS study protocol has been reviewed and approved by the relevant ethics committees and regulatory authorities at each research site, including the institutional review board at the University of Alabama at Birmingham, which serves as the lead site, and the DCC at RTI International (see online supplemental file 1 for current V.1.6, dated 13 July 2022). The DMC has reviewed the study protocol and will continue to review throughout the enrollment period. The study was registered (clinicaltrials.gov NCT03871491).

All research staff responsible for obtaining informed consent will be trained and certified in the protection of human subjects and the study-specific consent procedures. A model written informed consent form, developed according to the requirements of the US Office for Human Research Protection (OHRP), can be found as online supplemental appendix 2 of the trial protocol. The model consent may be modified by each site to conform to local standards, but the OHRP required elements must be maintained.

Potential risks and benefits to participation

There are several potential direct and indirect benefits of this trial. Emerging data suggest that intrapartum azithromycin reduces maternal and neonatal infection. As infections are a frequent cause of maternal and neonatal deaths, there is a possibility that mortality could be reduced at the participating sites, as well as worldwide.

An ongoing concern for peripartum and perinatal antibiotic prophylaxis is the selection of resistant organisms including azithromycin-resistant organisms

leading to resistant infections, and concern that disruption of gut and other flora (microbiome) in women and particularly in neonates may lead to adverse events including allergic reactions, rash and childhood asthma.^{16 28 37} There is a paucity of data to address concerns that disturbances in the establishment of the indigenous intestinal microbiome caused by antibiotic exposure in early life or CD, either directly or through modifications of breast microbiome, may increase risk of immune-mediated and inflammatory conditions later in life.^{38–41} In response to these global public health concerns, this protocol includes an ancillary study to monitor for antimicrobial resistance and changes to the maternal and newborn microbiome.

Dissemination

Following completion of primary data collection, we will disseminate findings through international meetings and high impact peer-reviewed journals. Per the National Institutes of Health (NIH) and Global Network Data Management and Sharing Policies, final research data will also be made publicly available through the NICHD Data and Specimen Hub system within 1 year of publishing the primary manuscript. All data will be deidentified and shared under the assurance of confidentiality and approval from the relevant IRBs. Trial results will also be shared with community members, health workers, health officials and other stakeholders from the areas where participants were recruited. If the results are positive, we will facilitate practice change at participating sites and approach the WHO to evaluate guideline updates based on study findings.

Author affiliations

¹RTI International, Research Triangle Park, North Carolina, USA

- ²The University of Alabama at Birmingham School of Medicine, Birmingham, Alabama, USA
- ³University of Zambia, University Teaching Hospital, Lusaka, Zambia
- ⁴University of Kinshasa, Kinshasa, Congo (the Democratic Republic of the)
- ⁵Kinshasa School of Public Health, Kinshasa, Congo (the Democratic Republic of the)
- ⁶Boston University School of Public Health, Boston, Massachusetts, USA
- ⁷Lata Medical Research Foundation, Nagpur, Maharashtra, India
- ⁸Datta Meghe Institute of Higher Education & Research (Deemed to be University), Wardha, Maharashtra, India
- ⁹KLE Academy of Higher Education and Research, Jawaharlal Nehru Medical College, Belgavi, Karnataka, India
- ¹⁰Community Health Sciences, The Aga Khan University, Karachi, Pakistan
- ¹¹Moi University School of Medicine, Eldoret, Kenya
- ¹²Instituto de Nutrición de Centroamérica y Panamá, Guatemala, Guatemala
- ¹³The University of Sydney, Sydney, New South Wales, Australia
- ¹⁴International Centre for Diarrhoeal Disease Research Bangladesh, Dhaka, Bangladesh
- ¹⁵Columbia University, New York, New York, USA
- ¹⁶The University of North Carolina at Chapel Hill School of Medicine, Chapel Hill, North Carolina, USA
- ¹⁷Indiana University School of Medicine, Indianapolis, Indiana, USA
- ¹⁸University of Colorado School of Medicine, Denver, Colorado, USA
- ¹⁹Office of Global Affairs, Thomas Jefferson University, Philadelphia, Pennsylvania, USA
- ²⁰University of Virginia, Charlottesville, Virginia, USA
- ²¹Eunice Kennedy Shriver National Institute of Child Health and Human Development, Bethesda, Maryland, USA

