Clinical Profile and Outcome of Group B Streptococcal Colonization in Mothers and Neonates in Ras Al Khaimah, United Arab Emirates: A Prospective Observational Study

Shatha Taher Salman AlZuheiri¹, Rajani Dube², Godfred Menezes³, Samar Qasem¹

¹Department of Obstetrics and Gynecology, Abdulla Bin Omran Hospital for Obstetrics and Gynecology, Departments of ²Obstetrics and Gynecology and ³Microbiology, RAK Medical and Health Sciences University, Ras Al-Khaimah, UAE

Abstract Background: Maternal Group B *Streptococcus* (GBS)/*Streptococcus agalactiae* colonization rates vary worldwide; however, no such recent data are available from the United Arab Emirates (UAE).

Objective: The objective of this study was to determine the prevalence of GBS colonization among pregnant women attending an antenatal clinic of a hospital in Ras Al Khaimah, UAE, along with the antibiotic sensitivity pattern, the clinical profile and pregnancy (maternal and fetal) outcome.

Methods: This prospective observational study routinely offered rectovaginal swab for GBS to all women attending the antenatal clinic at 35–37 weeks of pregnancy between January and December 2019. MASTASTREP kit and Vitek-2 identification system was used for culture and identification. Women with positive cultures were followed up for any maternal and neonatal complications and the use of intrapartum antibiotic prophylaxis (IAP).

Results: A total of 2295 women were included, of which 158 (6.9%) had positive cultures for GBS colonization. The carriage rate was higher in women without any risk factors for early-onset GBS disease (EOGBS) (P < 0.01). The GBS isolates were about 97% susceptible to linezolid and vancomycin, 90% to benzyl penicillin and 95% to ampicillin. Resistance to trimethoprim/sulfamethoxazole, clindamycin, erythromycin, and levofloxacin were about 77%, 57%, 57%, and 10%, respectively. Urinary tract infection in GBS colonized women were more common in those aged \leq 30 years (P = 0.009). Fetal outcome was favorable in women receiving IAP for GBS colonization. No neonate had culture proven EOGBS.

Conclusion: The prevalence of GBS colonization in pregnant women as well as the overall maternal and neonatal complications is low in Ras Al Khaimah, UAE. IAP is effective in preventing early-onset sepsis in newborn, and thus should be initiated in those with GBS colonization. The cultured GBS showed sensitivity to most antibiotics.

Keywords: Colonization, Group B Streptococcus, neonatal outcome, pregnancy outcome, sensitivity, UAE

Address for correspondence: Dr. Rajani Dube, RAK Medical and Health Science University, PO Box 11172, Ras Al-Khaimah, UAE. E-mail: rajani.dube@rakmhsu.ac.ae Submitted: 31-Mar-2021 Revised: 29-May-2021 Accepted: 24-Jun-2021 Published: 04-Sep-2021

Access this article online		
Quick Response Code:	Website:	
	www.sjmms.net	
	DOI: 10.4103/sjmms.sjmms_213_21	

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

For reprints contact: WKHLRPMedknow_reprints@wolterskluwer.com

How to cite this article: AlZuheiri ST, Dube R, Menezes G, Qasem S. Clinical profile and outcome of Group B streptococcal colonization in mothers and neonates in Ras Al Khaimah, United Arab Emirates: A prospective observational study. Saudi J Med Med Sci 2021;9:235-40.

INTRODUCTION

Group B *Streptococcus* (GBS)/*Streptococcus agalactiae* is a normal body commensal, colonizing the gut and vagina. Maternal GBS colonization rates vary worldwide, with a mean prevalence of about 18%.^[1,2] Currently, there is a lack of consensus regarding the association between risk factors, such as age, parity, education, socioeconomic status, and maternal factors (premature rupture of membranes [PROM], low birth weight and puerperal pyrexia), and GBS colonization.^[1-5]

In neonates, GBS infection is generally acquired by vertical transmission in the peripartum period and is a significant cause of sepsis and death in many countries.^[2,6-9] About 75% of the neonatal GBS infections present as pneumonia, septicemia or meningitis in the first 6 days of life.^[10] Meningitis causes long-term disabilities such as blindness, deafness, speech problems or learning impairments in up to 50% of the cases, and these are severe in about 25% of the neonates. In mothers, GBS is also a frequent cause of asymptomatic bacteriuria, urinary tract infection (UTI), chorio-amnionitis, postpartum endometritis, pneumonia, puerperal sepsis and bacteremia without a focus.^[11,12]

From the Middle East, studies have found the prevalence of maternal GBS colonization to range from 3.3% to 31.6%;^[13-15] however, there is no recent study from the United Arab Emirates (UAE) reporting the clinico-bacterial profile, prevalence, and antibiotic sensitivity pattern. Accordingly, this study was conducted to determine the prevalence of GBS colonization in pregnant women from UAE after 35 weeks of gestation as well as determine their clinical profile, the maternal and fetal outcome of those colonized with GBS, and the antibiotic sensitivity pattern of GBS.

