

Prevalence of group B streptococcus in pregnant women attending a tertiary hospital in Ghana in 2001

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Abstract

Title: Prevalence of group B streptococcus in pregnant women attending a tertiary hospital in Ghana in 2001

Background: Group B streptococci cause major perinatal bacterial infections including chorioamnionitis and endometritis in women and severe systemic infections in the newborn. Infections are a major cause of newborn deaths in Ghana but the organisms causing these infections are usually not identified. Group B streptococci colonization during pregnancy, a prerequisite to potentially preventable early onset disease in the newborn is believed to be rare in pregnant women in Ghana. We undertook this pilot study to assess the veracity of this belief.

Methods: General health education information on group B streptococcus was given to all women attending the Wednesday antenatal clinic at Korle Bu Teaching Hospital, Accra from April – June 2001. Anorectal and lower genital tract swab specimens were obtained from consecutive volunteers of 100 women in the third trimester of pregnancy. Specimens were inoculated onto selective Todd-Hewitt broth, incubated for 18 hours and sub-cultured onto sheep blood agar plate for group B streptococcus identification.

Findings: Of the 2420 antenatal consultations at the Wednesday clinic during the study period, 32.2% (781/2420) of the attendants were in the third trimester of pregnancy. There were 21 isolates (12 anorectal, 5 genital and 2 at both sites) in 19 women; 19% of the pregnant women were carriers of group B streptococcus.

Conclusion: The prevalence of group B streptococcus colonization in pregnant Ghanaian women is similar to that in other developed and developing countries. The rarity of isolates of group B streptococcus in Ghana may be due to inadequate microbiological methods.

Keywords: group B streptococcus, pregnancy, prevalence

Introduction

For over four decades, *Streptococcus agalactiae* also known as group B streptococcus (GBS) which commonly inhabits the genital, lower urinary and gastrointestinal tracts of pregnant women has been known as a leading cause of perinatal infections [1-2]. It causes chorioamnionitis and endometritis in women and pneumonia, septicemia and meningitis in the newborn. It has also been associated with increased risk of preterm delivery. Maternal carriage of GBS is a prerequisite for early onset disease in the newborn. Infections in the fetus are

transmitted vertically via an intact or ruptured amniotic membrane or during passage through a colonized birth canal [3]. Risk of neonatal infection is highest if mothers are colonized with GBS in the third trimester.

Newborn deaths are a major contributor to under 5 years child mortality in Ghana and a limiting factor to the achievement of millennium development goal 4. Infections are a major cause of stillbirths and newborn deaths in Ghana [4]. It is estimated that at least 32% of newborn deaths in Ghana are caused by infections; a significant number of these infections are acquired

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Authors' contribution:

All 3 authors worked together in the design of the study, NRK Damale recruited the subjects and collected the specimens, MJ Newman organized the laboratory investigations. CC Enweronu-Laryea wrote the draft manuscript and all 3 authors reviewed the manuscript before submission for publication

Conflict of interest: The authors have no conflict of interest to declare.

Funding: The authors received no funding for this work.

from the mother. Early onset neonatal infections are potentially preventable and effective intrapartum antimicrobial prophylaxis has been shown to significantly reduce the morbidity and case-fatality rate for infants at risk [5-6]. It is therefore important to know the antimicrobial susceptibility of organisms that colonize the genital tract of pregnant women over time.

GBS colonization of the genitourinary tract of pregnant Ghanaian women is believed to be rare because of traditional female hygienic practices and the rarity of the organism in isolates from local microbiological laboratories. A literature search revealed very little on GBS in Ghana. We conducted this pilot study using recommended guidelines for GBS microbial isolation to provide accurate baseline data on GBS carriage and antimicrobial sensitivity in pregnant Ghanaian women in 2001. We report our findings to stimulate further research in intrapartum antimicrobial prophylaxis and microbiological causes of newborn deaths in the region.

Materials and Methods

This cross-sectional study was conducted at the antenatal clinic of Korle Bu Teaching Hospital (KBTH) in Accra from April to June 2001. KBTH is a tertiary referral hospital for southern Ghana with approximately 900 deliveries per month. Antenatal clinics are held every weekday.

Participants were recruited from the Wednesday antenatal clinic of one of the investigators. About 180 women at all stages of pregnancy attend this clinic each week. General information about the relevance of GBS screening in perinatal care was offered to all women attending a routine morning antenatal care clinic. Consecutive volunteers who met the inclusion criteria (gestational age ≥ 28 weeks + no clinical complaint by the woman) were screened during their turn for consultation with the investigator until a total number of 100 women were recruited. Women with obstetric (e.g. abdominal pain, vaginal discharge) or other clinical (e.g. fever, urinary symptoms) complaints were excluded because the study was focused on GBS carriage in the normal population of pregnant women. A standard questionnaire was used to obtain basic demographic (age and marital status) and obstetric (gestational age and parity) data.

All specimens were collected by the investigator with a sterile cotton applicator. A lower vaginal specimen was taken about 2cm from the introitus and another sterile cotton swab was used to sweep the anorectum. Specimens were inoculated into Todd Hewitt broth (contains 15mg/ml gentamicin and 25 μ l/ml nalidixic acid) soon after collection at the clinic. After an overnight incubation at 37°C a standard loop (200 μ l) was used to transfer a loopful of broth and plated on 5% sheep blood agar. These plates were incubated in 5 – 10% CO₂ using candle extinction jar at 37°C for 24 hours, and then β -hemolytic colonies which were Gram-positive cocci and catalase-negative were

serotyped with Streptex (Plasmatec Laboratory, UK). All GBS isolates were tested for antimicrobial susceptibility to crystalline penicillin.

