

colonization was not associated with endometritis or wound infection among women who delivered by cesarean.

CONCLUSION: In contrast to earlier data prior to routine intrapartum antibiotic prophylaxis, colonization with GBS was associated with slightly lower odds of chorioamnionitis, but was not associated with postpartum wound infection or endometritis.

LEARNING OBJECTIVES: To understand the implications of intrapartum antibiotic prophylaxis for GBS on maternal infectious morbidity in pregnancy and the postpartum period.

29 Management of fever in labor after institution of a standardized order set at a maternity quaternary care center



C. Wang¹, I. Sirluck Schroeder¹, A. Sosa Cazales², R. Kim², A. Albert³, C. Elwood^{2,3}, J. Van Schalkwyk^{2,3}

¹UBC Faculty of Medicine, Vancouver BC, ²Department of Obstetrics and Gynecology, University of British Columbia, Vancouver, BC, ³Women's Health Research Institute, BC Women's Hospital, Vancouver, BC

OBJECTIVES: Maternal fever in labour (FIL) is common, and can be the first sign of chorioamnionitis and sepsis. Our institution initiated a standardized order set for the management of FIL. This study evaluated maternal and fetal parameters in women with FIL in order to determine trends in antibiotic use, the applicability of obstetrical scoring system for sepsis and outcomes for women.

METHODS: An interim analysis of the retrospective chart review on 510 patients between 2011-2016 for which the FIL protocol was initiated was performed. 403 charts were reviewed due to clinical concerns of increasing gram negative bacteremia with antimicrobial resistance and poor compliance with blood cultures being drawn. Antenatal history, intrapartum parameters and maternal/fetal outcomes were evaluated. Categorical variables were compared using Fisher's exact tests and continuous variables with Wilcoxon rank sum tests.

RESULTS: The median maternal age was 31.5 years (SD4.6 years), median gestational age 39.9 weeks (IQR39-40.6), and a majority of women were nulliparous (81.9%). The median maternal temperature at the time of first fever was 38.1°C (IQR38.0-38.3), and 90.8% report epidural use. Antibiotics were administered 74% of the time when the order set was initiated, with 95% of antibiotic administration being a combination of metronidazole and cefazolin. At time of first fever, 31.8% reported concurrent fetal tachycardia, and a larger proportion of those women were subsequently given antibiotics (88vs72%, p=0.0007). Conversely, a larger proportion of those without fetal tachycardia were given acetaminophen vs those without (82vs72% p=0.032). Only 16.7% of women has blood cultures drawn, however women with blood cultures had a slightly higher first temperature; 38.2°C (IQR 38.0-38.4) vs 38.1°C (IQR 38.0-38.2) in those without blood cultures (p=0.0004). Histologically diagnosed chorioamnionitis was associated with higher initial temperature, 38.2°C vs 38.1°C without chorioamnionitis (p=0.002). There was one maternal transfer to ICU and 3 fetal deaths.

CONCLUSION: Our preliminary results identified only 16.7% of women having blood cultures drawn with initiation of antibiotic therapy for FIL with higher initial maternal temperature correlating with an increased likelihood of them being drawn. We found that fetal tachycardia was associated with antibiotic use and less acetaminophen use. This interim analysis demonstrated that between 2011-2016 few women were routinely having blood cultures drawn with initiation of antibiotic therapy.

LEARNING OBJECTIVES: Evaluate standardized change in practice and management of maternal FIL at an institutional level and identify areas of improvement.

30 Simple and effective screening for Chagas disease at the prenatal intake visit



C. D. Yarrington¹, D. A. Hamer¹, E. Barnett¹, I. Camelo¹, J. H. Perez², M. Poorvu², J. R. Koehler³, N. S. Hochberg¹

¹Boston Medical Center, Boston MA, ²East Boston Neighborhood Health Center, East Boston MA, ³Childrens Hospital of Boston, Boston, MA

OBJECTIVES: An estimated 300,000 individuals have Chagas disease in the United States and Chagas is passed vertically at rates higher than syphilis, there is still not routine perinatal Chagas screening. We sought to evaluate the success of a simple screening program to identify maternal Chagas disease during prenatal care.

METHODS: This was a screening program of a cohort of women who had prenatal care at a community health center that serves a large Latinx population, all of whom delivered at a central hospital. Prenatal screening was done by asking all women at the prenatal intake where they were born, and if not born in a high risk area, whether they had ever spent 6 or more months continuously in Mexico, Central or South America. A positive answer to either or both questions prompted IgG screening. The practice of implementing routine screening for Chagas began mid-2017. We identified the women who delivered from February 2018 to March 2019, assuming initiation of prenatal care at twelve weeks and expecting a six month ramp up in universal screening. We evaluated the rate of identification of preliminary and confirmed positives in this cohort. Further, we evaluated connections with infectious disease and cardiology for the mother as well as CDC standard of care for evaluation of the newborn, when available.

RESULTS: A total of 619 women that delivered were screened for Chagas disease; IgG testing yielded 21 preliminary positive but confirmed negative results, 3 preliminary indeterminate results with subsequent negative confirmatory testing, and 3 confirmed positive results from the CDC. The prevalence of confirmed Chagas in the entire population was 0.5%. All three confirmed positive women have had normal cardiac evaluation. Of the three confirmed cases, one has delivered and both mother and infant have had follow up care in Infectious Disease.

CONCLUSION: A two question screen for Chagas disease risk at the initial prenatal visit is effective, and in select populations is high yield. Capitalizing on a time of insurance coverage and healthcare engagement, identification of Chagas in pregnancy can not only identify infants at risk of vertical transmission but may mitigate the complications of long term infection.

LEARNING OBJECTIVES: Learners will be able to describe a simple work flow that increases identification of Chagas in mothers as well as infants at risk of vertical transmission. Learners will be able to motivate their prenatal intake providers to add a single question to the new visit by empowering them with a realistic expectation of capturing women at risk and improving the health of both mother and child.

31 Successful linkage to Chagas care via screening at prenatal intake



C. D. Yarrington, M. Poorvu, C. S. Hochberg, D. W. Hamer

OBJECTIVES: While congenital Chagas occurs at rates higher than congenital syphilis, and prenatal care is a known time to capture chronic conditions while a woman is insured and engaged in care, there is still not routine perinatal screening. We sought to evaluate