METHODS

Study design and participants

This was a prospective observational study conducted at the Department of Obstetrics & Gynecology, Abdullah Bin Omran Hospital, Ras Al Khaimah, UAE, after receiving the ethical approval from the Ministry of Health and the local ethics committee.

All pregnant women attending the antenatal unit of the hospital between January 1 and December 31, 2019, were routinely offered rectovaginal swab for GBS at 35–37 weeks of pregnancy. In addition, in those who presented with prolonged PROM or preterm labor, high vaginal swab was done as part of the initial assessment. Urine culture was ordered for women presenting with symptoms of UTI. Written consent was obtained prior to the test, and maternal characteristics and antenatal risk factors were noted.

Women with GBS-positive cultures were followed up for antenatal complications, mode of delivery, gestational age at delivery and use of IAP. All the included GBS-positive women delivered at our hospital and their neonates were kept under observation for 48 hours. The neonates were examined by a pediatrician at birth and then at regular intervals during the hospital stay. Thereafter, the patients were followed-up telephonically weekly during the entire neonatal period for any symptoms. For any further neonatal complications, the interventions and outcomes were noted.

Specimen collection

Rectovaginal swabs were collected by brushing the lower vagina and rectum using a sterile cotton swab by obstetricians following universal precautions.

Identification of Group B Streptococcus

The isolates were identified as GBS based on being catalase-negative, Gram-positive cocci, CAMP positive,^[16] bacitracin resistant and their reaction with commercial latex-group-specific streptococcal typing system (MASTASTREP kit, Mast House, Merseyside, UK) and Vitek-2 identification system (BioMérieux, Inc., Durham, NC).

Antimicrobial susceptibility test

The antibacterial susceptibility of all GBS isolates was tested by automated susceptibility testing using Vitek-2 AST-03 card (BioMérieux, Inc., Durham, NC) according to Clinical Laboratory Standard Institute (CLSI, 2017).^[17] The antibacterial agents tested included benzyl penicillin, ampicillin, ceftriaxone, cefotaxime, clindamycin, erythromycin, levofloxacin, trimethoprim/ sulfamethoxazole, vancomycin, and linezolid.

Data analysis

Data were analyzed using SPSS version 25 (SPSS, Inc., Chicago, IL, USA). The numerical variables were expressed in percentages. Groups were compared by Chi-square analysis and a P value <0.05 was considered statistically significant.

RESULTS

A total of 3007 antenatal patients attended the hospital during the specified period, of which 2295 women agreed to undergo the test and gave consent. Of these, GBS colonization was found in 158 women (6.9%). All further

analysis only includes the GBS-positive women, unless stated otherwise.

The mean age was 30.67 ± 4.58 years (range: 19–43 years), with 79 women each aged ≤ 30 and >30 years. Further, 27 (17%) women were primigravida and most had parity ≥ 1 (n = 131; 82.9%). The mean body mass index (BMI) was 32.5 ± 12.1 kg/mt²; the majority were obese with a BMI of ≥ 30 (n = 105; 66.4%), while 13.2% (n = 21) had Class 3 obesity [Table 1].

Prevalence of GBS colonization among women with low vaginal samples (LVS) and/or rectal swab for routine screening was 22.7% (123 of 541). The GBS colonization in women with PROM or preterm labor was 1.1% (20 of 1749). One of the two women with a history of GBS colonization in a previous pregnancy had colonization. Urine culture was carried out in 29 women, of which 13 (44.8%) were positive.

