



HAL
open science

Severe bacterial neonatal infections in Madagascar, Senegal, and Cambodia: A multicentric community-based cohort study

Bich-Tram Huynh, Elsa Kermorvant-Duchemin, Rattanak Chheang, Frederique Randrianirina, Abdoulaye Seck, Elisoa Hariniaina Ratsima, Zafitsara Zo Andrianirina, Jean-Baptiste Diouf, Armya Youssef Abdou, Sophie Goyet, et al.

► To cite this version:

Bich-Tram Huynh, Elsa Kermorvant-Duchemin, Rattanak Chheang, Frederique Randrianirina, Abdoulaye Seck, et al.. Severe bacterial neonatal infections in Madagascar, Senegal, and Cambodia: A multicentric community-based cohort study. PLoS Medicine, 2021, 18 (9), pp.e1003681. 10.1371/journal.pmed.1003681 . pasteur-03357830

HAL Id: pasteur-03357830

<https://pasteur.hal.science/pasteur-03357830v1>

Submitted on 29 Sep 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License

RESEARCH ARTICLE

Severe bacterial neonatal infections in Madagascar, Senegal, and Cambodia: A multicentric community-based cohort study

Bich-Tram Huynh^{1,2*}, Elsa Kermorvant-Duchemin³, Rattanak Chheang⁴, Frederique Randrianirina⁵, Abdoulaye Seck⁶, Elisoa Hariniaina Ratsima⁵, Zafitsara Zo Andrianirina⁷, Jean-Baptiste Diouf⁸, Armya Youssouf Abdou^{1,2}, Sophie Goyet⁹, Véronique Ngo⁹, Siyin Lach⁹, Long Pring⁹, Touch Sok¹⁰, Michael Padget^{1,2}, Fatoumata Diene Sarr¹¹, Laurence Borand⁹, Benoit Garin^{12^{aa}}, Jean-Marc Collard^{12^{ab}}, Perlinot Herindrainy^{13^c}, Agathe de Lauzanne^{9^c}, Muriel Vray^{11^{ac}}, Elisabeth Delarocque-Astagneau^{1,14}, Didier Guillemot^{1,2,14}, On behalf of the BIRDY study group[†]

1 Université Paris-Saclay, UVSQ, Inserm, CESP, Anti-infective Evasion and Pharmacoepidemiology Team, Montigny-Le-Bretonneux, France, **2** Institut Pasteur, Epidemiology and Modelling of Antibiotic Evasion (EMAE), Paris, France, **3** AP-HP, Hôpital Necker-Enfants Malades, Department of Neonatal Medicine, Université de Paris, Paris, France, **4** Medical Biology Unit, Institut Pasteur du Cambodge, Phnom Penh, Cambodia, **5** Centre de Biologie Clinique, Institut Pasteur de Madagascar, Antananarivo, Madagascar, **6** Laboratory of Medical Biology, Institut Pasteur de Dakar, Dakar, Senegal, **7** Pediatric Ward, Centre Hospitalier de Soavinandriana, Antananarivo, Madagascar, **8** Centre Hospitalier Roi Baudouin, Guédiawaye, Dakar, Sénégal, **9** Epidemiology and Public Health Unit, Institut Pasteur du Cambodge, Phnom Penh, Cambodia, **10** Ministry of Health, Phnom Penh, Cambodia, **11** Epidemiology of Infectious Diseases Unit, Institut Pasteur de Dakar, Dakar, Senegal, **12** Experimental Bacteriology Unit, Institut Pasteur de Madagascar, Antananarivo, Madagascar, **13** Epidemiology Unit, Institut Pasteur de Madagascar, Antananarivo, Madagascar, **14** AP-HP, Paris Saclay, Public Health, Medical Information, Clinical Research, Le Kremlin-Bicêtre, France

[☞] These authors contributed equally to this work.

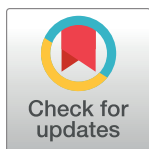
^{aa} Current address: CHU de Pointe-à-Pitre/les Abymes, Guadeloupe

^{ab} Current address: Experimental Bacteriology Unit, Institut Pasteur of Shanghai/Chinese Academy of Sciences, Shanghai, China

^{ac} Current address: Epidemiology of Emergent Diseases Unit, Institut Pasteur, Inserm, Paris, France

[†] Membership of the BIRDY study group is provided in the Acknowledgments.

* bich-tram.huynh@pasteur.fr



OPEN ACCESS

Citation: Huynh B-T, Kermorvant-Duchemin E, Chheang R, Randrianirina F, Seck A, Hariniaina Ratsima E, et al. (2021) Severe bacterial neonatal infections in Madagascar, Senegal, and Cambodia: A multicentric community-based cohort study. *PLoS Med* 18(9): e1003681. <https://doi.org/10.1371/journal.pmed.1003681>

Academic Editor: Quique Bassat, Instituto de Salud Global de Barcelona, SPAIN

Received: January 4, 2021

Accepted: June 1, 2021

Published: September 28, 2021

Copyright: © 2021 Huynh et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data Availability Statement: Data cannot be shared publicly because data privacy is subject to French, Cambodian, Senegalese and Malagasy regulations. Data are available from the Ethics Committee (contact via irb@pasteur.fr) for researchers who meet the criteria for access to confidential data.

Funding: This work was supported by the Department of International Cooperation of the Principality of Monaco (<https://en.gouv.mc/>)

Abstract

Background

Severe bacterial infections (SBIs) are a leading cause of neonatal deaths in low- and middle-income countries (LMICs). However, most data came from hospitals, which do not include neonates who did not seek care or were treated outside the hospital. Studies from the community are scarce, and few among those available were conducted with high-quality microbiological techniques. The burden of SBI at the community level is therefore largely unknown. We aimed here to describe the incidence, etiology, risk factors, and antibiotic resistance profiles of community-acquired neonatal SBI in 3 LMICs.

Methods and findings

The BIRDY study is a prospective multicentric community-based mother and child cohort study and was conducted in both urban and rural areas in Madagascar (2012 to 2018),

Government-Institutions/The-Government/Ministry-of-Foreign-Affairs-and-Cooperation/Department-of-International-Cooperation), MSDAVENIR (<http://www.msdavenir.fr/>) and Total foundation (<https://www.foundation.total.fr/>). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing interests: The authors have declared that no competing interests exist.

Abbreviations: aHR, adjusted hazard ratio; ARB, antibiotic-resistant bacteria; aTR, adjusted time ratio; CHAMPS, Child Health and Mortality Prevention Surveillance; CI, confidence interval; CoNS, coagulase-negative staphylococci; cSBI, confirmed severe bacterial infection; CSF, cerebrospinal fluid; ESBL, extended-spectrum β -lactamase; GBS, group B streptococcus; HR, hazard ratio; LBW, low birth weight; LMIC, low- and middle-income country; MALDI-TOF, matrix-assisted laser desorption/ionization time-of-flight; MCAR, missing completely at random; pSBI, possible severe bacterial infection; SBI, severe bacterial infection; STROBE, Strengthening the Reporting of Observational Studies in Epidemiology; VLBW, very low birth weight.

Cambodia (2014 to 2018), and Senegal (2014 to 2018). All pregnant women within a geographically defined population were identified and enrolled. Their neonates were actively followed from birth to 28 days to document all episodes of SBI. A total of 3,858 pregnant women (2,273 (58.9%) in Madagascar, 814 (21.1%) in Cambodia, and 771 (20.0%) in Senegal) were enrolled in the study, and, of these, 31.2% were primigravidae. Women enrolled in the urban sites represented 39.6% (900/2,273), 45.5% (370/814), and 61.9% (477/771), and those enrolled in the rural sites represented 60.4% (1,373/2,273), 54.5% (444/814), and 38.1% (294/771) of the total in Madagascar, Cambodia, and Senegal, respectively. Among the 3,688 recruited newborns, 49.6% were male and 8.7% were low birth weight (LBW). The incidence of possible severe bacterial infection (pSBI; clinical diagnosis based on WHO guidelines of the Integrated Management of Childhood Illness) was 196.3 [95% confidence interval (CI) 176.5 to 218.2], 110.1 [88.3 to 137.3], and 78.3 [59.5 to 103] per 1,000 live births in Madagascar, Cambodia, and Senegal, respectively. The incidence of pSBI differed between urban and rural sites in all study countries. In Madagascar, we estimated an incidence of 161.0 pSBI per 1,000 live births [133.5 to 194] in the urban site and 219.0 [192.6 to 249.1] pSBI per 1,000 live births in the rural site ($p = 0.008$). In Cambodia, estimated incidences were 141.1 [105.4 to 189.0] and 85.3 [61.0 to 119.4] pSBI per 1,000 live births in urban and rural sites, respectively ($p = 0.025$), while in Senegal, we estimated 103.6 [76.0 to 141.2] pSBI and 41.5 [23.0 to 75.0] pSBI per 1,000 live births in urban and rural sites, respectively ($p = 0.006$). The incidences of culture-confirmed SBI were 15.2 [10.6 to 21.8], 6.5 [2.7 to 15.6], and 10.2 [4.8 to 21.3] per 1,000 live births in Madagascar, Cambodia, and Senegal, respectively, with no difference between urban and rural sites in each country. The great majority of early-onset infections occurred during the first 3 days of life (72.7%). The 3 main pathogens isolated were *Klebsiella* spp. (11/45, 24.4%), *Escherichia coli* (10/45, 22.2%), and *Staphylococcus* spp. (11/45, 24.4%). Among the 13 gram-positive isolates, 5 were resistant to gentamicin, and, among the 29 gram-negative isolates, 13 were resistant to gentamicin, with only 1 *E. coli* out of 10 sensitive to ampicillin. Almost one-third of the isolates were resistant to both first-line drugs recommended for the management of neonatal sepsis (ampicillin and gentamicin). Overall, 38 deaths occurred among neonates with SBI (possible and culture-confirmed SBI together). LBW and foul-smelling amniotic fluid at delivery were common risk factors for early pSBI in all 3 countries. A main limitation of the study was the lack of samples from a significant proportion of infants with pSBI including 35 neonatal deaths. Without these samples, bacterial infection and resistance profiles could not be confirmed.

