



IV Antibiotic Audit

June 2014-July2014



Tracey Green, NNP
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Background

- ▶ “Suspected sepsis”
- ▶ One of the most common diagnoses made in the NICU
- ▶ Early-onset sepsis (EOS) a major cause of neonatal mortality and morbidity, particularly among preterm/very low-birth weight infants
- ▶ Low-incidence, high-consequence disease
- ▶ Clinical signs may be non-specific/absent in the immediate postnatal period.



Objectives

- ▶ To identify whether antibiotic use follows current antibiotic protocol
- ▶ To determine if antibiotic exposure has decreased following the implementation of the current antibiotic protocol
- ▶ Was maternal antibiotic cover appropriate according to current CWH guidelines
- ▶ Differentiate between well infants at risk of sepsis vs symptomatic babies with presumed sepsis



Policy: Early Onset Sepsis

- ▶ EOS defined: Infection within the first 72 hours following birth
- ▶ Treated with empiric antibiotics until sepsis is excluded
- ▶ Group B streptococcus (GBS) - most common cause alongside Escherichia coli.
- ▶ Maternal and infant clinical characteristics
- ▶ Infant laboratory values are utilised to determine newborns at risk



Risk Factors for EOS

- ▶ Prolonged rupture of membranes
- ▶ Maternal illness - eg. pyrexia >38.0 , raised CRP
- ▶ Pathogens (e.g. GBS, E. coli) present in maternal urine or high vaginal swab
- ▶ Prematurity < 37 weeks
- ▶ Fetal distress, tachycardia >160 bpm or neonatal depression
- ▶ Twin gestation

