Isolation of Group B Streptococci from Antenatal Women

Tupili Ramya*, K.H. Vasudeva Naidu and Usham Gangaram

Department of Microbiology, Sri Vekateshwara Medical College, Tirupathi, India

*Corresponding author

A B S T R A C T

Prevalence of Group-B streptococcal (GBS) infection varies from place to place and this organism is responsible for serious infections in newborns such as septicemia and meningitis. The present study was carried out to find out the prevalence & antibiotic sensitivity of GBS in primigravida & to identify the risk factors. 300 pregnant women were studied attending the antenatal clinics in Government Maternity Hospital, Tirupati. Two vaginal swabs were taken from each pregnant women. The two swabs were immediately transported to the laboratory for processing. Direct Gram stain was done from one swab and the other swab was inoculated onto sheep blood agar plate and incubated at 37ºC for 24-48 hours. Identification was done based on Gram staining, colony morphology, catalase reaction, CAMP test and Hippurate hydrolysis test & Bacitracin resistance. Of the 300 pregnant women screened, 7% were colonized by GBS. GBS colonization rate was higher among pregnant women in third trimester who were <25 years of age. All the isolates were sensitive to Penicillin, Ampicillin, ceftriaxone, levofloxacin and vancomycin. GBS colonization rate among pregnant women in third trimester of pregnancy is low in this area. Revised guidelines from the Centers of Disease Control and Prevention (CDC), 2010 for the prevention of perinatal GBS disease recommends that all pregnant women be screened for GBS carriage between 35 and 37 weeks of gestation and intrapartum antibiotic prophylaxis be given to colonized women at the time of labour onset or rupture of membranes.

Keywords
Group B streptococcus, Colonization, Drug resistance, Drug sensitivity.

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Introduction

Group-B streptococci or Streptococcus agalactiae, is a well recognized pathogen in veterinary medicine because of its role as causative agent of bovine mastitis. There has been increasing interest in Group-B streptococci (GBS) due to their association with and recognition as pathogens of neonatal infections like sepsis and meningitis (Hood, et al., 1961; Eickhoff, et al., 1964). They are present in 15-20% of pregnant women, in lower genital tract (Hoogkamp-Korstanje, et al., 1982). Vaginal colonization by GBS during pregnancy is associated with life threatening neonatal infections acquired during passage through the birth canal, and it has been the most frequent cause of purulent meningitis in infants aged less than 2 months (Baker, et al., 1973; Barton, et al., 1973).
Group-B streptococci has been associated with low-birth weight, premature delivery, preterm premature rupture of membranes, still birth and neonatal death suggesting that it has a marked predilection for pregnant women, fetus and neonates (Wheelver, 1966). The mother’s birth canal is the principle reservoir of this infectious agent for infants who develop the more severe syndrome with early onset (Franciosi, et al., 1973).

In India very few studies have been carried out, mainly to study the prevalence of GBS infections. The GBS colonization rate in the vaginal flora of pregnant women varied from 0.47 to 23.3% as reported by various workers (Arora, et al., 1994; Kishore, et al., 1986; Lakshmi, et al., 1998; Prakash, et al., 1976; Santosh Damyanti, et al., 1984; Uma Chaudhary, et al., 1981).

The factors controlling maternal GBS colonization and neonatal disease due to GBS are – age, number of previous pregnancies, phase of menstrual cycle, sexual activity, use of oral contraceptives (Baker, et al., 1977; Schauf, et al., 1976).

This study was undertaken in order to understand the magnitude of this problem in pregnant women in our area. The main aim of this study includes, to study the prevalence of group B Streptococcal infection from antenatal primi gravida attending antenatal clinic of government maternity Hospital attached to S.V. Medical college. And also to study drug sensitivity of group B streptococci to penicillin, ampicillin, ceftriaxone, Clindamycin, vancomycin.

**Material and Methods**

**Study Area and Period**

The present hospital based cross sectional study was carried out in the antenatal clinic & Department of Microbiology, S.V. Medical College, Tirupati, from June 2012 to May 2013.

**Source Population**

300 pregnant women were studied attending the antenatal clinics in Government Maternity Hospital, Tirupati. These women were in the age group of 19 years to 37 years. The inclusion criteria for study were the presence of signs & symptoms of pregnancy & willingness to participate in the study. Detailed history of each case was taken during their clinical examination. Group B streptococcal positive women were divided in four groups: Group I:≤ 20 years; Group II: 21 to 25 years ; Group III: 26to30 years and Group IV: above30years.