Conclusions

In this study, we observed a high incidence of neonatal SBI, particularly in the first 3 days of life, in the community of 3 LMICs. The current treatment for the management of neonatal infection is hindered by antimicrobial resistance. Our findings suggest that microbiological diagnosis of SBI remains a challenge in these settings and support more research on causes of neonatal death and the implementation of early interventions (e.g., follow-up of at-risk newborns during the first days of life) to decrease the burden of neonatal SBI and associated mortality and help achieve Sustainable Development Goal 3.

Author summary

Why was this study done?

- Severe bacterial infections (SBIs) are a leading cause of neonatal deaths in low- and middle-income countries (LMICs).
- In LMICs, most data come from studies conducted in hospitals. However, a significant proportion of neonates do not have access to the hospital or die before reaching it in these countries.
- The magnitude of SBI and its causative pathogens, consequences, and risk factors at the community level remain largely unknown in LMICs.

What did the researchers do and find?

- We conducted a prospective multicentric, rural and urban, community-based mother and child cohort study in Madagascar, Cambodia, and Senegal. All pregnant women within a geographically defined population were identified and invited to participate. Their neonates were actively followed from birth to 28 days.
- Among the 3,688 recruited newborns, 514 neonates presented an episode of SBI, with the great majority occurring during the first 3 days of life (54.7%). The incidence of possible severe bacterial infection (pSBI) varied from 78.3 [95% confidence interval [CI] 59.5 to 103] to 196.3 [176.5 to 218.2] cases per 1,000 live births and was the highest in Madagascar; the incidence of confirmed severe bacterial infection (cSBI) varied from 6.5 [2.7 to 15.6] to 15.2 [10.6 to 21.8] per 1,000 live births.
- The 3 main pathogens isolated were *Klebsiella* spp. (24.4%), *Escherichia coli* (22.2%), and *Staphylococcus* spp. (26.7%). Almost one-third of the isolates (31%) were resistant to both first-line drugs recommended by WHO for the management of neonatal sepsis.

What do these findings mean?

- Our findings show the high burden of neonatal SBI, particularly in the first 3 days of life, in the community of 3 LMICs in Asia and Africa.
- These findings also suggest that the current treatment strategy for the management of neonatal infection may be hindered by antimicrobial resistance.
- Our findings support the implementation of early interventions (e.g., follow-up of at-risk newborns during the first days of life) to decrease the burden of severe neonatal infection and mortality and help achieve Sustainable Development Goal 3.

Introduction

Neonatal mortality accounts for 40% of the global under-5 mortality, with the great majority occurring in low- and middle-income countries (LMICs) [1]. Severe infections are a leading

cause of neonatal deaths (>75%), with an estimate of 6.9 million cases of possible severe bacterial infection (pSBI) occurring yearly in these countries [2,3].

LMICs also combine a number of risk factors for the emergence and dissemination of antibiotic-resistant bacteria (ARB) leading to potential treatment failure and increased neonatal mortality [4].

Data on the burden and risk factors of neonatal severe bacterial infections (SBIs) and ARB infections are scarce in LMICs, particularly in Africa and Asia. Data that do exist are mainly provided by hospital-based investigations, which can miss a large proportion of newborns who did not seek care or were treated outside the hospital and for which microbiological information is not available [5,6]. Only few studies have been conducted at the community level with reliable clinical and microbiology data on SBI in these countries [5,7]. These gaps in knowledge prevent the implementation of simplified treatment of SBI in the community when referral is not possible as well as the improvement of prevention and treatment interventions of neonatal SBI at the community level [8].

Investigating community-acquired neonatal SBI in LMICs is extremely challenging regarding the specificities of the context. To address these challenges, we implemented a prospective cohort study including exhaustive community recruitment and active follow-up of pregnant women and their neonates in 3 LMICs. Here, we report the incidence, etiology, risk factors, and antibiotic resistance profiles of community-acquired neonatal SBI in Madagascar, Cambodia, and Senegal.

Methods

Study areas and design

The full protocol is described in detail in the Supporting information ([S1 Text](#)). In Madagascar (2012 to 2018), the study was implemented in 3 urban districts of Antananarivo (population: 14,997) and in the rural city of Moramanga (population: 17,159). In Cambodia (2014 to 2018), the study was conducted in 2 areas of Steung Meanchey, an urban district of Phnom Penh (population: 87,035) and 2 districts of rural Kampong Speu (79,000 inhabitants). In Senegal (2014 to 2018), the study took place in a district of Guédiawaye, a city near urban Dakar (population: 20,529), and Sokone, a rural city close to the Gambia border (population: 14,500).

Study recruitment included the exhaustive identification of pregnant women within the study area with the help of community healthcare workers who identified pregnant women eligible for inclusion. The study area was defined based on the expected numbers of live births and a target number of inclusions. Participants were enrolled during their third trimester of pregnancy and were actively monitored to include their newborn at birth. In addition, we optimized the exhaustiveness of live birth recruitment by also including neonates at the time of birth, without a prior inclusion phase of the mother if she was not identified during her pregnancy.

Inclusion criteria for pregnant women included routine residence in the study area with no plan to move away during the follow-up period and no opposition to the research being conducted or to the collection of biological samples.

At delivery, neonates who met the following criteria were enrolled: neonates born to parents living in the study area with no plan to move during the follow-up period; those whose legal guardians were informed and had no objection to the study procedures and collection of biological samples; and those for whom written consent was obtained from at least 1 legal guardian. Data collected were the mothers' sociodemographic, medical, and obstetric characteristics; delivery information; risk factors of infection at delivery; and the neonates' anthropometric measurements and Apgar scores.

After delivery, we performed both active and passive surveillance during the neonatal period to detect all symptoms of severe infection [9]. Surveillance visits were scheduled for the third day after delivery and then weekly until the first month of life. Passive surveillance consisted of asking the family to contact the investigators if the newborn presented symptoms suggestive of infection. All cases of suspected infections were investigated by a physician. Biological samples (C-reactive protein and complete blood count) as well as samples for bacterial identification including blood cultures were performed according to study guidelines and the physician's discretion. This study is reported as per the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) guideline ([S1 Checklist](#)).

Sample size

The sample size was determined in part by available funding, which was different for each country; we aimed to enroll 750, 800, and 2,000 newborns in Senegal, Cambodia, and Madagascar, respectively.

The primary objective of the BIRDY study was to estimate the incidence of SBI at the community level. At the time of study planning, published neonatal infection incidence ranged from 6.3 to 23 per 1,000 live births, depending on the study [10–14]. In addition, WHO considered that all-cause neonatal mortality rates to be underestimated by at least 20% [15]. Given that about 36% of these deaths are attributable to severe infections, we hypothesized that this underestimation of the number of deaths probably resulted in a similar underestimation of the incidence of bacterial infections [16]. After correction for this underestimation and assuming that the number of neonates at risk remained constant over the period considered, we could expect, with the targeted recruitment number, maximal incidences, by site, of between 8 per thousand (95% confidence interval (CI) [2.5 to 13.5]) and 28 per thousand (95% CI [17.6 to 37.4]) per 1,000 live births.

Bacteriologic analyses

All samples were immediately stored at a temperature between 0 and 4°C and transported within hours to the clinical laboratory at the Institut Pasteur of each country. All analyses were performed according to the guidelines from the French Society for Microbiology ([S1 Text](#)) [17]. For blood culture, fresh blood agar and chocolate media were used for bacterial isolation, and bacterial strains from the cultures were identified using API Gallery or matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometry. Antimicrobial susceptibility testing was done by disk diffusion in accordance with the French Society for Microbiology guidelines [17]. A double-disk synergy test was systematically done in all Enterobacteriaceae resistant to third-generation cephalosporin to identify extended-spectrum β -lactamase (ESBL) producers. *Escherichia coli* ATCC 25922 was used for quality control strains.

Case ascertainment

The neonatal period was defined from birth to 28 days (i.e., day 0 to 28) of life.

We defined pSBI cases as the presence of at least 1 sign among severe chest indrawing, tachypnea, hypo- or hyperthermia, lethargy, convulsions and poor feeding, and no pSBI nor hospitalization within the previous 7 days. Neonates presenting tachypnea as the only clinical sign were not counted as cases because of the low specificity of this sign alone [7].