UTI was significantly higher in women aged ≤ 30 years that those aged >30 years (P = 0.009), but there was

Table 1: Maternal and neonatal profile

Parameter	Value
Age (years)	
≤30	79
>30	79
BMI (kg/m ²)	
<25	12
25-29.9	41
30-34.9	52
35-39.9	32
≥40	21
Parity index	
Primigravida	27
Parity 1 or above	131
Pregnancy complications	
Diabetes*	41
Anemia	8
Hypertension**	7
Others***	17
IUGR	11
Type of delivery	
Vaginal delivery	127
Elective LSCS	11
Emergency LSCS	20
Birth weight (g)	
<2500	11
2500-3500	115
>3500	32
Postpartum complications	
Endometritis	3
Symptomatic urinary tract infection	1

*Gestational diabetes (n=37) and pregestational diabetes (n=4),

**Hypertension is defined as gestational hypertension + preeclampsia;
***Others: Obstetric cholestasis=1, Twin pregnancy=2, Hypo/
Hyperthyroidism=5, Toxoplasmosis under treatment=1, Rh negative
pregnancy=3, Genital wart=1, Treated micro-invasive cervical
carcinoma=1, polyhydramnios=1, Gestational thrombocytopenia=2.
BMI – Body mass index, IUGR – Intrauterine growth restriction, LSCS
– Lower segment Caesarean section

no statistical difference between primigravida and multipara (P = 0.34) or lean and obese (P = 0.08) women. The most common pregnancy-associated complication was diabetes 25.9% (n = 41), while 10.7% (n = 17) of women had other complications including thyroid dysfunction, gestational thrombocytopenia, and blood group Rh-negative pregnancy. The mean birth weight of neonates was 3186 ± 470 g. Among neonates requiring admission, risk factors in were present in six mothers, with the most common being gestational diabetes mellitus (n = 3). Two cases had intrauterine growth restriction and one had cholestasis of pregnancy. Maternal antenatal complications were not different between neonates who developed sepsis (and were admitted to NICU) and those who were asymptomatic [Table 2].

Of the 158 GBS-colonized women, 117 received antibiotic treatment for IAP (74%) and 41 women did not receive IAP due to elective lower segment Caesarean section for other indications or maternal refusal. Women in the IAP group primarily received crystalline penicillin (n = 95; 81.2%), while those allergic to penicillin, received clindamycin (n = 22, 18.8%).

Two women had intrapartum maternal fever without prolonged PROM; both received IAP, and their neonates did not show signs of sepsis. Postpartum complications in GBS-colonized mother were postpartum endometritis (n = 3) and UTI (n = 1), while chorioamnionitis and puerperal sepsis were not seen in any women.

Antibiotic sensitivity profile

The most common antibiotics to which GBS were susceptible were linezolid and vancomycin (97.4% for both) followed by ceftriaxone and cefotaxime (96.2% for both), ampicillin (95.6%), and benzyl penicillin (90.5%). Resistance to clindamycin, erythromycin, levofloxacin, and trimethoprim/ sulfamethoxazole were found to be 57.6%, 57%, 10.1%, and 77.2%, respectively [Table 3].

Neonatal outcome

A total of 12 neonates were admitted to the NICU. The most common age at admission was <6 hours (n = 9; 75%) and most common presentation was respiratory distress (n = 8; 66.6%). Only two cases were culture-positive sepsis (*Staphylococcus aureus*); among others with features of sepsis, the culture were negative.

Five neonates received ampicillin and gentamycin; one of them had hyperbilirubinemia and was managed with phototherapy along with antibiotics. Neonates with *S. aureus* sepsis received vancomycin. One neonate was admitted with bronchopneumonitis at 3 weeks of age and received erythromycin, while four were managed with observation. The mean hospital stay was 6.9 ± 3.2 days and there was no neonatal mortality. Seven cases had received IAP (penicillin); one child with intrauterine growth restriction and prematurity had features of sepsis, but the culture was negative. There was no difference between the groups whose mothers did and did not receive IAP in terms of lengths of hospital stay (P = 0.921) or need for admission in the first hour of life (P = 0.198) [Table 4].

