Utility of Hematological Scoring System in Diagnosis of Neonatal Sepsis

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ABSTRACT

Background: Neonatal septicemia is defined as a clinical syndrome characterized by systemic signs and symptoms caused by a bacterial infection and gives positive blood culture in the first month of life. It is associated with high morbidity and mortality, but early diagnosis and treatment significantly improve the outcomes. The present study brings out a quick and cost-effective Hematological Scoring System that enables early diagnosis of neonatal sepsis.

Method: This study was conducted in the Department of Pathology of Sree Gokulam Medical College and Research Foundation, Venjaramoodu, Thiruvananthapuram over 1 year. Eighty neonates with clinical suspicion of sepsis were studied with respect to their peripheral smear findings, blood culture and C-reactive protein levels.

Result: Among the eighty cases, twelve neonates were culture positive. Male gender and late-onset of sepsis were the significant risk parameters. Escherichia coli and Staphylococcus aureus were the most common isolated organisms. Of the different parameters studied I:T ratio and Absolute neutrophil count showed the highest specificity and immature neutrophil count had the highest sensitivity. Forty-one percent of neonates had a high Hematological Scoring System score. The specificity of Hematological Scoring System with a score > 5 was 92%.

Conclusion: Hematological Scoring System is a cost-effective, rapid and easy to perform screening test that can be used to rule out sepsis thus avoiding unnecessary administration of antibiotics to unaffected babies. It should be adopted as a routine screening procedure by minimally qualified rural doctors with minimal resources to ensure appropriate action immediately for children with high index of suspicion of sepsis.

Keywords: Blood Culture, WBC Count, Early Onset Sepsis (EOS), I/T Ratio

Introduction

Neonatal septicaemia is defined as a clinical syndrome characterized by systemic signs and symptoms due to generalized bacterial infection with positive blood culture in the first four weeks of life.[1] Accurate clinical diagnosis of neonatal septicaemia is an involved task. The early signs and symptoms are non-specific; however, treatment has to be started immediately, an outcome of septicaemia in neonate largely depends on its early identification and treatment.[2] Although blood culture is considered to be the gold standard for diagnosis, the inherent requirement of sophisticated laboratory equipment makes it less available for resource-poor countries. In addition, blood culture requires a minimum of 48-72 hours and yields positive results in only 30-40% of cases.[3] Also the inability to isolate a microbial pathogen does not exclude sepsis.[4] Therefore, the necessity for a rapid and cost-effective test, with minimal time consumption is prevalent. An ideal diagnostic test for neonatal sepsis should have maximum sensitivity and specificity. In recent years, various investigators have evaluated some highly sensitive and specific inflammatory markers to diagnose neonatal sepsis. Although these markers are sensitive and specific, they also require sophisticated and expensive methodologies thereby making it impractical for developing countries.[5]

A comprehensive study of hematological parameters together as a combination of tests provided early predictions of neonatal septicemia. This combination study resulted in both increased sensitivity and specificity, thereby aiding in early treatment with appropriate antibiotics.[6] Rodwell et. al. developed a seven-point hematologic scoring system based on the WBC count, total and immature neutrophil counts and ratios, degenerative changes in neutrophils and thrombocytopenia. This approach proved to be effective with 96% sensitivity and 99% negative predictive value was obtained.[6]

The present study evaluated the effectiveness of haematological profile in early diagnosis of neonatal sepsis.

Materials and Methods

This was a prospective cross-sectional study conducted in the Department of Pathology, over a period of 1 year from
November 2014 to November 2015 with prior approval from institutional Ethical committee. Neonates up to thirty days of age were enrolled in the study if there were clinical suspicion of sepsis or any potential risk factors for sepsis like prematurity or prolonged rupture of membrane included in the study. Peripheral smear from severely jaundiced neonates, due to blood group incompatibilities, were not included in the study. Blood sample (0.5 -1.0 ml) obtained by peripheral venipuncture was collected in EDTA vial. A good quality peripheral smear prepared within 1 hour of venepuncture, stained with Leishman and Giemsa stains were examined under oil immersion with 100x magnification. In addition total leukocyte and platelet counts were counted by ADVIA 2120i automated system. Differential leukocyte counts were performed on Leishman and Giemsa stained blood smears by counting 100 cells. Immature neutrophils (promyelocyte, myelocyte, metamyelocyte and band forms) and degenerative morphological changes in neutrophils including toxic granulations, vacuolations and Dohle bodies were noted. The hematological findings were analysed according to HSS of Rodwell et al. HSS assigns a score of 1 for each of seven findings significantly associated with sepsis. Abnormal total leukocyte count, abnormal PMN count, elevated immature to total (I: T) PMN ratio, immature to mature(I:M) PMN ratio, platelet count(<150,000/mm3) and degenerative changes in PMNs. An abnormal total PMN count is assigned score of two instead of 1, if no mature polymorphs are seen on the peripheral smear to compensate for the low I:M ratio. Score of two and <2 was interpreted as sepsis is unlikely, score 3-4 asepsis is possible and score 5 and above as sepsis is very likely.