We defined confirmed severe bacterial infection (cSBI) as presence of at least 1 sign cited above and a positive culture from blood or cerebrospinal fluid (CSF) or urine (bacterial and leukocyte counts $>10^5$ and 10^4 , respectively) or pus from umbilicus/subcutaneous abscess in case of soft tissue or omphalitis-associated sepsis.

All cases of infection were reviewed by an epidemiologist (BTH), a neonatologist (EKD), and a microbiologist (BG or JMC) to classify them as cSBI or pSBI and to exclude nonsevere cases or clinically nonsignificant contaminant.

Early-onset infections were defined as cases occurring between 0 and 6 days and late-onset infections as cases occurring between 7 and 28 days. The time when day 0 became day 1 was set at midnight.

Multidrug-resistant infections were those caused by pathogens resistant to >1 agent in >3 antibacterial categories [18].

Statistical analyses

Incidence estimation was planned a priori. The analysis plan for risk factors determination of SBI was developed prospectively, with later adjustments, such as running a Cox model with a stratification for covariate violating the proportional hazards assumption according to reviewers' suggestions, and no data-driven changes took place to the analysis plan.

Incidence rates of cSBI and pSBI were expressed as neonatal infections per 1,000 live births.

Maternal characteristics and obstetrical and neonatal outcomes were compared between countries by the use of Fisher exact or χ^2 tests for proportions and *t* tests for mean values. Very low birth weight (VLBW) and low birth weight (LBW) were defined as a [birth weight](#) of less than 1,500 g and less than 2,500 g, regardless of [gestational age](#). Dystocic delivery is defined as [abnormal labor](#) resulting from abnormalities of the power (uterine contractions or maternal expulsive forces), the passenger (position, size, or presentation of the fetus), or the passage (pelvis or soft tissues).

We investigated in each country if LBW and factors related to delivery (foul-smelling amniotic fluid, premature rupture of membranes, dystocic delivery, and cesarean section) (exposures of primary interest) were associated with early pSBI, as it represented the majority of the overall burden and considered site, sex of the newborns, and maternal and pregnancy characteristics as potential confounding factors. We first conducted a univariate analysis to describe the relationship between the outcome and the exposures of primary interest and potential confounding factors. Then, we performed a multivariable analysis to present adjusted analysis for the exposures of interest. In univariate analysis, the associations with risk of early pSBI were estimated using Cox proportional hazards models and expressed as hazard ratios (HRs) with 95% CI. The proportional hazards assumption was tested using Schoenfeld residuals for each independent variable and outcome pair.

For multivariable analysis, factors associated with the outcome were entered in the initial model if *p*-value was less than 0.20 in univariate analysis. A stepwise backward procedure was then performed to identify factors independently associated with the outcome (*p* < 0.05). If a variable violated the proportional hazards assumption, this variable was considered as a stratification factor in the Cox proportional hazard model. To document the direction and the strength of the variable that violated the proportional hazards assumption, an additional parametric regression model with an accelerated failure time approach and a log-normal distribution was used.

Two variables were highly correlated: foul-smelling amniotic fluid and premature rupture of the membranes; we therefore kept the variable that was the most associated with the considered outcome (foul-smelling amniotic fluid) to avoid multicollinearity in multivariable analysis.

In Senegal, 14% of neonates had missing values for variables related to risk factors at delivery (foul-smelling amniotic fluid, premature rupture of membranes, and dystocic delivery), mainly for logistic reasons. Consequently, missing data were classified as missing completely at random (MCAR). We performed complete case analyses that provide unbiased estimates

under MCAR assumption and also made sensitivity analyses incorporating worst- and best-case scenarios [19,20]. In the worst scenario, infants were classified as infants born to mothers with foul-smelling amniotic fluid and/or premature rupture of membranes and/or dystocic delivery depending on which variable was missing. In the best scenario, infants were classified as infants born to mother with no foul-smelling amniotic fluid and/or no premature rupture of membranes and/or who had a eutocic delivery.

All analyses were performed using Stata version 15 (Stata, College Station, Texas, United States of America). All *p*-values were from 2-sided tests, and results were deemed statistically significant at $p < 0.05$.

Ethical statement

The study was approved by the ethics committees of Madagascar (068-MSANP/CE), Senegal (SEN 14–20), Cambodia (108 NEHCR), and the Institutional Review Board of Institut Pasteur (IRB/2016/08/03), France. Written informed consent was given by all participants.

Results

A total of 3,858 pregnant women were enrolled in the study: 2,273 (58.9%) in Madagascar, 814 (21.1%) in Cambodia, and 771 (20%) in Senegal (Fig 1). Women enrolled in the urban sites represented 39.6%, 45.5%, and 61.9% of the total in Madagascar, Cambodia, and Senegal, respectively. Maternal and neonatal characteristics by country are presented in Table 1. A total of 3,688 (95.6%) babies were born in the frame of the study. Among the 3,688 recruited newborns, 49.6% were male, and 75.6% were born in a healthcare facility. There were 10.6% (221/2,078), 4.1% (33/792), and 9.6% (68/705) LBW in Madagascar, Cambodia, and Senegal, respectively. Among them, 50% (108/216), 39.4% (13/33), and 12.7% (7/55) were born prematurely (before 37 weeks gestation) in Madagascar, Cambodia, and Senegal, respectively. There were 73 infants (2%) lost to follow-up (42 in Madagascar, 2 in Cambodia, and 29 in Senegal). In total, 3,414 (92.6%) newborns (1,953 in Madagascar, 776 in Cambodia, and 671 in Senegal) had a complete follow-up during the neonatal period.

Samples

A total of 514 cases of SBI (42 culture confirmed and 472 pSBI) were identified. Among these, 247 had at least 1 microbiological sample (215 blood cultures, 101 urine tests, and 26 lumbar punctures). The proportion of cases with no microbiological sample taken was 50% (186/372), 58.3% (49/84), and 55.2% (32/58) in Madagascar, Cambodia, and Senegal, respectively (Fig 2). The proportion of unexplored pSBI was high in the rural site in the 3 countries (53.6%, 68.4%, and 61.5% in Madagascar, Cambodia, and Senegal, respectively), although significantly different only in Madagascar when compared to the urban site ($p = 0.048$). Among SBI cases, those who experienced dystocic delivery were less likely to have a microbiological sample taken compared to those born without delivery complication (60.56% versus 47.64% ($p = 0.05$)) in Madagascar. Also, neonates with SBI who needed resuscitation at birth were less likely to be sampled compared to those who did not need resuscitation in Senegal (85.7% versus 47.9% ($p = 0.06$)), although this difference was not statistically significant.

Incidence of severe bacterial neonatal infections

pSBIs. The overall incidence of pSBI was 196.3 per 1,000 live births [95% CI = 176.5 to 218.2], 110.1 [88.3 to 137.3], and 78.3 [59.5 to 103] in Madagascar, Cambodia, and Senegal, respectively.