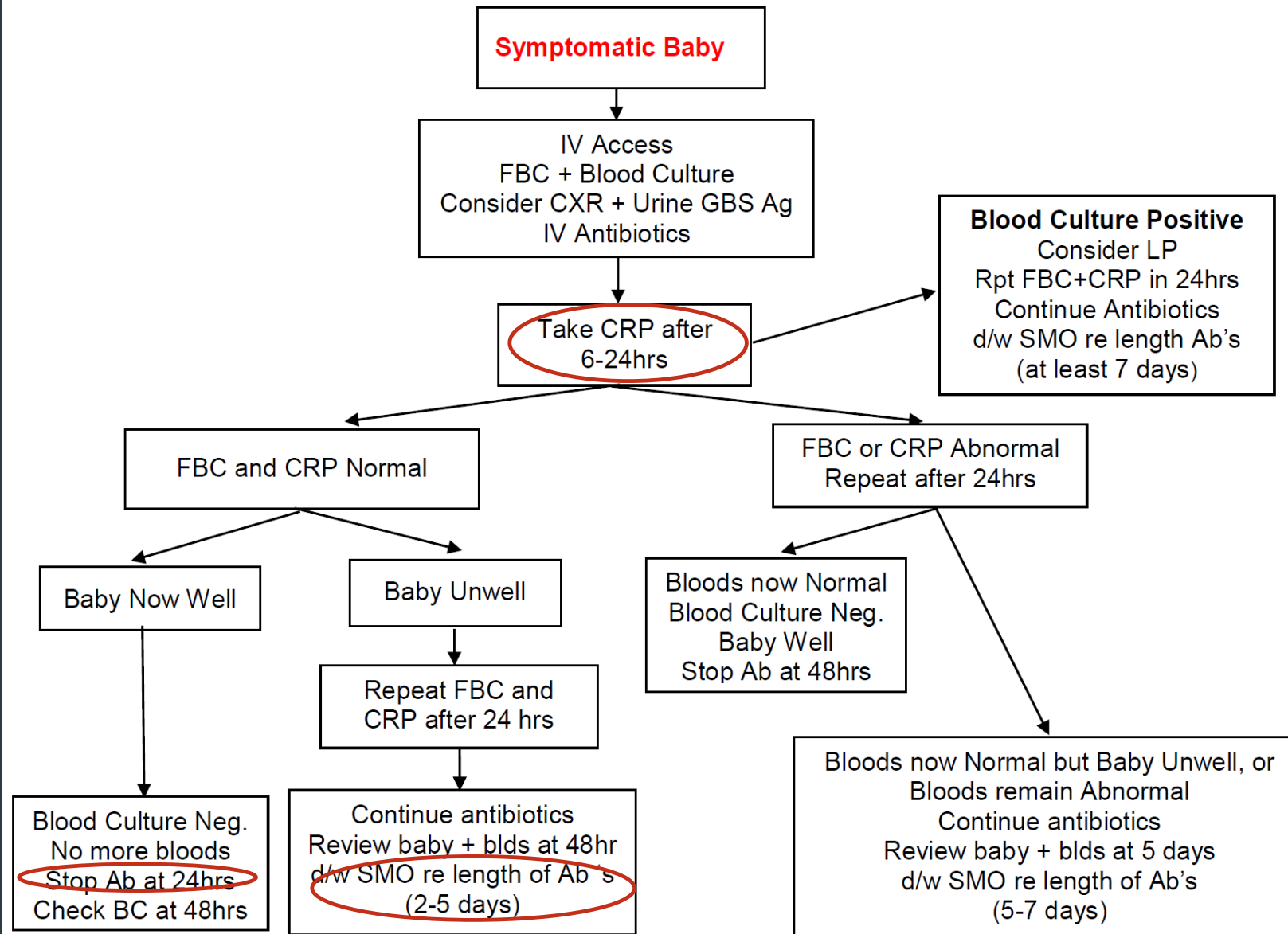


Criteria for Commencing Antibiotics

- ▶ All newborn infants with early respiratory distress
 - ▶ No infant should be untreated after 4 hrs of age
- ▶ Temperature instability
- ▶ Apnoea, especially new onset or increased frequency or severity in a premature infant
- ▶ Listlessness, lethargy, pallor, mottling and irritability
- ▶ Jaundice if it develops unusually rapidly
- ▶ Ileus (abdominal distension or bilious vomiting/nasogastric aspirate)
- ▶ Previously healthy baby who refuses to feed



Management of the Symptomatic Baby at Risk of Sepsis



Method

- ▶ A retrospective audit of clinical records for all infants admitted to the CWH NICU (established from the NICU database), requiring antibiotic treatment during the first 72hrs of life between June 2014 & July 2014.