C-reactive protein levels were also recorded from Immunology laboratory and quantitative evaluation of CRP done.

Blood samples for culture and sensitivity was sent from NICU to Department of Microbiology in specifically provided sterile blood culture bottles and reports obtained after 72 hours. The diagnosis of sepsis was confirmed when there were positive findings in blood culture.

Data Analysis
Data analysis was carried out in Microsoft excel. Data was filtered, coded and analysed using Chi-square test for quantitative variables in SPSS version 16 (Statistical Package for Social Sciences). In all the tests p-value of less than 0.05 was taken to be statistically significant and values less than 0.01 was taken highly significant.

Result
Among the 80 cases studied, 12 cases indicated a positive culture while the remaining 85% indicated a negative culture. The two groups were analyzed for baseline parameters, bacteriological Profile and hematologic Parameters (Table 1,2).

Males predominated females (3:1) in the culture-positive group. Late-onset sepsis (more than 72 hours) was more frequent (58%). There was no significant impact of birth weight with both groups being represented equally. Pre-term babies were under-represented in the study (9/80) and thus contributed to only 33% of the culture-positive cases.

The most common causative organism in the study were E. coli and Staphylococcus aureus (Fig 4) each of which attributed to 25% followed by Streptococcus pneumonia and Group B beta Streptococci (GBS) which constituted 16% each.

Abnormal TLC (Leukocytosis/Leukopenia) was seen in 67% of the culture-proven cases of which >50% showed leukocytosis (Fig. 1) and the remaining indicated leucopenia. 34% of culture-positive babies had a normal leukocyte count.

Absolute Neutrophil Count (<2000) was seen in 66% of the culture- positive cases whereas it was seen only in 6% of the culture-negative group.

Immature Neutrophil Count- 75% culture positive cases had an elevated immature neutrophil count.

Both I:T (< 0.2) and I:M ratio (>0.3) were significantly associated with culture positive group. (P-value <.002)

Degenerative changes All the culture positive cases showed degenerative changes in neutrophils. The most common degenerative changes seen were toxic granulations followed by cytoplasmic vacuolations (Fig. 2).

Platelet Count- None of the culture-positive cases showed platelet count <1.5L but 8% of culture-negative neonates exhibited thrombocytopenia (Fig. 3)

C Reactive Protein- Only 9% of culture positive cases gave nonreactive CRP and found to be statistically significant (p-value 0.011).

HSS- 41% of culture-positive neonates have a Hematological score of >5, which is suggestive of sepsis. 33% of culture- positive shows a score which indicates probable sepsis and 26% cases gives a score suggesting that sepsis is unlikely. Only 7% of culture-negative cases indicated a hematological score suggestive of sepsis.

Discussion
Neonatal sepsis is a serious illness with high morbidity as well as mortality. Early diagnosis with prompt antibiotic therapy can significantly improve the outcome. So, a
Table 1: Hematological profile.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Culture Positive (N=12)</th>
<th>Culture Negative (N = 68)</th>
<th>p Value &lt; 0.05</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abnormal TLC</td>
<td>8 (67%)</td>
<td>11 (16%)</td>
<td>YES</td>
</tr>
<tr>
<td>ANC &lt; 2000</td>
<td>8 (67%)</td>
<td>4 (06%)</td>
<td>YES</td>
</tr>
<tr>
<td>Increased immature PMN</td>
<td>9 (75%)</td>
<td>11 (16%)</td>
<td>YES</td>
</tr>
<tr>
<td>I:T &gt; 0.2</td>
<td>9 (75%)</td>
<td>13 (19%)</td>
<td>YES</td>
</tr>
<tr>
<td>I:M &gt; 0.3</td>
<td>7 (58%)</td>
<td>9 (13%)</td>
<td>YES</td>
</tr>
<tr>
<td>Degenerative changes</td>
<td>12 (100%)</td>
<td>44 (65%)</td>
<td>YES</td>
</tr>
<tr>
<td>Platelet &lt; 1,50,000/cu.mm</td>
<td>0 (0%)</td>
<td>6 (9%)</td>
<td>NO</td>
</tr>
<tr>
<td>CRP -reactive</td>
<td>11 (92%)</td>
<td>35 (51%)</td>
<td>YES</td>
</tr>
<tr>
<td>H.S.S &gt; 3</td>
<td>9 (75%)</td>
<td>20 (29%)</td>
<td>YES</td>
</tr>
</tbody>
</table>

Table 2: Statistical analysis of hematologic parameters.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abnormal Leukocyte Count</td>
<td>66%</td>
<td>83%</td>
<td>42%</td>
<td>93%</td>
</tr>
<tr>
<td>ANC</td>
<td>66%</td>
<td>94%</td>
<td>66%</td>
<td>94%</td>
</tr>
<tr>
<td>INC</td>
<td>75%</td>
<td>83%</td>
<td>45%</td>
<td>95%</td>
</tr>
<tr>
<td>I:T</td>
<td>40%</td>
<td>94%</td>
<td>75%</td>
<td>80%</td>
</tr>
<tr>
<td>I:M</td>
<td>55%</td>
<td>86%</td>
<td>46%</td>
<td>92%</td>
</tr>
<tr>
<td>Degenerative change</td>
<td>100%</td>
<td>35%</td>
<td>21%</td>
<td>100%</td>
</tr>
<tr>
<td>Thrombo-cytopenia</td>
<td>0%</td>
<td>91%</td>
<td>0%</td>
<td>84%</td>
</tr>
<tr>
<td>CRP</td>
<td>91%</td>
<td>48%</td>
<td>23%</td>
<td>97%</td>
</tr>
<tr>
<td>H.S.S &gt; 3</td>
<td>41%</td>
<td>92%</td>
<td>50%</td>
<td>90%</td>
</tr>
</tbody>
</table>

Table 3: Comparing degenerative changes in present study with other studies.