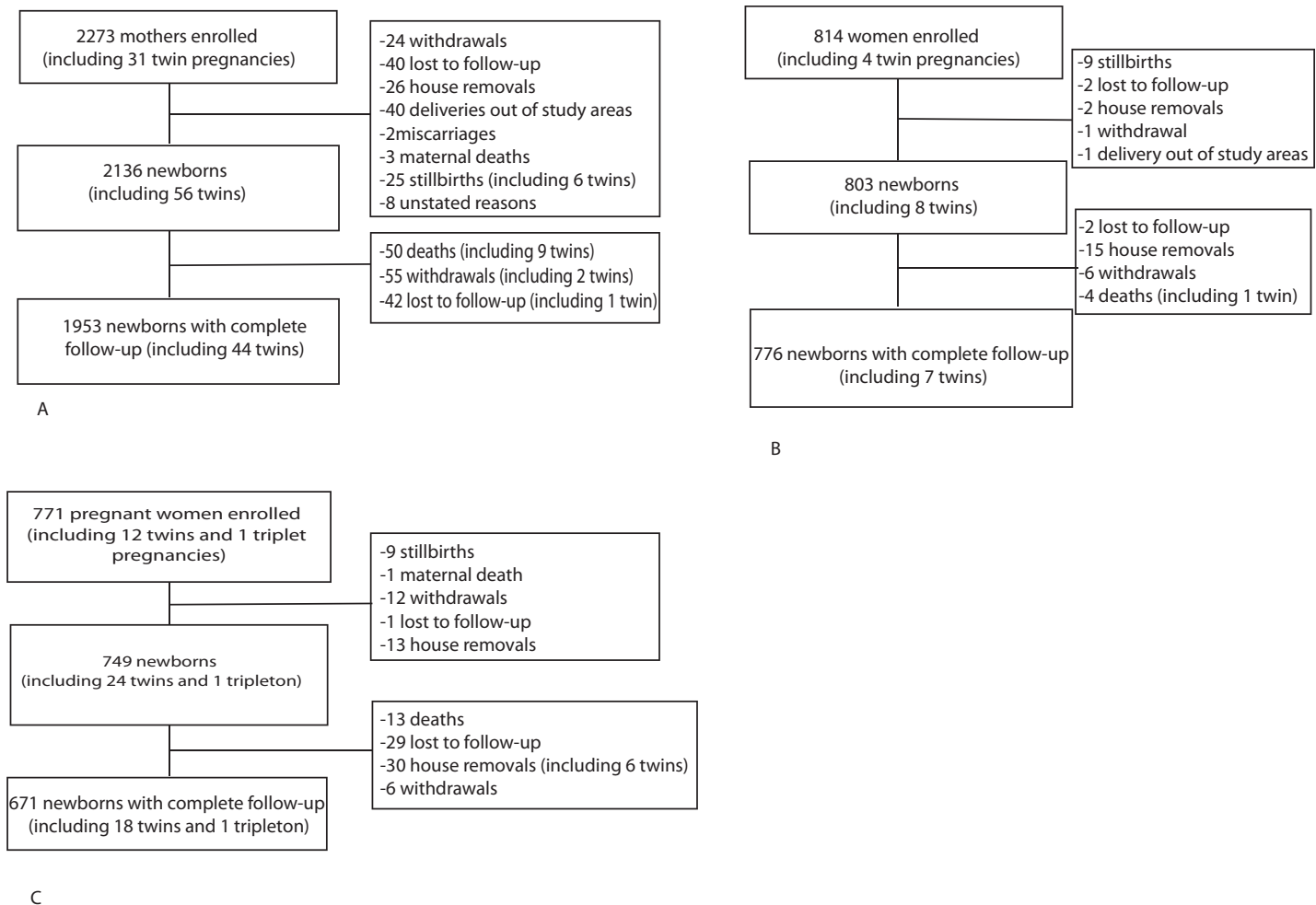


Fig 1. Flowchart of the study population. (A) Madagascar, (B) Cambodia, and (C) Senegal.

<https://doi.org/10.1371/journal.pmed.1003681.g001>

The incidence of pSBI differed between urban and rural sites in all study countries. In Madagascar, we estimated an incidence of 161.0 pSBI per 1,000 live births [133.5 to 194.0] in the urban site and 219.0 [192.6 to 249.1] pSBI per 1,000 live births in the rural site ($p = 0.008$). In Cambodia, estimated incidences were 141.1 [105.4 to 189.0] and 85.3 [61.0 to 119.4] pSBI per 1,000 live births in urban and rural sites, respectively ($p = 0.025$), while in Senegal, we estimated 103.6 [76.0 to 141.2] pSBI and 41.5 [23.0 to 75.0] pSBI per 1,000 live births in urban and rural sites, respectively ($p = 0.006$).

The early pSBI incidences were estimated at 115.9 per 1,000 live births [101 to 133], 59.9 [44.4 to 80.8], and 63 [46.3 to 85.5] in Madagascar, Cambodia, and Senegal, respectively. The incidence of late pSBI ranged from 15.4 [8.3 to 28.5] in Senegal to 80.9 [68.6 to 95.4] per 1,000 live births in Madagascar (Fig 3, S1 Table).

cSBIs. A total of 42 neonates were classified as having culture-confirmed severe infection ($n = 30$ in Madagascar, $n = 5$ in Cambodia, and $n = 7$ in Senegal). The overall incidence rates of cSBI per 1,000 live births were 15.2 [10.6 to 21.8], 6.5 [2.7 to 15.6], and 10.2 [4.8 to 21.3] in Madagascar, Cambodia, and Senegal, respectively, with no difference between the 3 countries and between urban and rural sites in each country (S1 Table).

Table 1. Characteristics of the mothers and their neonates.

	Madagascar	Cambodia	Senegal	P
Mothers	N = 2,273	N = 814	N = 771	
Urban site	39.6 (900/2,273)	45.5 (370/814)	62.6 (477/762)	<0.001
Age, median (SD) [range]	25 (6.5) [14–48]	27.0 (5.4) [14–46]	27.9 (6.5) [15–49]	0.20
Education				
Absence/primary school	24.8 (564/2,273)	54.7 (445/814)	73.5 (560/762)	<0.001
Partial secondary school	52.3 (1,188/2,273)	31.8 (259/814)	18.0 (137/762)	
Complete secondary or higher	22.9 (521/2,273)	13.5 (110/814)	8.5 (65/762)	
Electricity	75.4 (1,714/2,273)	99.5 (809/813)	97.1 (740/762)	<0.001
Latrine				
Inside the house	7.3 (166/2,273)	50.3 (409/813)	51.1 (387/757)	<0.001
Outside the house	92.7 (2,106/2,273)	49.7 (405/813)	48.9 (370/757)	
Household members median [range]	4 [2–14]	5 [2–21]	10 [2–21]	0.03
Primigravidae	32.8 (746/2,273)	33.8 (275/814)	23.5 (179/762)	<0.001
Twin pregnancy	1.4 (31/2,273)	0.4 (4/814)	1.6 (13/771)	0.07
At least 1 antenatal visit at enrollment	97.6 (2,218/2,273)	98.6 (761/772)	99.3 (722/727)	<0.001
Skilled birth attendant	72.2 (1,528/2,117)	99.4 (787/791)	95.3 (726/762)	<0.001
Hospitalization during pregnancy	2.6 (54/2,098)	3.2 (26/812)	3.0 (21/687)	0.45
Babies	N = 2,136	N = 803	N = 749	
Male	50.7 (1,080/2,128)	48.0 (381/794)	50.2 (376/749)	0.29
Birth weight, median (SD) [range]	2,969.1 (469) [1,000–4,900]	3,104.1 (437.3) [1,500–4,600]	3,049.6 (497) [1,000–5,000]	0.10
LBW^a	10.6 (221/2,078)	4.1 (33/792)	9.6 (68/705)	<0.001
VLBW^b	0.7 (15/2,078)	0 (0/792)	0.2 (2/705)	0.02
Preterm^c	15.5 (329/2,121)	2.2 (17/777)	4.5 (31/689)	<0.001
Delivery in healthcare facilities	60.2 (1,276/2,119)	98.6 (782/793)	98.5 (733/744)	<0.001
Cesarean section	9.3 (198/2,135)	12.5 (99/794)	3.4 (25/739)	<0.001
Premature rupture of the membranes	2.5 (54/2,120)	0.5 (4/769)	4.3 (27/630) ^d	<0.001
Fetid amniotic fluid	7.8 (166/2,119)	2.1 (16/769)	18.3 (115/630) ^d	<0.001
Dystocic delivery	8.5 (181/2,119)	4.6 (35/769)	2.2 (14/630) ^d	<0.001
Exclusive breastfeeding at day 7	89 (1,826/2,051)	77.1 (586/760)	94.8 (688/726)	0.03

^a LBW is defined as a birth weight < 2,500 g.

^b VLBW is defined as a birth weight < 1,500 g.

^c Birth before 37 weeks gestation.

^d In Senegal, data related to delivery (premature rupture of the membranes, fetid amniotic fluid, and dystocic delivery) are for 630 neonates among 749 due to missing data.

Data are % (n/N) unless stated otherwise.

LBW, low birth weight; VLBW, very low birth weight.

<https://doi.org/10.1371/journal.pmed.1003681.t001>

The early-onset severe neonatal infections rates ranged from 3.9 [1.3 to 12.1] in Cambodia to 12.3 [8.2 to 18.3] per 1,000 live births in Madagascar, and late-onset infection rates ranged from 1.4 [0.2 to 10.3] in Senegal to 3.1 [1.4 to 6.8] per 1,000 live births in Madagascar.

The great majority of early-onset infections occurred during the first 3 days of life (16/24 in Madagascar, 3/3 in Cambodia, and 5/6 in Senegal).

In the 3 countries, all estimates of pSBI incidences largely surpassed those of cSBI: 12.9, 16.9, and 7.7 times greater in Madagascar, Cambodia, and Senegal, respectively. Among the 514 episodes of SBI, 54.7% occurred during the first 3 days (Fig 4).

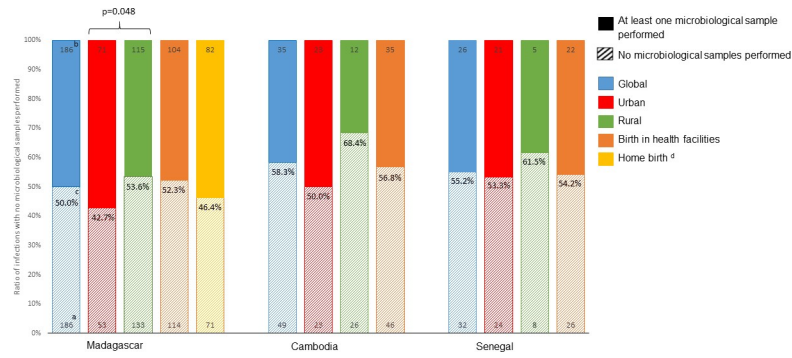


Fig 2. Ratio of SBI without microbiological samples performed. Total SBI include both culture-confirmed SBI and pSBI. (a) Numbers at the bottom of each bar represent the number of neonatal SBI with no microbiological samples performed. (b) Numbers at the top of each bar represent the number of neonatal SBI with at least 1 microbiological sample performed among blood culture, urine test, and lumbar puncture. (c) Percentages in the middle of each bar represent the ratio of neonatal SBI with no microbiological samples performed (number of neonatal SBI without microbiological samples performed/ overall number of neonatal SBI). (d) Only in Madagascar (>98% of the pregnant women delivered in healthcare facilities in Cambodia and Senegal). pSBI, possible severe bacterial infection; SBI, severe bacterial infection.