 Table 2: Association of parameters with culture positivity and maternal complications on neonatal outcome

Parameters	Р
Age and culture positive UTI	0.009
Parity and culture positive UTI	0.336
BMI and culture positive UTI	0.077
Presence of other maternal complications in babies admitted	0.819
to NICU compared to babies who were asymptomatic	

BMI – Body mass index, NICU – Neonatal intensive care unit, UTI – Urinary tract infection

Table 3: Antibiotic sensitivity

Antibacterial agent	Susceptible, n (%)
Vancomycin	154 (97.5)
Linezolid	154 (97.5)
Ceftriaxone	152 (96.2)
Cefotaxime	152 (96.2)
Ampicillin	151 (95.6)
Benzyl penicillin	143 (90.5)
Levofloxacin	142 (89.9)
Erythromycin	68 (43)
Clindamycin	67 (42.4)
Trimethoprim/sulfamethoxazole	36 (22.8)

Table 4: Neonatal complications (n=12)

Parameters	Value
Mean±SD gestational age at birth	38 weeks and 4 days±0.757
Mean±SD birth weight (g)	2937.1±353.8
Intrapartum antibiotic prophylaxis	
Received	7
Not received	5
Symptoms	
Respiratory distress	8
Floppy baby with bradycardia	2
Fever	1
Cyanosis	1
Vomiting	1
Mean ± SD age at admission to NICU (h)	9.5±21.3
Treatment received	
Ampicillin + Gentamicin	5
Vancomycin	2
Erythromycin	1
Supportive measures and observation	4
only	
Mean±SD length of hospital stay	6.1±3.2

SD - Standard deviation, NICU - Neonatal intensive care unit

DISCUSSION

The prevalence of maternal GBS colonization rate in this study was 6.9%. In a meta-analysis that included 78 studies published between 1997 and 2015 with 73,791 pregnant women from 37 countries, the overall mean prevalence was 17.9%, with the highest being in Africa (22.4%) followed by Americas (19.7%), Europe (19.0%) and Southeast Asia (11.1%).^[1] Earlier studies from UAE reported prevalence of 10.1% and 21.5%, which is higher than that found in the current study.^[18,19] Recent studies from Middle Eastern countries also show significant variance in the prevalence of GBS colonization.^[13-15,18-24] The varying rates of colonization can be partly explained by difference in socioeconomic groups, education level and acceptance of medical and diagnostic facilities.

The Royal College of Obstetricians and Gynaecologists recommend that clinicians should be aware of risk factors that increases the risk of delivering a child with EOGBS disease. In our study, the prevalence of GBS carriage was significantly higher among those with no risk factors (123/541) than those with at least one of the risk factors (22/1754) (P < 0.00001). A recent study also suggested that testing for GBS with PCR tests among pregnant women with risk factors for EOGBS would reduce the use of IAP by two-thirds compared with the risk-based approach alone.^[25] However, in our study, no neonate had culture-proven EOGBS, and no further analysis was possible due to the small sample size.

The UTI and GBS bacteriuria in a pregnant woman is a marker of heavy genital tract colonization and is an indication of IAP.^[3] In this study, UTI in GBS colonized women was found to be is significantly higher in younger women (aged ≤ 30 years), which is in accordance with another study.^[26] UTI was not found to be related to parity or BMI in our study. Such comparisons were not found in the other studies.

The type of postpartum complications in our study (endometritis and UTI) is in accordance with another study^[3] but is in contrast with others.^[27,28] The association of maternal GBS colonization with adverse pregnancy outcomes such as prematurity, low birth weight, PROM, longer duration of labor reported in earlier studies were not found in the present study.^[3] Further, GBS was not isolated in any of the neonates born to mothers receiving IAP. This is in accordance with other studies and indicates that IAP likely prevents early-onset sepsis.^[3,29]

Although the drug of choice for GBS infection is penicillin, erythromycin and clindamycin are used in those allergic to penicillin. In the current study, resistance of GBS isolates was about 57% for erythromycin, which is much higher than that reported in the literature from different countries (0%-37%).^[29,30] Similarly, clindamycin resistance in our study was 58%, while previous studies from different countries have found it to be 17%-50%.^[29-31] Resistance to penicillin in our study was 10%, which is lower than that recently reported from Iran (about 17%).^[32]

The prospective study design allowed collection of all relevant data regarding the pregnancy outcome and antibiotic sensitivity profile, which is a strength of this study. A major limitation of the study was the relatively small sample size. Finally, as most cases were in women without any risk factors for GBS, the authors recommend a universal screening for GBS in high-income countries such as the UAE to further optimize outcomes.

CONCLUSION

The prevalence of maternal GBS colonization in pregnant women from Ras Al Khaimah, UAE, was found to be low, with carriage significantly higher in women without any risk factors for EOGBS than those with risk factors. GBS was sensitive to most antibiotics, but resistance to erythromycin was high. The study found overall maternal complications were low, and fetal outcome was favorable in women receiving IAP for GBS colonization.