Acknowledgements The authors would like to acknowledge the A-PLUS research teams for their many contributions to this work: Kinshasa School of Public Health, Kinshasa, Democratic Republic of Congo: Michel Kalonji and Miyalu Junior; University of North Carolina, Chapel Hill, North Carolina, USA: Jackie Patterson; University Teaching Hospital, Lusaka, Zambia: Ernest Banda, Mwansa Chimfwembe, Ruth Nakazwe and Abigail Mwapule; Instituto de Nutrición de Centroamérica y Panamá, Guatemala City, Guatemala: Maynor Manrique; University of Colorado School of Medicine, Denver, CO, USA: Jamie Westcott; International Centre for Diarrhoeal Disease Research, Dhaka, Bangladesh: Qazi Sadeq-ur-Rahman, Amita Farzana, Farhana Jahan, Zarin Tasnim Maliha and Lolit Singh; University of Virginia, Charlottesville, Virginia, USA: Chris Chisolm; KLE Academy Higher Education and Research, J N Medical College Belagavi, Karnataka, India: Sheetal Harakuni, Manjunath Somannavar and Kadappa Beniawadi; Thomas Jefferson University, Philadelphia, Pennsylvania, USA: Frances Jaeger; Aga Khan University, Karachi, Pakistan: Zaheer Habib, Imran Ahmed, Farnaz Naqvi, Najja Ghanchi and Saleem Jessani; Lata Medical Research Foundation, Nagpur, India: Vaishali Khedikar, Savita Bhargav, Samreen Sadaf and Chaitali Gedam; Moi University School of Medicine, Eldoret, Kenya: Milsort Kemboi, Anderson Misati and Kevin Otieno; Indiana School of Medicine, University of Indiana, Indianapolis, Indiana, USA: Sheri Bucher; RTI International, Research Triangle Park, North Carolina, USA: Anna Aceituno, Kay Jackson, Jean Kim, Janet Moore, Suchita Parepalli, Marissa Trotta and Alexis Williams. Bill & Melinda Gates Foundation—Laura Lamberti.

Contributors AT and WAC conceived of the study and developed the protocol, with input from EMC, TN, JH-F, MK-T and PLH. TN developed the statistical analyses plan. JH-F wrote the first draft of the manuscript and subsequent revisions, with critical feedback from WAC, AT, EMC, MK-T and TN. EC, MMwenechanya, TM, AL, ATK, GL, PLH, AP, PKD, KK, SSG, AK, MMetgud, SS, SST, FE, PN, AS, LF, MMazariegos, SMB, RH, MSS, RG, MB, CB, EAL, OAE, NFK, RD and WAP contributed to the refinement and finalisation of the study protocol and trial implementation. All authors contributed to the preparation of this manuscript and have reviewed and approved the final version.

Funding The A-PLUS study is supported through institutional grants from the Eunice Kennedy Shriver National Institute of Child Health and Human Development (NICHD) (RTI International (U01 HD040636), University of North Carolina at Chapel Hill (U10 HD076465), University of Alabama at Birmingham (U10 HD078437), University of Colorado (U10 HD076474), Thomas Jefferson University (U10 HD076457), Columbia University (U10 HD078438), Boston University (U10 HD078439), Indiana University (U10 HD076461)) and a grant from the Foundation for the National Institutes of Health (MCCL19APT) through the Maternal, Newborn & Child Health Discovery & Tools initiative of the Bill & Melinda Gates Foundation (BMGF) (INV-008973). The views expressed in this manuscript are those of the

authors and do not necessarily represent the views of the NICHD; the National Institutes of Health; or the US Department of Health and Human Services or the BMGF.

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.

Provenance and peer review Not commissioned; externally peer reviewed.

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

Jennifer Hemingway-Foday <http://orcid.org/0000-0001-9776-5306>
 Avinash Kavi <http://orcid.org/0000-0002-2176-4697>
 Sarah Saleem <http://orcid.org/0000-0002-6797-8631>
 Paul Nyongesa <http://orcid.org/0000-0002-2896-4720>
 Sk Masum Billah <http://orcid.org/0000-0002-8690-6932>
 Md Shahjahan Siraj <http://orcid.org/0000-0001-6925-324X>
 Carl Bose <http://orcid.org/0000-0002-8727-8640>
 Osayame A Ekhaquere <http://orcid.org/0000-0002-8926-3709>
 Elizabeth McClure <http://orcid.org/0000-0001-8659-5444>