<https://doi.org/10.1371/journal.pmed.1003681.g002>

Pathogens isolated and antibacterial resistance

Pathogens. Among the 42 neonates with cSBI, 45 pathogens were identified including 3 samples in which 2 pathogens were isolated (S2 Table). The 3 main pathogens isolated were *Klebsiella* spp. (11/45, 24.4%) *E. coli* (10/45, 22.2%), and *Staphylococcus* spp. (11/45, 24.4%).

Overall, we identified 5 cases of meningitis and 37 cases of bacteremia/sepsis including 7 secondary to urinary tract infection, 5 secondary to omphalitis, and 1 secondary to a muscular abscess.

Antibacterial resistance. Among the 45 bacteria isolated, 42 were tested for antimicrobial drug susceptibility (29/32 gram-negative and 13/13 gram-positive isolates) (S3 Table). More than half of isolates with resistance data (23/42, 54.8%) were resistant to at least 1 of the first-line drugs recommended for the management of neonatal sepsis (ampicillin and gentamicin), with 13/42 (31%) resistant to both drugs (8 *Klebsiella* spp., 4 *E. coli*, and 1 *Enterobacter cloacae*).

Overall, 18 isolates were identified as multidrug resistant (44.8% (13/29), 20% (1/5), and 50% (4/8) in Madagascar, Cambodia, and Senegal, respectively), which led to an estimation of

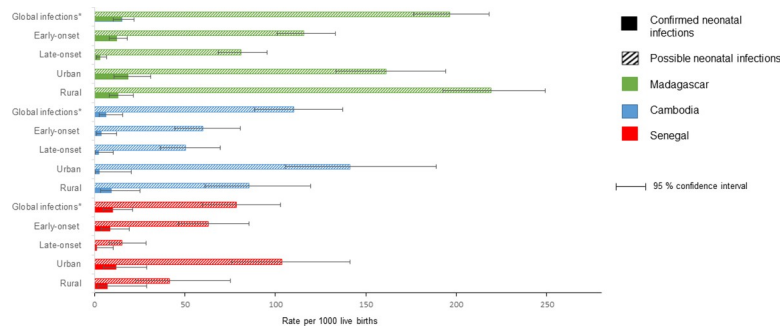


Fig 3. Incidence of severe neonatal infections (per 1,000 live births). Early-onset infections are defined as cases occurring between 0 and 6 days and late-onset infections as cases occurring between 7 and 28 days.* Global infections incidence is defined as early-onset and late-onset infections together.

<https://doi.org/10.1371/journal.pmed.1003681.g003>

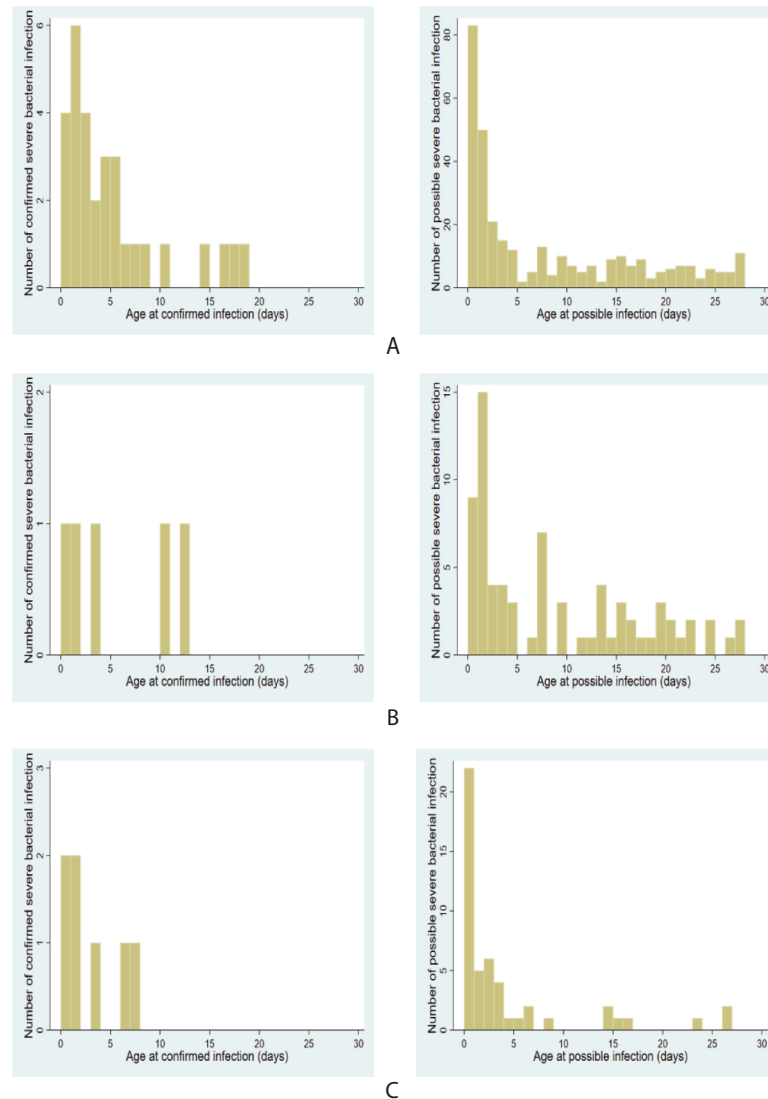


Fig 4. Timing of cSBI and pSBI among neonates. (A) Madagascar, (B) Cambodia, and (C) Senegal. cSBI, confirmed severe bacterial infection; pSBI, possible severe bacterial infection.

<https://doi.org/10.1371/journal.pmed.1003681.g004>

the overall incidence rate for multidrug-resistant severe neonatal infection of 5.0 [3.1 to 8.0] cases per 1,000 live births.

Almost half of the gram-negative isolates were resistant to cefotaxime (48%, 12/25) and to gentamicin (46.4%, 13/28), and more than one-third showed resistance to ciprofloxacin (34.5%, 10/29). No resistance was detected toward imipenem. Moreover, 11 Enterobacteriaceae were ESBL producers: 8 in Madagascar and 3 in Cambodia.

Among the 5 *Staphylococcus aureus* and the 5 coagulase-negative staphylococci (CoNS) isolated, 1 and 2 strains were resistant to methicillin, respectively.

Clinical presentation and outcomes

Among the 472 neonates with pSBI, the most prevalent clinical signs were fever (51%), feeding difficulties (28.2%), lethargy (17.5%), irritability (12.7%), and difficult breathing (13.5%). For

neonates presenting cSBI, fever (61.9%), feeding difficulties (31.8%), irritability (31.8%), lethargy (22.7%), difficult breathing (15.9%), prolonged capillary refill (>3 seconds) (11.9%), and cyanosis (9.1%) were the most frequent clinical signs.

There were 3 deaths among neonates with cSBI caused by *E. cloacae* at day 15 in Senegal, *Klebsiella pneumoniae* at day 5, and *Acinetobacter baumannii* at day 20 in Madagascar. All 3 deaths occurred among hospital births and were due to resistant bacteria that required imipenem.

Among neonates with pSBI, 26, 2, and 7 deaths occurred in Madagascar, Cambodia, and Senegal, respectively. Among these neonates, infection fatality ratio of early-onset infection was 11.4% (23/201), [CI = 7.4 to 16.7%] in Madagascar, 4.7% (2/43), [CI = 0.6 to 15.8%] in Cambodia, and 4.9% (2/41), [CI = 0.6 to 16.5%] in Senegal, with no statistically significant difference between the 3 countries ($p = 0.22$). Infection fatality ratio of late-onset infection was 2.1% (3/141), [CI = 0.4 to 6%] in Madagascar and 50% (5/10), [CI = 18.7 to 81.3%] in Senegal; no death occurred among late-onset pSBI in Cambodia.

Factors associated with early pSBI

We found in all 3 countries that neonates with foul-smelling amniotic fluid at delivery (adjusted hazard ratio (aHR) = 2.2 [1.71 to 3.4], $p < 0.001$ in Madagascar, aHR = 3.94 [2.05 to 7.59], $p < 0.001$ in Senegal, and 5.5 [2.1 to 14.4], $p = 0.001$ in Cambodia) presented a higher risk of early pSBI (Table 2). Also, LBW babies (aHR = 4.52; 95% CI [3.36 to 6.1], $p < 0.001$ in Madagascar, aHR = 2.48 [1.03 to 5.95], $p = 0.04$ in Senegal) were at increased risk of early pSBI compared to normal weight babies (Table 2). In Cambodia, the variable “LBW” violated the proportional hazards assumption and was considered as a stratification variable. To document the direction and the strength of “LBW” in Cambodia, a parametric regression model with an accelerated failure time approach and a log-normal distribution were used (S4 Table). In the adjusted analysis, in neonates with LBW, time to onset of SBI was about 10 times shorter than in neonates with normal birth weight (adjusted time ratio (aTR): 0.11 [0.05 to 0.24], $p < 0.001$).

Complete case analyses and sensitivity analyses (worst- and best-case scenarios) yielded comparable results except for LBW, which was not significantly associated with early pSBI in the worst-case scenario in Senegal (S5 Table).