Acknowledgement

The authors are thankful to Dr. Manal Abdulfatah (Director) and Ms. Alphansomma (Laboratory Technician), Pure Health Laboratory, Saqr Hospital, Ras Al-Khaimah, UAE, for their continued help and support throughout the study as well as to Pure Health Laboratory for the administrative support.

Ethical considerations

This study was approved by the Ministry of Health and Prevention Research Ethics Committee/RAK Research and Ethics of Research Sub-Committee at RAK Medical and Health Sciences University, Ras Al-Khaimah, UAE, on December 12, 2018 (Approval no.: MOHAP/RAK/ SUBC/REC/2018/49-2018-MOH-DR). The study was conducted in accordance with the Declaration of Helsinki, 2013. All participants provided written informed consent before sample collection.

Peer review

This article was peer-reviewed by two independent and anonymous reviewers.

Financial support and sponsorship Nil.

Conflicts of interest

There are no conflicts of interest.

REFERENCES

- Kwatra G, Cunnington MC, Merrall E, Adrian PV, Ip M, Klugman KP, et al. Prevalence of maternal colonisation with Group B Streptococcus: A systematic review and meta-analysis. Lancet Infect Dis 2016;16:1076-84.
- Dechen TC, Sumit K, Ranabir P. Correlates of vaginal colonization with Group B streptococci among pregnant women. J Glob Infect Dis 2010;2:236-41.
- Puopolo KM, Lynfield R, Cummings JJ; COMMITTEE ON FETUS AND NEWBORN, COMMITTEE ON INFECTIOUS DISEASES. Management of infants at risk for Group B streptococcal disease. Pediatrics 2019;144:e20191881.
- Khalil MR, Uldbjerg N, Thorsen PB, Møller JK. Risk-based approach versus culture-based screening for identification of Group B streptococci among women in labor. Int J Gynaecol Obstet 2019;144:187-91.
- Kim EJ, Oh KY, Kim MY, Seo YS, Shin JH, Song YR, et al. Risk factors for Group B Streptococcus colonization among pregnant women in Korea. Epidemiol Health 2011;33:e2011010.
- Chen J, Fu J, Du W, Liu X, Rongkavilit C, Huang X, *et al.* Group B streptococcal colonization in mothers and infants in western China: Prevalences and risk factors. BMC Infect Dis 2018;18:291.
- Muller-Pebody B, Johnson AP, Heath PT, Gilbert RE, Henderson KL, Sharland M, *et al.* Empirical treatment of neonatal sepsis: Are the current guidelines adequate? Arch Dis Child Fetal Neonatal Ed 2011;96:F4-8.
- Depani SJ, Ladhani S, Heath PT, Lamagni TL, Johnson AP, Pebody RG, et al. The contribution of infections to neonatal deaths in England and Wales. Pediatr Infect Dis J 2011;30:345-7.
- O'Sullivan CP, Lamagni T, Patel D, Efstratiou A, Cunney R, Meehan M, et al. Group B streptococcal disease in UK and Irish infants younger than 90 days, 2014-15: A prospective surveillance study. Lancet Infect Dis 2019;19:83-90.
- Heath PT, Schuchat A. Perinatal Group B streptococcal disease. Best Pract Res Clin Obs Gyn 2007;21:411-24.
- 11. Plumb J, Clayton G. Group B *Streptococcus* infection: Risk and prevention. Pract Midwife 2013;16:27-30.
- Bedford H, de Louvois J, Halket S, Peckham C, Hurley R, Harvey D. Meningitis in infancy in England and Wales: Follow up at age 5 years. BMJ 2001;323:533-6.
- Hadavand S, Ghafoorimehr F, Rajabi L, Davati A, Zafarghandi N. Frequency of Group B streptococcal colonization in pregnant women aged 35- 37 weeks in clinical centers of Shahed University, Tehran, Iran. Iran J Pathol 2015;10:120-6.
- Musleh J, Al Qahtani N. Group B *Streptococcus* colonization among Saudi women during labor. Saudi J Med Med Sci 2018;6:18-22.
- Clouse K, Shehabi A, Suleimat AM, Faouri S, Khuri-Bulos N, Al Jammal A, *et al.* High prevalence of Group B *Streptococcus* colonization among pregnant women in Amman, Jordan. BMC Pregnancy Childbirth 2019;19:177.
- Rahbar M, Hajia M, Mohammadzadeh M. Urinary tract infections caused by GBS in adult women: Survey of 11,800 urine culture results. Iran J Pathol 2012;7:32-7.
- Wayne PA. Performance Standards for Antimicrobial Susceptibility Testing. 27th ed. Wayne, PA, USA: CLSI Supplement M100; Clinical and Laboratory Standards Institute; 2017.
- 18. Amin A, Abdulrazzaq YM, Uduman S. Group B streptococcal serotype

AlZuheiri, et al.: Maternal GBS colonization in UAE

distribution of isolates from colonized pregnant women at the time of delivery in United Arab Emirates. J Infect 2002;45:42-6.