Results

Clinical records for 58 babies were reviewed

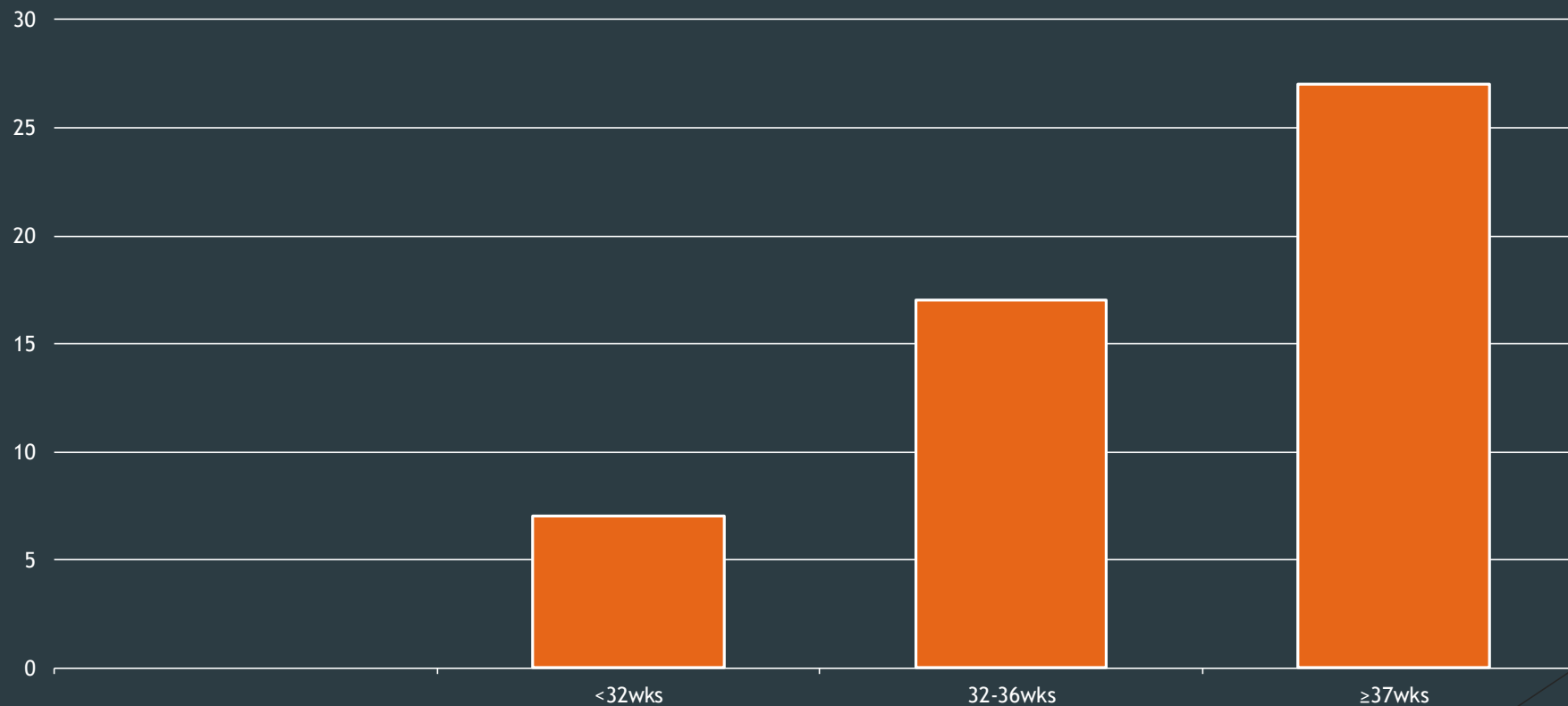
- ▶ 7 excluded leaving 51 babies
 - ▶ 4 Surgical
 - ▶ 1 Day 5 admission excluded
 - ▶ 1 Day 10 Rhinovirus from paediatrics excluded
 - ▶ 1 Transfer from Dunedin at 2wks of age
- ▶ Median birth weight was 2910g (Quartiles 2255-3840g).



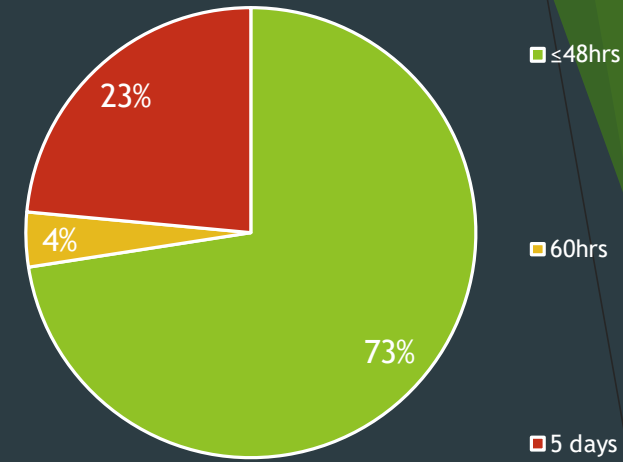
Results

	<32wks	32-36wks	>37wks
Received Antibiotics (51)	7	10	34
Admitted No Antibiotics (30)	1	16	13
Maternal Abs in labour	5	6	15
Confirmed Maternal Chorioamnionitis	2	0	4