Discussion

Our results showed high incidences of SBI among neonates in 3 different LMICs in Asia and Africa using a prospective community-based approach. A great majority of SBI occurred during the first 3 days of life and rates of antibiotic resistance to recommended first-line treatments were also high, raising questions about current recommendations for the management of neonatal sepsis (ampicillin and gentamicin).

We observed a relatively high estimation of pSBI incidence, up to more than 200 cases per 1,000 live births. A recent community-based cohort with a similar design was conducted in Pakistan, India, and Bangladesh from 2011 to 2014 by Saha and colleagues [7]. However, the authors included cases of severe infections occurring until 59 days of life, whereas we considered cases during the first 28 days of life [21]. The authors found an estimation of pSBI ranging from 78.3 to 112.6 episodes per 1,000 live births. If we take into account the different follow-up time between the 2 studies, our estimation of pSBI incidence is higher, particularly in Madagascar.

Saha and colleagues performed extensive molecular diagnosis and could attribute a cause in only 28% of the episodes (16% bacterial and 12% viral). It is therefore also likely that in our

Table 2. Risk factors analysis of possible severe bacterial neonatal infections.

	Madagascar N = 2,050				Senegal N = 630				Cambodia ^d N = 769			
	Univariate analysis		Multivariable analysis		Univariate analysis		Multivariable analysis		Univariate analysis		Multivariable analysis	
	HR [95% CI]	p	Adjusted HR ^a [95% CI]	p	HR [95% CI]	p	aHR ^b [95% CI]	p	HR [95% CI]	p	aHR ^c [95% CI]	p
Site (urban site as reference)	0.9 [0.75–1.3]	0.88			0.58 [0.29–1.14]	0.12			0.8 [0.46–1.45]	0.48		
Education												
Absence/primary school	Ref				Ref				Ref			
Partial secondary school	0.81 [0.58–1.11]	0.19			0.82 [0.31–2.12]	0.68			0.83 [0.44–1.58]	0.57		
Complete secondary or higher	1.1 [0.77–1.58]	0.59			1.44 [0.56–3.74]	0.45			0.55 [0.19–1.56]	0.26		
Primigravidae	1.36 [1.03–1.79]	0.03			0.94 [0.43–2.04]	0.87			1.5 [0.86–2.76]	0.14		
Twins pregnancy	3.99 [2.5–6.4]	0.001			- ^e				7.03 [1.7–29]	0.007		
Hospitalization during pregnancy	2.02 [1.1–3.7]	0.02			0.87 [0.12–6.3]	0.89			0.7 [0.1–5.1]	0.72		
Skilled birth attendant	0.95 [0.7–1.31]	0.77			0.6 [0.08–4.3]	0.61			- ^f			
Sex of newborn (boys as reference)	0.72 [0.55–0.94]	0.02			0.9 [0.5–1.7]	0.80			0.8 [0.45–1.4]	0.45		
Delivery out of healthcare facilities	0.8 [0.6–1.04]	0.10			- ^e				- ^f			
LBW	4.35 [3.28–5.8]	0.001	4.52 [3.36–6.1]	<0.001	2.39 [1.0–5.7]	0.05	2.48 [1.03–5.95]	0.04	- ^d			
Cesarean section	1.65 [1.14–2.4]	0.001			1.7 [0.23–12.4]	0.53			1.34 [0.6–3]	0.48		
Fetid amniotic fluid	3.2 [2.3–4.44]	0.001	2.2 [1.71–3.4]	<0.001	3.78 [1.99–7.16]	0.001	3.94 [2.05–7.6]	<0.001	6.01 [2.38–15.2]	<0.001	5.5 [2.1–14.4]	0.001
Dystocic delivery	4.15 [3.08–5.58]	0.001	3.2 [2.38–4.5]	<0.001	2.8 [0.67–11.6]	0.16			2.2 [0.77–6.04]	0.14		

^a Multivariable analysis in Madagascar: adjusted on site (rural versus urban) (1.2 [0.88–1.55], $p = 0.28$) and sex of the newborn (0.65 [0.49–0.86], $p = 0.002$).

^b Multivariable analysis in Senegal: adjusted on site (rural versus urban) (0.51 [0.25–1.01], $p = 0.05$) and sex of the newborn (1.06 [0.55–2.01], $p = 0.87$).

^c Multivariable analysis in Cambodia: adjusted on site (rural versus urban) (0.91 [0.5–1.65], $p = 0.75$) and sex of the newborn (0.7 [0.38–1.26], $p = 0.23$).

^d For analysis in Cambodia: LBW is incorporated in the model as a stratification factor.

^e No possible estimation as no cases of pSBI occurred among twins and one case among neonates who delivered out of healthcare facilities.

^f No possible estimation as no cases of pSBI occurred among neonates who delivered out of healthcare facilities and among those who delivered with unqualified healthcare workers.

aHR, adjusted hazard ratio; CI, confidence interval; HR, hazard ratio; LBW, low birth weight; pSBI, possible severe bacterial infection.

<https://doi.org/10.1371/journal.pmed.1003681.t002>

study a significant proportion of the episodes were not of bacterial origin or of infectious origin as signs and symptoms of severe infection in neonates are subtle and nonspecific with some clinical signs of severe infection overlapping with noninfectious severe events (e.g., intrapartum related complication, prematurity . . .) [22].

Our estimates of cSBI, ranging from 6.5 to 15.2 cases per 1,000 live births, is 6 to 15 times higher than in high-income countries (<1 per 1,000 live births in the USA) [23]. Although the definition of the neonatal period differed between the 2 studies, our results are in line with estimates of Saha and colleagues, which ranged from 10.61 to 23.46 cases per 1,000 live births.

Taking into account the challenges of addressing this research question in LMICs, our methodology provided reliable estimates of both pSBI and cSBI, with the true incidence of SBI probably falling between these two. These figures underline the high burden of SBI in settings where SBI management is challenged by limited provision of basic supportive care and respiratory support and the lack of skilled staff.

Our findings also suggest that the first week and especially the first 3 days of life are critical with the majority of infections occurring during that period [7,24]. In South Asia, Saha and colleagues showed that incidence of pSBI was higher in the first 3 days with an increase in incidence density rate of roughly 13.5 times. Thus, this finding reinforces the need for interventions targeting the perinatal period.

In concordance with previous studies conducted in LMICs, we found that the most predominant bacteria responsible for neonatal infections were *Klebsiella* spp., *E. coli*, and *Staphylococcus* spp. [7,11,13,14,25–27]. Among the latter, a significant proportion of CoNS were identified in our study as a causative organism for early-onset infections with strains sensitive to vancomycin and teicoplanin unlike high-income countries where CoNs are responsible for late-onset sepsis acquired in hospital settings and resistant to vancomycin and teicoplanin. Our results suggest that neonates might also get infected due to unhygienic conditions during labor or nonoptimal cord care and not solely in nosocomial environments. These findings reinforce the importance of hygienic measures during delivery and for perinatal care. In Madagascar, particularly, where one-third of the pregnant women delivered at home, the use of clean birth kits should be encouraged. We did not find as Saha and colleagues that *Ureaplasma* spp. was a predominant causative bacteria as we did not culture them. Indeed, *Ureaplasma* spp. do not grow on standard media. Also, we did not perform molecular diagnosis. Further investigation including systematic testing of *Ureaplasma* spp. in neonates with pSBI are needed to confirm its burden compared to the others bacteria.

We did not identify any group B streptococcus (GBS) culture-confirmed infections, although our methodology maximized the chances to document such infections, which often occur at the early stage of the neonatal period. GBS infection might have occurred among the 35 deaths with pSBI, since only 50% of neonates with pSBI underwent microbiological investigation. Indeed, Saha and colleagues showed that the attributed proportion of GBS for pSBI was higher among babies who died than among survivors. Also, Taylor and colleagues recently conducted a study in 7 African countries aiming at determining causes of under-5 mortality with minimally invasive tissue sampling (Child Health and Mortality Prevention Surveillance (CHAMPS) study) [27]. This innovative methodology allowed the authors to capture blood samples after early death, and their preliminary results showed that GBS was a major causative bacteria among neonates with sepsis who died, along with *A. baumannii* (probably mostly hospital acquired), *K. pneumoniae*, and *E. coli*. Our findings highlight the challenge in estimating the burden of GBS as infected neonates might deteriorate very quickly in LMIC settings. This question is all the more important to investigate as a vaccine for GBS carriage is currently under development.

One important result of our study is that almost one-third of the overall isolates were resistant to both drugs recommended by WHO to manage neonatal sepsis (ampicillin and gentamicin). All these infections were caused by gram-negative bacilli, particularly *Klebsiella* spp., and almost half of the isolates also showed resistance to third-generation cephalosporins. No carbapenem resistance was identified.

Susceptibility of causative bacteria to drugs recommended by WHO is unclear with some studies showing results consistent with ours, whereas other show moderate resistance [7,11,27,28]. These findings suggest that the situation might be heterogeneous depending on the country, and revised recommendations should be adapted locally. Importantly, we found a

relatively low incidence of multidrug-resistant infections (5 cases/1,000 live births) and did not identify carbapenemase-producing *Enterobacteriaceae*. These results suggest that antibiotic resistance to last-resort drugs is still limited in community settings in the study areas of these 3 countries, and it is still time for action to contain the spread of these strains.