**Data Collection, Processing and Analysis**

Two low vaginal swabs were taken aseptically prior to first pelvic examination. The swabs were immediately transported to the Microbiology laboratory. One swab was used for direct gram staining and the other swab was inoculated on to sheep blood agar containing 5% sheep blood; it was incubated at 37°C for 24-48 hours. Identification was done based on gram stain, colony morphology, catalase reaction, CAMP test, hippurate hydrolysis test. 

**The Presumptive Diagnosis of GBS was based on the following Criteria**

Direct gram staining showing gram positive cocci arranged in pairs and short chains. The colony appearance of GBS on sheep blood agar at 24 hours is usually grey, smooth, shiny, convex, moist, regular, soft and mucoid in appearance and about 1 mm in diameter, often surrounded by a small hazy zone of beta hemolysis. The confirmation of GBS was made by
subculturing colony from the blood agar on to chocolate agar to check the catalase activity. A clean (grease free) glass slide was taken, 1 or 2 drops of H2O2 (3%) was put on the slide. Using a clean glass rod, a colony was picked and dipped into H2O2. Staphylococci produce catalase while streptococci does not. Effervescence is seen in positive catalase test whereas no effervescence is seen in negative catalase test. Other confirmatory tests carried out were CAMP and Hippurate hydrolysis test.

Antimicrobial sensitivity of the GBS was done by the Kirby-Bauer disc diffusion method. Fresh sub-cultures of GBS were used after overnight growth (16 hours) on blood agar plate.

The inoculum was prepared by suspending several of the colonies in sterile phosphate buffered saline (pH 7.2) to achieve a turbidity of 0.5 McFarland standard. This resulted in a suspension containing approximately $1-2 \times 10^8$ CFU/ml. A sterile cotton swab was dipped into the bacterial suspension, elevated above the liquid and rotated several times against the inside wall of the tube to remove excess of the inoculum.

This swab was streaked evenly in three different directions onto the blood agar containing 5% sheep blood. Five antibiotic discs were employed namely Penicillin Penicillin (10 μg/disc), Ampicillin (10 μg/disc), Clindamycin (2 μg/disc), Ceftriaxone (30 μg/disc), Vancomycin (30 μg/disc). The data was analyzed and interpreted.

**Results and Discussion**

In the present study, out of 300 women, 21 (7 %) showed GBS colonization (figure 1). According to Table 1, most of the isolates were from the age group ≤ 20 years (8.73%), followed by age group 21 to 25 (7.18%). By applying chi square test relationship between GBS colonization and age group was not statistically significant.

As per the Table 2, most of the GBS isolated from the third trimester of pregnancy (10.19%). Out of 10 samples from the third trimester of pregnancy, one sample turned to be positive. This indicates that third trimester of pregnancy is a risk factor for the colonization of GBS, compared to the other trimester.

All the group-B streptococci showed 100% sensitivity to Ampicillin, Penicillin, Ceftriaxone, Vancomycin and 90.47% sensitivity to Clindamycin.

A total of 300 primigravida women who were attending to antenatal clinic included in this study. These included females in the age group of 19 years to 37 years. Figure 1: Shows the prevalence of GBS is 7.0% in the present study.

Motlova, et al., (2004) showed that 29.3% of pregnant women were colonized by GBS. In a study done by Kavitha P Konikkara, et al., (2008) showed the GBS carriage rate 12%. In comparison to the above studies, our study shows the incidence rate of 7.0%, which is lower.

But when compared to the studies done by AA Kulkarni, et al., (2001) and Vijayan Sharmila, et al., (2011), which shows the colonization rate of 2.52% and 2.3% respectively. Our incidence rate was found to be higher.

The incidence rate in our study (7%) correlates with the study done by Annie Rajaratnam, et al., (2013), which also showed the colonization rate of 8.3%. And also correlates with study done by Vinay
Hazare, et al., (2012), which showed the colonization rate of 7.5%.

The reasons for varying results may be attributed to the fact that GBS maternal colonization varies from place to place. Other factors that may have contributed to this variation include socioeconomic factors, variation in clinical practices of samples collected and the technique used for sampling. Ethnic and genetic factors might play a role in variation of the rates of infection with GBS.

According to Table 1, in the present study most of the isolated strains were from cases belonging to age group less than and equals to 20 years (8.73%) followed by the age group of 21-25 years (7.18%), thus in the present study, most of the GBS colonized pregnant women were from teenage and younger age group.