REFERENCES

- World Health Organization. WHO recommendations for the prevention and treatment of maternal peripartum infections; 2015. Available: <https://www.who.int/publications/i/item/9789241549363> [Accessed 22 Aug 2018].
- Liu L, Oza S, Hogan D, *et al*. Global, regional, and national causes of child mortality in 2000–13, with projections to inform post-2015 priorities: an updated systematic analysis. *Lancet* 2015;385:430–40.
- Fleischmann-Struzek C, Goldfarb DM, Schlattmann P, *et al*. The global burden of paediatric and neonatal sepsis: a systematic review. *Lancet Respir Med* 2018;6:223–30.
- Fleischmann C, Reichert F, Cassini A, *et al*. Global incidence and mortality of neonatal sepsis: a systematic review and meta-analysis. *Arch Dis Child* 2021;106:745–52.
- Baguiya A, Bonet M, Cecatti JG, *et al*. Perinatal outcomes among births to women with infection during pregnancy. *Arch Dis Child* 2021;106:946–53.
- Brizuela V, Cuesta C, Bartolelli G, *et al*. Availability of facility resources and services and infection-related maternal outcomes in the WHO global maternal sepsis study: a cross-sectional study. *Lancet Glob Health* 2021;9:e1252–61.
- World Health Organization. Statement on maternal sepsis. 2017. Available: http://www.who.int/reproductivehealth/publications/maternal_perinatal_health/maternalsepsis-statement/en/ [Accessed 22 Aug 2018].
- African Neonatal Sepsis Trial (AFRINEST) group, Tshefu A, Lokangaka A, *et al*. Simplified antibiotic regimens compared with injectable procaine benzylpenicillin plus gentamicin for treatment of neonates and young infants with clinical signs of possible serious bacterial infection when referral is not possible: a randomised, open-label, equivalence trial. *Lancet* 2015;385:1767–76.
- Baqui AH, Saha SK, Ahmed A, *et al*. Safety and efficacy of alternative antibiotic regimens compared with 7 day injectable procaine benzylpenicillin and gentamicin for outpatient treatment of neonates and young infants with clinical signs of severe infection when referral is not possible: a randomised, open-label, equivalence trial. *Lancet Glob Health* 2015;3:e279–87.