One striking result of the study is the gap between pSBI and cSBI incidences, pSBI being 8 to 17 times more frequent than cSBI. This gap illustrates the challenge of microbiological diagnosis in LMIC settings. Indeed, despite a prospective design and freely available access to biological analyses, less than 50% of neonates with pSBI underwent microbiological investigation, which was at the attending clinician's discretion. Our results suggest that microbiological diagnosis is largely underused, particularly in rural areas and most likely in critically ill babies. Specifically, we observed that lumbar puncture was not widely practiced routinely, although the protocol specified indications of CSF sampling and despite extensive refresh trainings. These findings are worrisome for optimal management of severe neonatal infection in LMICs and strengthen the pressing need for the development of innovative rapid diagnostic tests easily usable at bedside in order to identify causative pathogens and associated susceptibility pattern. Also, this bottleneck is essential to overcome in order to provide high-quality data on cause and antimicrobial resistance to meet research and surveillance needs.

LBW and foul-smelling amniotic fluid at delivery were common risk factors for early pSBI in all 3 countries. LBW can be either due to intrauterine growth retardation or prematurity. The latter has been shown to increase the risk of developing a SBI due to immaturity of the immune system and of the skin and intestine epithelial barriers [28–30]. Foul-smelling amniotic fluid often reflects ascending maternal infection from colonizing pathogens of perineum or genitourinary tract, which might be transmitted to the neonates perinatally. This result is consistent with previous studies conducted in others LMICs [31]. These findings are important as these 2 clinical signs are easily detectable by nonskilled healthcare workers, and their surveillance could be part of an intervention to identify at-risk newborns during the perinatal period, especially in settings where access to health facilities is limited.

Our study draws its strength from its rigorous methodology designed for exhaustive identification of neonatal SBI in a geographic area along with appropriate and timely diagnosis to overcome challenges in investigating community-acquired neonatal SBI in 3 LMICs of Asia and Africa.

However, we acknowledge some limitations. First, despite intensive efforts, samples could not be collected from the 35 babies with pSBI who died, most of the time because the clinical situation was too critical to sample them. Therefore, we cannot confirm that these babies died from a bacterial infection. Moreover, more than half of these newborns were LBW, indicating that others factors, such as prematurity, may have contributed to these deaths. It is also likely that mortality due to SBI in the community is higher in reality than in research settings where facilitated access to healthcare facilities, adequate treatment, and active surveillance may impact outcomes. In this context, innovative approaches such as in the CHAMPS study are important to decipher causes of mortality [27].

Second, blood cultures in neonates are particularly challenging to draw, requiring adequate blood volume in aseptic conditions. In rural areas, particularly in Senegal (6 hours), transportation time of the samples to the laboratory might have reduced the bacterial yield, resulting in negative results. Although continuous trainings to ensure blood sampling quality were delivered throughout the study, including recommendations on adequate blood volume and aseptic sampling techniques, the incidence rate of cSBI may be underestimated in our study, as already mentioned above. Also, we cannot exclude that some gut pathogens on the baby's skin might have contaminated the blood cultures.

Finally, our findings might not be generalizable to others areas within the same country or in another LMIC. However, given the challenges in investigating neonatal sepsis at the community level in LMICs, our findings add important values to the scientific community and to policymakers of each country where data on neonatal sepsis are scarce.

This study has important policy and research implications. Our findings show the high burden of neonatal SBI in the community of 3 LMICs in Asia and Africa and also suggests that the current treatment strategy for the management of neonatal infection could be hindered by antimicrobial resistance. Our study shows that SBI is a critical issue but that even in a research setting, microbiological diagnosis of SBI remains a challenge in LMICs, impeding not only neonatal SBI management but also research and resistance surveillance. Also, our findings shows that the first 3 days are particularly vulnerable and support the implementation of early interventions (e.g., follow-up of at-risk newborns during the first days of life).

Conclusions

The BIRDY study uses an innovative community-based approach to inform knowledge on the burden of neonatal SBI in 3 LMICs in Asia and Africa. Results from this study will help to improve prevention and treatment interventions at the community level, to decrease neonatal severe infection and associated mortality, and to help achieve Sustainable Development Goal 3.

Supporting information

S1 Checklist. STROBE checklist. STROBE, Strengthening the Reporting of Observational Studies in Epidemiology.

(DOC)

S1 Text. Study protocol.

(DOCX)

S1 Table. Global (both early- and late-onset infections), early, and late incidences of culture-confirmed and possible severe neonatal infections according to site.

(DOCX)

S2 Table. Pathogens isolated in neonatal culture-confirmed severe infections.

(DOCX)

S3 Table. Susceptibility pattern of isolated bacteria in culture-confirmed neonatal infections.

(DOCX)

S4 Table. Risk factors analysis of possible severe bacterial neonatal infection in Cambodia with parametric regression model, with an accelerated failure time approach.

(DOCX)

S5 Table. Sensitivity analysis: risk factors analysis of pSBI in Senegal. pSBI, possible severe bacterial infection.

(DOCX)

Acknowledgments

We are grateful to all the mothers and their newborns for their participation. We thank all physicians, laboratory staff, field interviewers, and community workers for their involvement in this project. We thank all collaborators of the BIRDY program: Rasoanaivo Fanjalalaina

Vololonirina, Andrianonimiadana Lova Maminirina, Rabearitiana Lydos, Rasoloson Dimitri, Ratsima Hariniaina Elisoa, Volahasina Antsa Tanjona, Volahasina Antso Fenitra, Randriamamonjarison Aina Nirina, Andriatahina Todisoa, Marguerite Diatta, and Joseph Faye. We thank Yoann Madec for his advice in the statistical analysis. We also thank the Department of International Affairs and the [Centre de Recherche Translationnelle](#) - Coordination Clinique of Institut Pasteur for their support in the management of the study.

Author Contributions

Conceptualization: Bich-Tram Huynh, Elsa Kermorvant-Duchemin, Elisabeth Delarocque-Astagneau, Didier Guillemot.

Data curation: Bich-Tram Huynh.

Formal analysis: Bich-Tram Huynh, Elsa Kermorvant-Duchemin.

Funding acquisition: Bich-Tram Huynh, Elisabeth Delarocque-Astagneau, Didier Guillemot.

Investigation: Bich-Tram Huynh, Rattanak Chheang, Frederique Randrianirina, Abdoulaye Seck, Elisoa Hariniaina Ratsima, Zafitsara Zo Andrianirina, Jean-Baptiste Diouf, Armya Youssouf Abdou, Sophie Goyet, Véronique Ngo, Siyin Lach, Long Pring, Touch Sok, Fatoumata Diene Sarr, Laurence Borand, Perlinot Herindrainy, Agathe de Lauzanne, Muriel Vray.

Methodology: Bich-Tram Huynh, Elsa Kermorvant-Duchemin, Michael Padget, Benoit Garin, Elisabeth Delarocque-Astagneau, Didier Guillemot.

Supervision: Bich-Tram Huynh, Sophie Goyet, Véronique Ngo, Touch Sok, Michael Padget, Fatoumata Diene Sarr, Laurence Borand, Benoit Garin, Jean-Marc Collard, Perlinot Herindrainy, Agathe de Lauzanne, Muriel Vray, Elisabeth Delarocque-Astagneau.

Validation: Bich-Tram Huynh, Benoit Garin, Jean-Marc Collard, Agathe de Lauzanne.

Writing – original draft: Bich-Tram Huynh.

Writing – review & editing: Bich-Tram Huynh, Elsa Kermorvant-Duchemin, Rattanak Chheang, Frederique Randrianirina, Abdoulaye Seck, Elisoa Hariniaina Ratsima, Zafitsara Zo Andrianirina, Jean-Baptiste Diouf, Armya Youssouf Abdou, Sophie Goyet, Véronique Ngo, Siyin Lach, Long Pring, Touch Sok, Michael Padget, Fatoumata Diene Sarr, Laurence Borand, Benoit Garin, Jean-Marc Collard, Perlinot Herindrainy, Agathe de Lauzanne, Muriel Vray, Elisabeth Delarocque-Astagneau, Didier Guillemot.