Gestational Age of Babies Requiring Antibiotics: Mean Gestational 37wks

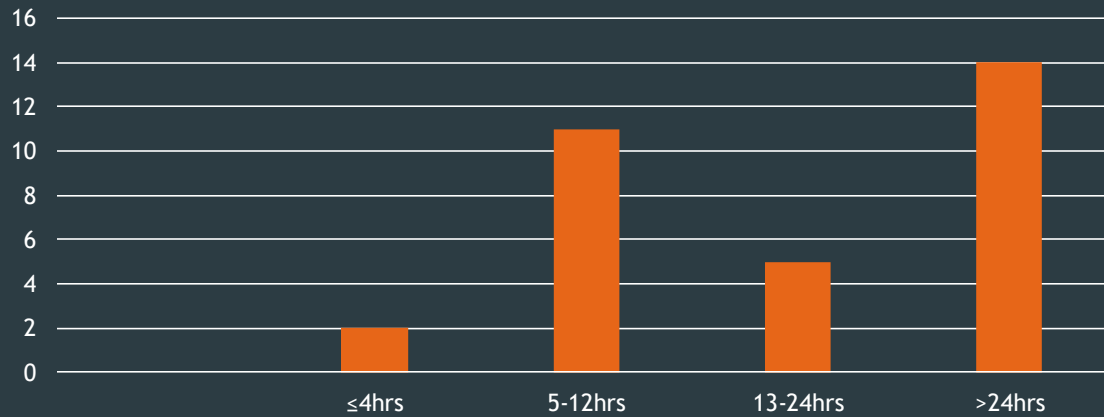


Neonatal Antibiotic Exposure



Duration of Antibiotics	<32wks	32-36wks	≥37wks
24 hrs	5	5	8
36 hrs		4	5
48 hrs	2	6	2 + x2 5doses
3-5 days		2	10

Duration of Respiratory Support (Ventilation, CPAP, Oxygen)



