



Outbreak of a Late-Onset Group B Streptococcus Cluster due to Probable Horizontal Transmission in a Level 3 Neonatal Intensive Care Unit

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Introduction

- Group B Streptococcus (GBS) is a major pathogen among neonates and young infants.
 - Early onset --occurs within the first week of life, associated with pregnancy-related risk factors.
 - Late-onset --occurs after the first week of life through 3 months, sporadic.
 - Outbreaks of late-onset (GBS) are rare.
- ❖ In February 2017, our 20 bed Neonatal Intensive Care Unit (NICU) identified 2 preterm infants with late-onset GBS infections occurring within a 5 day period, prompting an outbreak investigation and report of a probable link.

Methods

Case Identification

- ❖ A case was defined as culture-confirmed invasive GBS infection in a preterm infant <37 weeks gestational age (GA).
- Laboratory records for the 4 weeks prior to and 12 weeks following the index case were reviewed to identify additional cases.

Analysis of GBS Isolates

- Antibiograms compared
- ❖ All 3 isolates referred to NYS DOH Wadsworth Lab for pulse field gel electrophoresis (PFGE).

Other Investigation

- Hand hygiene compliance
- Nursery conditions census, floor plan, incubator proximity
- Breast milk handling & storage
- Disinfection of reusable equipment
- Environmental Services (EVS) procedures

Results

Case Reports

Case 1 (Index Case)

- ❖ 34 6/7 week preterm male, birthweight 855 g, born by caesarian section to a mother with unknown GBS status who developed fever on day of life (DOL) 25, followed by erythema & induration c/w cellulitis below the umbilicus and extending over the groin and buttocks.
- Blood culture grew GBS
- CSF culture was no growth
- Hip/pelvic x-ray, ultrasound, MRI c/w cellulitis plus myositis left anterior thigh muscles, no evidence of septic arthritis.

Case 2

- ❖ 27 5/7 week preterm female birthweight 860 g, born by vaginal delivery to a GBS-negative mother, who developed lethargy and apnea requiring intubation on DOL 39.
- ❖ Blood culture grew GBS (Pen MIC ≤ 0.06)
- CSF culture also positive for GBS.

Lab Record Review

No additional GBS cases

Analysis of GBS Isolates

<u>Antibiograms</u>

- Isolates from the 2 infants had the same susceptibility patterns.
- All were penicillin susceptible, erythromycin resistant and clindamycin susceptible without inducible resistance.

PFGE

PFGE patterns for the 3 isolates were indistinguishable and the isolates were considered to be the same strain.

Results

Hand Hygiene Compliance

Compliance measured by anonymous observer had fallen to <90% at the time of the outbreak.</p>

Nursery Conditions

- Average daily census was high, more than 30% above recommended occupancy, causing overcrowding, decreased distance between incubators.
- Incubators of Case 1 & Case 2 were adjacent

Breast Milk Handling & Storage

No lapses in breast milk storage protocol were identified.

Interventions/Control Measures

- Contact precautions
- -- maintained until repeat cultures negative, antibiotics completed.
- Breast milk for the affected infants was stored in a separate refrigerator.
- Enhanced focus on hand hygiene
- Focus on cleaning of reusable equipment
- Focus on compliance with EVS procedures
- CleanTrace/ATP as a monitoring aid and rapid feedback and follow up for failures

Conclusions

- No environmental factor or clear mode of transmission were identified; however, horizontal transmission was highly suspected.
- The average NICU census during this time was 27, exceeding the maximum of 20 patients.
- High census and possible transient hand carriage of GBS by the staff may have played a role in the transmission of GBS.
- Adherence to hand hygiene and isolation practices were essential in containing the outbreak and preventing further transmissions.

References

- 1. Franciosi RA, Knostman JD, Zimmerman RA. Group B streptococcal neonatal and infant infections. J Pediatr 1973;82:707-18.
- 2. Paredes A, Wong P, Mason E, Taber L, Barrett F. Nosocomial transmission of group B streptococci in a newborn nursery. Pediatrics 1977;59:679-82.
- B. MacFarquhar JK, Jones TF, Woron AM, et al. Outbreak of late-onset group B streptococcus in a neonatal intensive care unit. Am J Infect Control 2010;38:283-8.
- Kotiw M, Zhang GW, Daggard G, Reiss-Levy E, Tapsall JW, Numa A. Late onset and recurrent neonatal group B streptococcal disease associated with breast-milk transmission. Pediatr Dev Pathol 2003;6:251-6.
- 5. Green PA, Singh KV, Murray BE, Baker CJ. Recurrent group B streptococcal infections in infants: Clinical and microbiological aspects. J Pediatr 1994;125:931-8.
- . Wang LY, Chen CT, Liu WH, Wang YH. Recurrent neonatal group B streptococcal disease associated with infected breast milk.Clin Pediatr (Phila) 2007;46:547-9.