These findings are in correlation with that of Vinay Hazare, et al., (2012), Tsering Chomu Dechen, et al., (2010), the reasons for the predisposition of women younger than 21 years of age to vaginal colonization with GBS is less apparent. This relationship could be the result of age related development of local or humoral immunological responses that interfere with mucosal attachment and/ or persistence of GBS.

Table-2: Shows most of the GBS isolated from the third trimester of pregnancy (10.19%). This indicates third trimester of pregnancy is a risk factor for the colonization of GBS, compared to the other trimester. According to Fatemi, et al., (2008), 7 (8.8%) of 80 women with any kind of antibiotic therapy during third trimester and 61 (24.4%) of 250 in those who were not receiving antibiotic, the results of culture were positive. Baker, et al., (1977) found that the colonization rate almost doubles between the second trimester and delivery.

According to Fashina, et al., (2008), Out of 100 women tested, 90 were positive in third trimester and 10 were in the second trimester and cultures were negative in the first trimester. In contrast, Hansen, et al., (2004) did not find any significant variation in the prevalence of GBS during pregnancy.

Revised guidelines from the Centers of Disease Control and Prevention (CDC), 2010 for the prevention of perinatal GBS disease recommends that all pregnant women be screened for GBS carriage between 35 and 37 weeks of gestation and intrapartum antibiotic prophylaxis be given to colonized women at the time of labour onset or rupture of membranes.

Figure-2: shows antibiotic sensitivity of 21 GBS isolates from mothers. All the strains were 100% sensitive to Ampicillin, Penicillin, Ceftriaxone, Vancomycin and 90.47% sensitive to Clindamycin.

Arora, et al., (1994) reported that all the 60 GBS were susceptible to Penicillin, Ampicillin, & Erythromycin. Vinay Hazare, et al., (2012) showed that all the 15 strains were 100% sensitive to ampicillin, erythromycin and penicillin followed by chloramphenicol (66.6%). A marked resistance was observed with Gentamicin (100%) and Kanamycin (80%).
Table 1 Isolation of GBS in Different Age groups

| Age group | No. of primigravida | GBS positive | Percent |
|-----------|---------------------|--------------|---------|
| ≤ 20 years | 103                 | 9            | 8.73    |
| 21 – 25   | 167                 | 12           | 7.18    |
| 26 – 30 years | 23               | 0            | 0.0     |
| >30       | 7                   | 0            | 0.0     |
| Total     | 300                 | 21           | 7.0     |

χ²=0.927  p=0.818  Insignificant

Table 2 Isolation of GBS in different trimesters

| trimester | No. of primigravida | GBS Positives | Percent |
|-----------|---------------------|---------------|---------|
| First     | 34                  | 0             | 0.0     |
| Second    | 60                  | 0             | 0.0     |
| Third     | 206                 | 21            | 10.19   |
| Total     | 300                 | 21            | 7.00    |

χ²=6.928  p=0.031  significant

Fig. 1 Prevalence of GBS in Primigravida

Fig. 2 Antibiotic Sensitivity testing of 21 GBS isolates

Fashina, et al., (2008) reported, the sensitivity pattern of Streptococcus agalactiae showed that 100% were sensitive to Penicillin G and Erythromycin. Eighty percent (80%) were sensitive to Ampicillin, Vancomycin and Augmentin, while Tetracycline was markedly resisted with a sensitivity of 30%.
According to Jannati, et al., (2012) all isolates were susceptible to ampicillin, vancomycin and penicillin. 96.7% and 93% of isolates were susceptible to erythromycin and clindamycin respectively and 83.9%, 14.2%, 12.5% isolates were resistant to Cotrimoxazole ciprofloxacin and ceftriaxone respectively.

Recommendation

Most of the clinicians are not aware about the incidence of GBS in their area. So an attempt was made to find out the incidence of GBS in pregnant women which would help the clinicians to diagnose the GBS infections.

A low colonization rate of 7% was found among the pregnant women in third trimester of pregnancy in this part of study area. Among them higher colonization rate was found in pregnant women in third trimester of pregnancy who were ≤ 20 years of age and primi gravidae. According to our study, risk factors for GBS colonization includes age less than 20 years and third trimester.

As per Revised guidelines from the CDC, 2010 for the prevention of perinatal GBS disease recommends that all pregnant women be screened for GBS carriage between 35 and 37 weeks of gestation and intrapartum antibiotic prophylaxis be given to colonized women at the time of labour onset or rupture of membranes”.

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