All <32wk infants required respiratory support

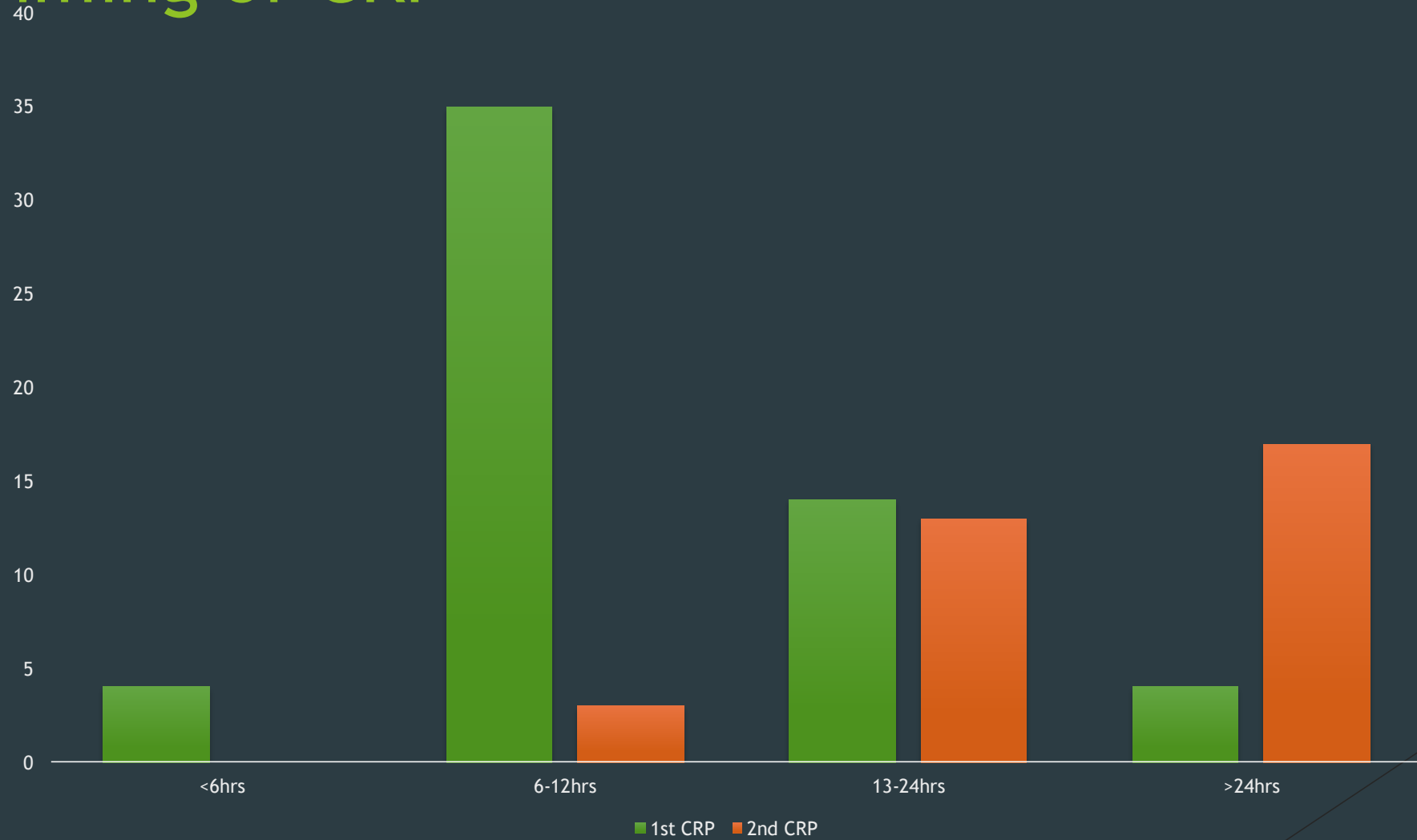
80% <37wk infants

63 % ≥37wk infants

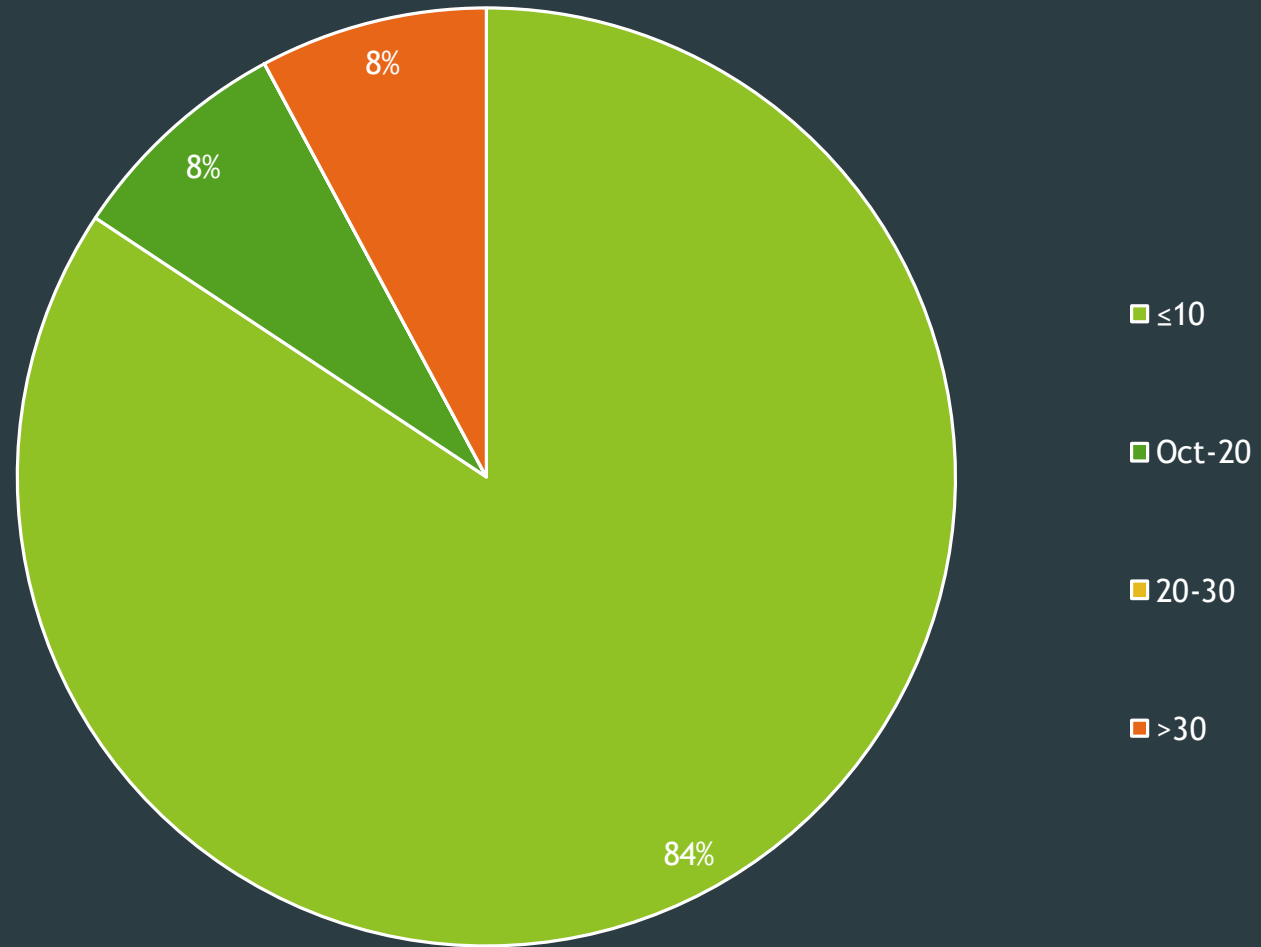
Other Reasons:

Tachypnoea, meconium exposure, thrombocytopenia, dehydration, CXR changes

Timing of CRP



1st CRP Result



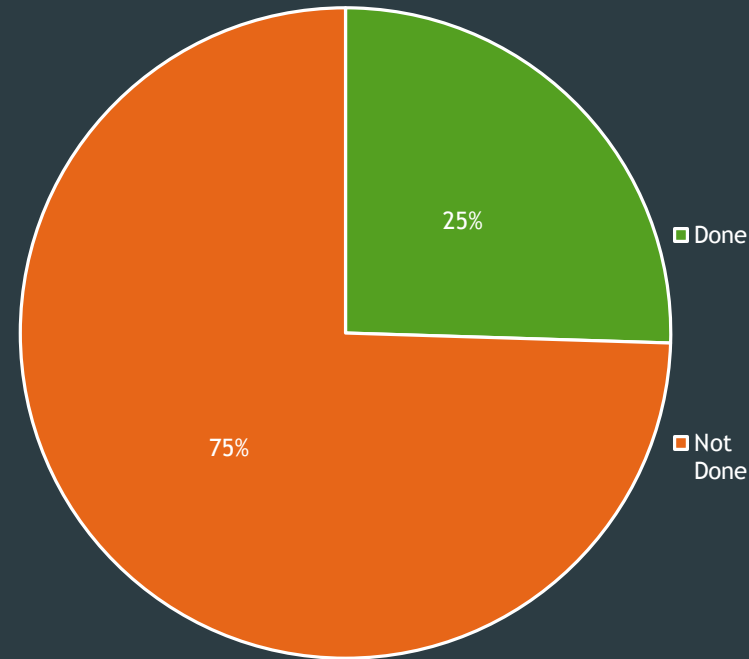
CRP Considerations

- ▶ An acute phase reactant synthesised within 6-8 hours in response to tissue injury
- ▶ Non-infectious processes can also elevate the CRP
- ▶ Levels peak at 24-48 hours
- ▶ A normal CRP at the start of an illness/at birth lacks the sensitivity to rule out sepsis
- ▶ If taken at >6hrs the sensitivity improves to >90%
- ▶ A level of <10mg/L is considered normal and has a negative predictive value of 99% for EOS