- Sidky I, Thomas M. Prevalence of Group B streptococcal infection colonisation in pregnant women and their offspring in the Middle East. J Obstet Gynaecol 2002;22:179-80.
- Hakim M, Jabour A, Anton M, Hakim M, Kheirallah S. Screening Arab Israeli pregnant women for Group B *Streptococus* by the AmpliVue GBS assay: Are the rates higher than the national average? Isr Med Assoc J 2018;20:291-4.
- Ghaddar N, Alfouzan W, Anastasiadis E, Al Jiser T, Itani SE, Dernaika R, *et al.* Evaluation of chromogenic medium and direct latex agglutination test for detection of Group B *Streptococcus* in vaginal specimens from pregnant women in Lebanon and Kuwait. J Med Microbiol 2014;63:1395-9.
- Moraleda C, Benmessaoud R, Esteban J, López Y, Alami H, Barkat A, et al. Prevalence, antimicrobial resistance and serotype distribution of Group B Streptococcus isolated among pregnant women and newborns in Rabat, Morocco. J Med Microbiol 2018;67:652-61.
- Khan MA, Faiz A, Ashshi AM. Maternal colonization of Group B Streptocecus: Prevalence, associated factors and antimicrobial resistance. Ann Saudi Med 2015;35:423-7.
- Alp F, Findik D, Dagi HT, Arslan U, Pekin AT, Yilmaz SA. Screening and genotyping of Group B *Streptococcus* in pregnant and non-pregnant women in Turkey. J Infect Dev Ctries 2016;10:222-6.
- Rosenberg LR, Normann AK, Henriksen B, Fenger-Gron J, Møller JK, Khalil MR. Risk-based screening and intrapartum Group B *Streptococcus* polymerase chain reaction results reduce use of antibiotics during labour. Dan Med J 2020;67:A06200460.

- Bidgani S, Navidifar T, Najafian M, Amin M. Comparison of Group B streptococci colonization in vaginal and rectal specimens by culture method and polymerase chain reaction technique. J Chin Med Assoc 2016;79:141-5.
- Ngonzi J, Bebell LM, Bazira J, Fajardo Y, Nyehangane D, Boum Y, et al. Risk factors for vaginal colonization and relationship between bacterial vaginal colonization and in-hospital outcomes in women with obstructed labor in a Ugandan regional referral hospital. Int J Microbiol 2018;2018:6579139.
- Goel N, Wattal C, Gujral K, Dhaduk N, Mansukhani C, Garg P. Group B *Streptococcus* in Indian pregnant women: Its prevalence and risk factors. Indian J Med Microbiol 2020;38:357-61.
- Dilrukshi GN, Kottahachchi J, Dissanayake DM, Pathiraja RP, Karunasingha J, Sampath MK, *et al.* Group B *Streptococcus* colonisation and their antimicrobial susceptibility among pregnant women attending antenatal clinics in tertiary care hospitals in the Western Province of Sri Lanka. J Obstet Gynaecol 2021;41:1-6.
- Sharmila V, Joseph NM, Arun Babu T, Chaturvedula L, Sistla S. Genital tract Group B streptococcal colonization in pregnant women: A South Indian perspective. J Infect Dev Ctries 2011;5:592-5.
- Castellano-Filho DS, da Silva VL, Nascimento TC, de Toledo Vieira M, Diniz CG. Detection of Group B *Streptococcus* in Brazilian pregnant women and antimicrobial susceptibility patterns. Braz J Microbiol 2010;41:1047-55.
- Daramroodi AK, Keshavarzi F. The investigation of antibiotic resistance and rapid detection of Group B *Streptococcus* (Bca) from vaginal specimens of pregnant women by colony PCR method. J Bas Res Med Sci 2018;5:27-32.