<table>
<thead>
<tr>
<th>Author</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
</tr>
</thead>
<tbody>
<tr>
<td>Narasimha A (2011)</td>
<td>68.4%</td>
<td>66.6%</td>
<td>66.6%</td>
<td>40%</td>
</tr>
<tr>
<td>Makkar M (2013)</td>
<td>78.1%</td>
<td>94.4%</td>
<td>92.5%</td>
<td>82.9%</td>
</tr>
<tr>
<td>Supreetha (2015)</td>
<td>53%</td>
<td>89%</td>
<td>68%</td>
<td>82%</td>
</tr>
<tr>
<td>Present</td>
<td>100%</td>
<td>35%</td>
<td>21%</td>
<td>100%</td>
</tr>
</tbody>
</table>

Fig. 1: Peripheral blood film showing increased number of immature neutrophils (Leishman stain; X1000).

Fig. 2: Peripheral blood film showing Cytoplasmic vacuolation in neutrophil (Leishman stain; X1000).
screening test is necessary for the detection and treatment of neonatal sepsis. Blood culture which is considered the gold standard has very low sensitivity due to pre- and post-analytic factors and is also not available within the therapeutic window. Our study showed a culture positivity of 12% which is similar to other studies. The culture-positive cases were correlated with various hematologic parameters, CRP reactivity and HSS of Rodwell et al to assess the sensitivity, specificity PPV, NPV and their utility in the early diagnosis of sepsis.

Baseline characteristics: Male gender and low birth weight were more frequently associated with sepsis, similar to studies of Buch et al and Varsha et al. This is attributed to factors regulating synthesis of IgM which is situated on the X-chromosome and the low maternal IgM seen in low birth weight babies as proposed by Buch et al. Age of onset of sepsis- Among culture-positive cases only 42% of cases were in the early neonatal period (within 72 hours). This correlates with the study by Nandy et al (48 hrs. or 72 hrs.). Late onset sepsis occurs due to low birth weight, prematurity, admission in intensive care unit, mechanical ventilation, invasive procedures and administration of parenteral fluids.

Pre-term infants were underrepresented in our study (12%) 33% of whom were culture positive comparable to the studies done by Sriram et al and Supreetha. Hematological parameters- Among the hematologic parameters Immature Neutrophil Count was the best performing index with high sensitivity, specificity and NPV, similar to studies by Makkar et al and Khair et al. Degenerative changes and CRP were highly sensitive but lacked specificity as they were seen in culture negative cases too. I:T >0.2 and IM >0.3 were highly specific (94% and 86% respectively) with a high NPV (95% and 80%) similar to studies of Narasimha et al Nandy et al and Makkar et al respectively. Presence of degenerative changes in PMNs showed stress induced leucopoiesis. Degenerative changes in neutrophils exhibited high sensitivity but lacked specificity for sepsis similar to that of Supreetha et al. While many studies have seen a positive association between sepsis and thrombocytopenia none of the culture positive cases in our study indicated a low platelet count. According to Ghosh et al platelet count in isolation is not a reliable predictor of sepsis since thrombocytopenia is common in first week of life. Moreover, culture positive babies with thrombocytopenia do badly. But in our study since none of the culture positive babies showed thrombocytopenia and responded well to treatment. This indicates that thrombocytopenia has a possible prognostic value in culture positive cases.

Studies by Sriram et al have shown a high NPV with an ANC <2000. Present study showed 66%, 94%, 66%, 94% of sensitivity, specificity, PPV, NPV for an ANC < 2000. Out of the 12 culture positive cases 41% of cases had a score had score >5, 33% of cases had score 3 & 4. Thus, with a cut off score of > 2 HSS was highly specific (92%) with a NPV of (92%) indicating a HSS of < 2 can safely rule out sepsis and avoid unnecessary antibiotic treatment and resultant complications.

**Conclusion**

The results of the study indicate that HSS can be reliably and effectively used as a screening test to rule out sepsis.
in a neonate where clinical features often overlap with other conditions due to the non-specific symptoms. This avoids unnecessary administration of antibiotics to unaffected babies. HSS increases the diagnostic accuracy of complete blood count and standardizes its interpretation. Moreover, HSS is a cost effective, easily performed by basic qualification doctors in any rural areas with minimal resources. This will ensure immediate action for children with high index of suspicion of sepsis.

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Competing Interests
The authors declare that there is no conflict of interest.

References

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