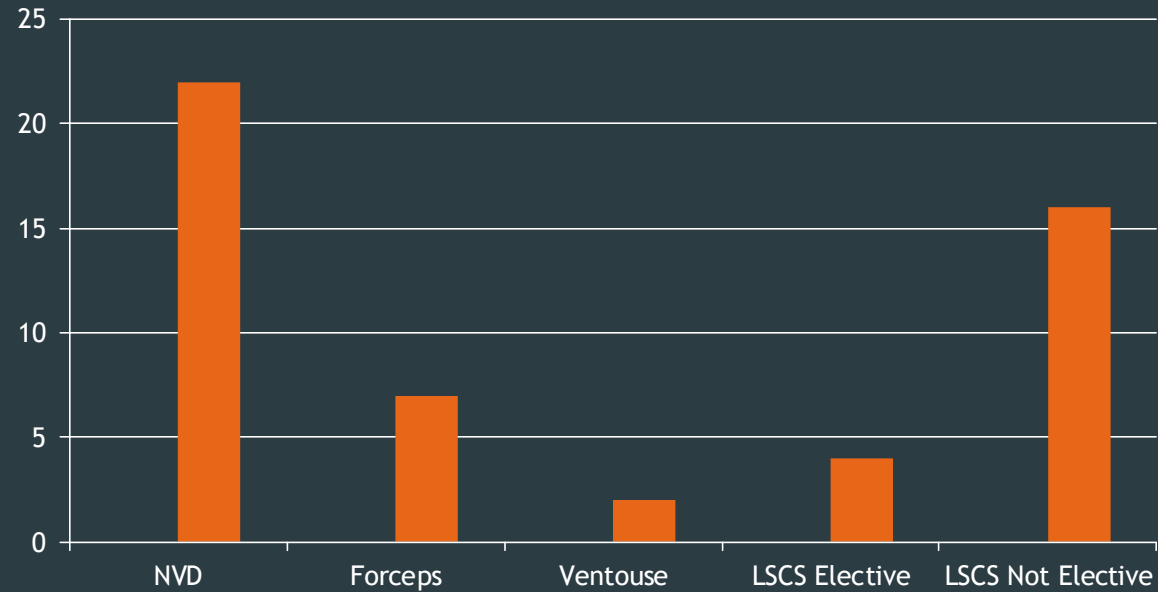


GBS Antigen Protocol

- ▶ A urine should be sent for GBS antigen
- indicates systemic GBS infection
- ▶ If antibiotics are stopped after 24-48hrs
and the baby is on the postnatal ward
then it can be omitted



Delivery Type



Non Elective LSCS

- ▶ X2 Abruptio
- ▶ X2 Breech
- ▶ X1 set twins bulging membranes,
- ▶ x1 transverse, thinning scar
- ▶ X1 FTP
- ▶ X1 Post dates not in labour

Electives LSCS

- ▶ X1 GDM on insulin
- ▶ X1 severe IUGR 32wks
- ▶ X2 39wks

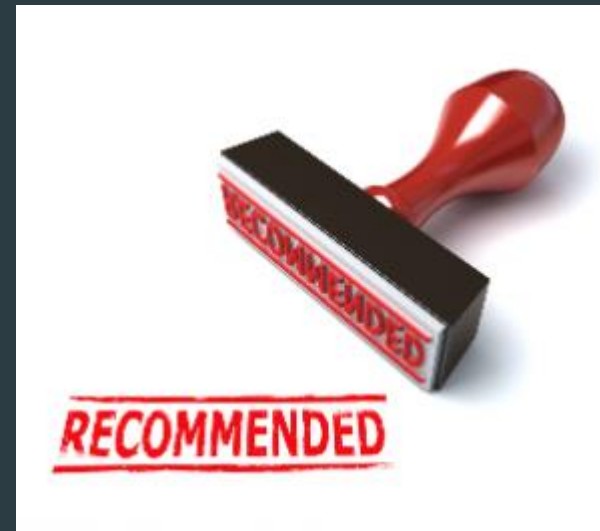
Summary: Clinical Implications

- ▶ Early antibiotics for many are correct response
- ▶ Overall clinical picture important factor
- ▶ Reduced antibiotic exposure without compromising safety
- ▶ Appropriate and safe antibiotic treatment
- ▶ Decreased unnecessary intervention and antibiotic resistance.
- ▶ Joint responsibility for ensuring antibiotic review
- ▶ Decrease cost without increased risk to neonates.
- ▶ Reduce staff workload.



Future Recommendations

- ▶ Capture postnatal babies requiring antibiotics
- ▶ Reduce antibiotics exposure further
- ▶ Protocol review
- ▶ Determine Nationwide practices
- ▶ Stop antibiotics in timely manner
- ▶ Determine urine GBS antigen requirement
- ▶ Utilise evidence from this audit to further review current practice



International Data

- ▶ 36hrs minimum cover
- ▶ Use of Sepsis Calculators
- ▶ Limited research and supporting evidence determining appropriate IV antibiotic course duration
- ▶ NICE guidelines recommend treating neonates with risk factors but clinically well for 36hrs



Conclusion

- ▶ Clinical notes of 58 infants were reviewed of which 51 meet the required criteria
- ▶ Essentially antibiotics were given according to the new protocol
- ▶ Improvements can be made
- ▶ There is an ability to decrease antibiotic exposure while maintaining safety