Categorical variables were compared using chi square test. Differences were considered significant if $p < 0.05$.

Results

There were 2420 antenatal consultations at the Wednesday clinic by women at all stages of pregnancy during the study period. About 32% (781/2420) of the clinic attendants during the period of study were in the third trimester of pregnancy. Even though all the volunteers had no complaint at the time of recruitment, 21 of the 100 women were found to have abnormal vaginal discharge by the investigator during gynecological examination.

Table 1 shows the characteristics of the 100 participants. There was no difference in the marital status, age, parity and gestational age of women colonized with GBS and those who were not. There were 21 isolates from nineteen participants, 12 harbored the organism only in the anorectum, 5 had it only in the genital tract and 2 had GBS at both sites. Nineteen percent (19%) of the participants were colonized with GBS. All isolates were susceptible to crystalline penicillin.

TABLE 1 Characteristics of subjects and group B streptococcus carriage

Characteristics	Number of subjects	Number group B streptococcus positive (%)	p value*
Maternal age (years)			
20 - 34	84	16 (19.1)	1.00
≥ 35	16	3 (18.8)	
Marital status			
Married	85	15 (18.8)	0.476
Unmarried	15	4 (26.6)	
Parity			
Primigravida	45	8 (17.8)	0.804
≥ 1 child	55	11 (20)	
Gestational age (weeks)			
28 - 34	37	8 (21.6)	0.608
≥ 35	63	11 (17.5)	
Vaginal discharge			
Present	21	4 (19.1)	1.00
Absent	79	15 (19.0)	

* Chi square test

Discussion

We have shown that 19% of Ghanaian women in the third trimester of pregnancy in 2001 were colonized with penicillin-sensitive GBS. The prevalence of GBS carriage in our study is similar to that described by other studies [7-9] from other parts of the world. Colonization in pregnancy is intermittent but maternal colonization in the third trimester especially after 35 weeks gestation is predictive of neonatal colonization and infection, 17.5% of participants at ≥ 35 weeks gestation were colonized in this study. Current guidelines for cost-effective screening and chemoprophylaxis are from 35 weeks gestation of pregnancy [10]. We screened women from 28 weeks gestation because we could not find any published work on GBS colonization or its association with premature births in Ghana.

This pilot study has the inherent limitations of studies with volunteer participants. Though the findings of our work are clinically and epidemiologically relevant no conclusions can be made based on our data because of the small sample size. The inclusion criteria for the study were based on the women's perception of abnormal symptoms and women with a complaint of abnormal vaginal discharge were excluded. Even though none of the 100 women had any clinical complaint at recruitment, 21% of them were found to have abnormal-looking vaginal discharge during the sampling procedure. This is not an unusual observation in routine clinical practice and sometimes what the obstetrician observes as suspected abnormality yield normal flora on microbiological testing. There was no difference in GBS carriage in these women compared to those without vaginal discharge in this study.

No conclusions can be made on the correlation of colonization with maternal age and marital status, gestational age, parity or vaginal discharge because of the small number of subjects in this cohort. However other studies [11-12] have not found a strong association between these factors and GBS colonization. Vertical transmission of GBS to the newborn occurs in 40 – 73% of culture positive women but only 2% of these infants develop early-onset disease [13]. However, the risk of disease can increase up to 35 fold in the event of risk factors including prolonged rupture of membranes, intrapartum fever and low birth weight. These risk factors are not uncommon in Ghanaian pregnant women.

The microbiological findings of our study are similar to those of Badri *et al* [14] and Quinlan *et al* [15] who also cultured more GBS from the anorectum than the genital tract. This lower yield of the genital tract can be explained by our sampling of only the lower vagina. In the study by Anthony *et al* [16] the cotton swab for sampling the lower vagina was used to swab the periurethral area and medial aspects of labia majora and they obtained almost identical isolates from the anorectal and genital cultures. Cultures from both sites are recommended because the anorectum may provide a reservoir for intermittent genitourinary tract re-colonization. All the isolates of GBS in our study

were susceptible to penicillin G as observed in other studies [17-19]. However, the susceptibility pattern of GBS in Ghanaian women presently may be different from the findings of this study in 2001 because of increased use and misuse of antibiotics inherent in many developing countries.

There is no established protocol for screening or prevention of GBS disease in Ghana. The common practice for microbiological testing in Ghanaian pregnant women involves the collection of a high vaginal swab specimen that is transported to a laboratory for plating on solid media. This method fails to identify GBS in many women harboring the organism. We have shown that maternal carriage of GBS is not rare in women attending antenatal clinic at KBTH. The supposed rarity of GBS in Ghana is most likely due to inadequate microbiological methods. A large multicentre study using the recommended microbiological methods and wider spectrum of antimicrobial susceptibility of isolates is recommended.

Acknowledgement

The authors wish to express their gratitude to Francis Codjoe of the Department of Microbiology of KBTH for his technical assistance and Elizabeth Amui of the Department of Obstetrics for her help in the recruitment of the women.

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