References

1. Liu L, Oza S, Hogan D, Chu Y, Perin J, Zhu J, et al. Global, regional, and national causes of under-5 mortality in 2000–15: an updated systematic analysis with implications for the Sustainable Development Goals. *Lancet*. 2016; 388:3027–35. [https://doi.org/10.1016/S0140-6736\(16\)31593-8](https://doi.org/10.1016/S0140-6736(16)31593-8) PMID: 27839855
2. Seale AC, Blencowe H, Manu AA, Nair H, Bahl R, Qazi SA, et al. Estimates of possible severe bacterial infection in neonates in sub-Saharan Africa, south Asia, and Latin America for 2012: a systematic review and meta-analysis. *Lancet Infect Dis*. 2014; 14:731–41. [https://doi.org/10.1016/S1473-3099\(14\)70804-7](https://doi.org/10.1016/S1473-3099(14)70804-7) PMID: 24974250
3. Rudd KE, Johnson SC, Agesa KM, Shackelford KA, Tsoi D, Kievlan DR, et al. Global, regional, and national sepsis incidence and mortality, 1990–2017: analysis for the Global Burden of Disease Study. *Lancet*. 2020; 395:200–11. [https://doi.org/10.1016/S0140-6736\(19\)32989-7](https://doi.org/10.1016/S0140-6736(19)32989-7) PMID: 31954465
4. Laxminarayan R, Matsoso P, Pant S, Brower C, Røttingen J-A, Klugman K, et al. Access to effective antimicrobials: a worldwide challenge. *Lancet*. 2016; 387:168–75. [https://doi.org/10.1016/S0140-6736\(15\)00474-2](https://doi.org/10.1016/S0140-6736(15)00474-2) PMID: 26603918

5. Huynh B-T, Padget M, Garin B, Herindrainy P, Kermorvant-Duchemin E, Watier L, et al. Burden of bacterial resistance among neonatal infections in low income countries: how convincing is the epidemiological evidence? *BMC Infect Dis*. 2015; 15:127. <https://doi.org/10.1186/s12879-015-0843-x> PMID: 25888320
6. Herbert HK, Lee ACC, Chandran A, Rudan I, Baqui AH. Care seeking for neonatal illness in low- and middle-income countries: a systematic review. *PLoS Med*. 2012; 9:e1001183. <https://doi.org/10.1371/journal.pmed.1001183> PMID: 22412355
7. Saha SK, Schrag SJ, El Arifeen S, Mullany LC, Shahidul Islam M, Shang N, et al. Causes and incidence of community-acquired serious infections among young children in south Asia (ANISA): an observational cohort study. *Lancet*. 2018; 392:145–59. [https://doi.org/10.1016/S0140-6736\(18\)31127-9](https://doi.org/10.1016/S0140-6736(18)31127-9) PMID: 30025808
8. African Neonatal Sepsis Trial (AFRINEST) group, Tshetu A, Lokangaka A, Ngaima S, Engmann C, Esamai F, 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. [https://doi.org/10.1016/S0140-6736\(14\)62284-4](https://doi.org/10.1016/S0140-6736(14)62284-4) PMID: 25842221
9. Young Infants Clinical Signs Study Group. Clinical signs that predict severe illness in children under age 2 months: a multicentre study. *Lancet*. 2008; 371:135–42. [https://doi.org/10.1016/S0140-6736\(08\)60106-3](https://doi.org/10.1016/S0140-6736(08)60106-3) PMID: 18191685
10. Zaidi AKM, Huskins WC, Thaver D, Bhutta ZA, Abbas Z, Goldmann DA. Hospital-acquired neonatal infections in developing countries. *Lancet*. 2005; 365:1175–88. [https://doi.org/10.1016/S0140-6736\(05\)71881-X](https://doi.org/10.1016/S0140-6736(05)71881-X) PMID: 15794973
11. Zaidi AKM, Thaver D, Ali SA, Khan TA. Pathogens associated with sepsis in newborns and young infants in developing countries. *Pediatr Infect Dis J*. 2009; 28:S10–8. <https://doi.org/10.1097/INF.0b013e3181958769> PMID: 19106757
12. Berkley JA, Lowe BS, Mwangi I, Williams T, Bauni E, Mwarumba S, et al. Bacteremia among children admitted to a rural hospital in Kenya. *N Engl J Med*. 2005; 352:39–47. <https://doi.org/10.1056/NEJMoa040275> PMID: 15635111
13. Seale AC, Mwaniki M, Newton CRJC, Berkley JA. Maternal and early onset neonatal bacterial sepsis: burden and strategies for prevention in sub-Saharan Africa. *Lancet Infect Dis*. 2009; 9:428–38. [https://doi.org/10.1016/S1473-3099\(09\)70172-0](https://doi.org/10.1016/S1473-3099(09)70172-0) PMID: 19555902
14. Vergnano S, Sharland M, Kazembe P, Mwansambo C, Heath PT. Neonatal sepsis: an international perspective. *Arch Dis Child Fetal Neonatal Ed*. 2005; 90:F220–4. <https://doi.org/10.1136/adc.2002.022863> PMID: 15846011
15. World Health Organization. Neonatal and perinatal mortality: country, regional and global estimates. World Health Organization; 2006. Available from: <https://apps.who.int/iris/handle/10665/43444>
16. Lawn JE, Cousens S, Zupan J, Lancet Neonatal Survival Steering Team. 4 million neonatal deaths: when? Where? Why? *Lancet*. 2005; 365:891–900. [https://doi.org/10.1016/S0140-6736\(05\)71048-5](https://doi.org/10.1016/S0140-6736(05)71048-5) PMID: 15752534
17. Société Française de Microbiologie. Référentiel en microbiologie médicale. 6th ed. Société Française de Microbiologie; 2018.
18. Magiorakos A-P, Srinivasan A, Carey RB, Carmeli Y, Falagas ME, Giske CG, et al. Multidrug-resistant, extensively drug-resistant and pandrug-resistant bacteria: an international expert proposal for interim standard definitions for acquired resistance. *Clin Microbiol Infect*. 2012; 18:268–81. <https://doi.org/10.1111/j.1469-0691.2011.03570.x> PMID: 21793988
19. Pedersen AB, Mikkelsen EM, Cronin-Fenton D, Kristensen NR, Pham TM, Pedersen L, et al. Missing data and multiple imputation in clinical epidemiological research. *Clin Epidemiol*. 2017; 9:157–66. <https://doi.org/10.2147/CLEP.S129785> PMID: 28352203
20. Bartlett JW, Harel O, Carpenter JR. Asymptotically Unbiased Estimation of Exposure Odds Ratios in Complete Records Logistic Regression. *Am J Epidemiol*. 2015; 182:730–6. <https://doi.org/10.1093/aje/kvv114> PMID: 26429998
21. Islam MS, Baqui AH, Zaidi AK, Bhutta ZA, Panigrahi P, Bose A, et al. Infection Surveillance Protocol for a Multicountry Population-based Study in South Asia to Determine the Incidence, Etiology and Risk Factors for Infections Among Young Infants of 0 to 59 Days Old. *Pediatr Infect Dis J*. 2016; 35:S9–15. <https://doi.org/10.1097/INF.0000000000001100> PMID: 27070072
22. Wynn JL. Defining neonatal sepsis. *Curr Opin Pediatr*. 2016; 28:135–40. <https://doi.org/10.1097/MOP.0000000000000315> PMID: 26766602
23. Weston EJ, Pondo T, Lewis MM, Martell-Cleary P, Morin C, Jewell B, et al. The burden of invasive early-onset neonatal sepsis in the United States, 2005–2008. *Pediatr Infect Dis J*. 2011; 30:937–41. <https://doi.org/10.1097/INF.0b013e318223bad2> PMID: 21654548

24. Huynh B-T, Kermorvant-Duchemin E, Herindrainy P, Padget M, Rakotoarimanana FMJ, Feno H, et al. Bacterial Infections in Neonates, Madagascar, 2012–2014. *Emerg Infect Dis*. 2018; 24:710–7. <https://doi.org/10.3201/eid2404.161977> PMID: 29553312
25. Aiken AM, Mturi N, Njuguna P, Mohammed S, Berkley JA, Mwangi I, et al. Risk and causes of paediatric hospital-acquired bacteraemia in Kilifi District Hospital, Kenya: a prospective cohort study. *Lancet*. 2011; 378:2021–7. [https://doi.org/10.1016/S0140-6736\(11\)61622-X](https://doi.org/10.1016/S0140-6736(11)61622-X) PMID: 22133536
26. Downie L, Armiento R, Subhi R, Kelly J, Clifford V, Duke T. Community-acquired neonatal and infant sepsis in developing countries: efficacy of WHO's currently recommended antibiotics—systematic review and meta-analysis. *Arch Dis Child*. 2013; 98:146–54. <https://doi.org/10.1136/archdischild-2012-302033> PMID: 23142784
27. Taylor AW, Blau DM, Bassat Q, Onyango D, Kotloff KL, Arifeen SE, et al. Initial findings from a novel population-based child mortality surveillance approach: a descriptive study. *Lancet Glob Health*. 2020; 8:e909–19. [https://doi.org/10.1016/S2214-109X\(20\)30205-9](https://doi.org/10.1016/S2214-109X(20)30205-9) PMID: 32562647
28. Soman M, Green B, Daling J. Risk factors for early neonatal sepsis. *Am J Epidemiol*. 1985; 121:712–9. <https://doi.org/10.1093/aje/121.5.712> PMID: 4014163
29. Belachew A, Tewabe T. Neonatal sepsis and its association with birth weight and gestational age among admitted neonates in Ethiopia: systematic review and meta-analysis. *BMC Pediatr*. 2020; 20:55. <https://doi.org/10.1186/s12887-020-1949-x> PMID: 32020850
30. Simonsen KA, Anderson-Berry AL, Delair SF, Davies HD. Early-onset neonatal sepsis. *Clin Microbiol Rev*. 2014; 27:21–47. <https://doi.org/10.1128/CMR.00031-13> PMID: 24396135
31. Chan GJ, Lee AC, Baqui AH, Tan J, Black RE. Risk of early-onset neonatal infection with maternal infection or colonization: a global systematic review and meta-analysis. *PLoS Med*. 2013; 10:e1001502. <https://doi.org/10.1371/journal.pmed.1001502> PMID: 23976885