- 10 Mir F, Nisar I, Tikmani SS, *et al.* Simplified antibiotic regimens for treatment of clinical severe infection in the outpatient setting when referral is not possible for young infants in Pakistan (simplified antibiotic therapy trial [SATT]): a randomised, open-label, equivalence trial. *Lancet Glob Health* 2017;5:e177–85.
- 11 Zaidi AKM, Tikmani SS, Sultana S, *et al.* Simplified antibiotic regimens for the management of clinically diagnosed severe infections in newborns and young infants in first-level facilities in Karachi, Pakistan: study design for an outpatient randomized controlled equivalence trial. *Pediatr Infect Dis J* 2013;32 Suppl 1:S19–25.
- 12 World Health Organization. WHO statement on Caesarean section rates. 2015. Available: http://www.who.int/reproductivehealth/publications/maternal_perinatal_health/cs-statement/en/ [Accessed 22 Aug 2018].
- 13 Mackeen AD, Packard RE, Ota E, *et al.* Timing of intravenous prophylactic antibiotics for preventing postpartum infectious morbidity in women undergoing cesarean delivery. *Cochrane Database Syst Rev* 2014;12:CD009516.
- 14 Small FM, Grivell RM. Antibiotic prophylaxis versus no prophylaxis for preventing infection after cesarean section. *Cochrane Database Syst Rev* 2014;2014:CD007482.
- 15 Gyte GM, Dou L, Vazquez JC, *et al.* Different classes of antibiotics given to women routinely for preventing infection at caesarean section. *Cochrane Database Syst Rev* 2014;11:CD008726.
- 16 Tita ATN, Rouse DJ, Blackwell S, *et al.* Evolving concepts in antibiotic prophylaxis for cesarean delivery: a systematic review. *Obstet Gynecol* 2009;113:675–82.
- 17 Harrison MS, Saleem S, Ali S, *et al.* A prospective, population-based study of trends in operative vaginal delivery compared to cesarean delivery rates in low- and middle-income countries, 2010–2016. *Am J Perinatol* 2019;36:730–6.
- 18 Tita ATN, Szychowski JM, Boggess K, *et al.* Adjunctive azithromycin prophylaxis for cesarean delivery. *N Engl J Med* 2016;375:1231–41.
- 19 Oluwalana C, Camara B, Bottomley C, *et al.* Azithromycin in labor lowers clinical infections in mothers and newborns: a double-blind trial. *Pediatrics* 2017;139:e20162281.
- 20 Hume-Nixon M, Quach A, Reyburn R, *et al.* A systematic review and meta-analysis of the effect of administration of azithromycin during pregnancy on perinatal and neonatal outcomes. *EClinicalMedicine* 2021;40:101123.
- 21 Hallamaa L, Cheung YB, Maleta K, *et al.* Child health outcomes after presumptive infection treatment in pregnant women: a randomized trial. *Pediatrics* 2018;141:e20172459.
- 22 Fitzgibbon A, Clooney L, Broderick D, *et al.* Erythromycin compared to amoxicillin and azithromycin for antimicrobial prophylaxis for preterm premature rupture of the membranes: a retrospective study. *J Obstet Gynaecol* 2021;41:569–72.
- 23 Sutton AL, Acosta EP, Larson KB, *et al.* Perinatal pharmacokinetics of azithromycin for cesarean prophylaxis. *Am J Obstet Gynecol* 2015;212:812.
- 24 Russell NJ, Seale AC, O'Driscoll M, *et al.* Maternal Colonization with group B Streptococcus and Serotype distribution worldwide: systematic review and meta-analyses. *Clin Infect Dis* 2017;65:S100–11.
- 25 Rac H, Gould AP, Eiland LS, *et al.* Common bacterial and viral infections: review of management in the pregnant patient. *Ann Pharmacother* 2019;53:639–51.
- 26 Harper LM, Kilgore M, Szychowski JM, *et al.* Economic evaluation of adjunctive azithromycin prophylaxis for cesarean delivery. *Obstet Gynecol* 2017;130:328–34.
- 27 McClure EM, Garces AL, Hibberd PL, *et al.* The global network maternal newborn health registry: a multi-country, community-based registry of pregnancy outcomes. *Reprod Health* 2020;17:184.
- 28 Chan A-W, Tetzlaff JM, Gøtzsche PC, *et al.* SPIRIT 2013 explanation and elaboration: guidance for protocols of clinical trials. *BMJ* 2013;346:e7586.
- 29 Bonet M, Souza JP, Abalos E, *et al.* The global maternal sepsis study and awareness campaign (GLOSS): study protocol. *Reprod Health* 2018;15:16.
- 30 Albright CM, Has P, Rouse DJ, *et al.* Internal validation of the sepsis in obstetrics score to identify risk of morbidity from sepsis in pregnancy. *Obstetric Anesthesia Digest* 2018;38:8.
- 31 Bowyer L, Robinson HL, Barrett H, *et al.* SOMANZ guidelines for the investigation and management sepsis in pregnancy. *Aust N Z J Obstet Gynaecol* 2017;57:540–51.
- 32 World Health Organization. Guideline: managing possible serious bacterial infection in young infants when referral is not feasible. 2015. Available: https://apps.who.int/iris/bitstream/handle/10665/181426/9789241509268_eng.pdf [Accessed 18 Aug 2020].
- 33 Pasha A, McClure EM, Saleem S, *et al.* A prospective cause of death classification system for maternal deaths in low and middle-income countries: results from the global network maternal newborn health registry. *BJOG* 2018;125:1137–43.
- 34 Garces AL, McClure EM, Pérez W, *et al.* The global network neonatal cause of death algorithm for low-resource settings. *Acta Paediatr* 2017;106:904–11.
- 35 Goudar SS, Goco N, Somannavar MS, *et al.* Institutional deliveries and stillbirth and neonatal mortality in the global network's maternal and newborn health registry. *Reprod Health* 2020;17:179.
- 36 Hoffman MK, Goudar SS, Kodkany BS, *et al.* A description of the methods of the aspirin supplementation for pregnancy indicated risk reduction in nulliparas (ASPIRIN) study. *BMC Pregnancy Childbirth* 2017;17:135.
- 37 Roca A, Oluwalana C, Bojang A, *et al.* Oral azithromycin given during labour decreases bacterial carriage in the mothers and their offspring: a double-blind randomized trial. *Clin Microbiol Infect* 2016;22:565.
- 38 Roberts S, Maccato M, Faro S, *et al.* The Microbiology of post-cesarean wound morbidity. *Obstet Gynecol* 1993;81:383–6.
- 39 Rosene K, Eschenbach DA, Tompkins LS, *et al.* Polymicrobial early postpartum endometritis with facultative and anaerobic bacteria, genital Mycoplasmas, and Chlamydia trachomatis: treatment with piperacillin or cefoxitin. *J Infect Dis* 1986;153:1028–37.
- 40 Andrews WW, Shah SR, Goldenberg RL, *et al.* Association of post-cesarean delivery endometritis with colonization of the chorioamnion by ureaplasma urealyticum. *Obstet Gynecol* 1995;85:509–14.
- 41 Keski-Nisula L, Kirkinen P, Katila ML, *et al.* Amniotic fluid U. Urealyticum colonization: significance for maternal peripartur infections at term. *Am J Perinatol* 1997;14:151–6.