HYPERGLYCEMIA: A BAD PROGNOSTIC FACTOR FOR NEONATE.

Muhammad Anwar¹, Muhammad Asghar Ali², Ali Hammad³

ABSTRACT… Objectives: The study was aimed to find out morbidity and mortality associated with hyperglycemia among neonates at our setting. Study Design: Observational Case-Control study. Setting: Department of Neonatology, Civil Hospital, Bahawalpur. Period: From 1st July 2019 to 31st December 2019. Material & Methods: The study included 194 neonates (97 each for cases and controls). Data like age (days), gender and weight were noted for all while presence of morbidities like IVH, NEC or infections were also recorded. Outcome among study participants of both groups was noted in terms of length of hospital stay and discharged/expired. Results: Out of a total of 194 study participants, majority, 114 (58.8%) were male, 101 (52.1%) had age between 1 to 7 days, and 84 (43.3%) had birth weight below 1.5 kilograms. Significantly more neonates had morbidity. Significantly more cases, 57 (58.8%) expired in comparison to 38 (39.2%) controls (p value 0.006). Length of hospital stay was also significantly more prolonged among cases. Conclusion: Neonates having hyperglycemia not only are at significantly increased risk of having morbidities and mortality.

Key words: Birth Weight, Hyperglycemia, Morbidity, Mortality, Neonates.

INTRODUCTION

Hyperglycemia is a commonly noticed metabolic abnormality among sick neonates.¹ General criteria for diagnosis of neonatal hyperglycemia is considered to be glucose levels above 125 mg/dl (6.9 mmol/L) or serum glucose concentration of more than or equal to 150 mg/dl (8.3 mmol/L).² Intervention is recommended with neonates presenting two consecutive readings (at least two hours apart) of blood glucose levels above 252 mg/dl (14 mmol/L) or even a single reading of more than 216 mg/dl (12 mmol/L) in the presence of glycosuria.³

The 1st week of life, more specifically, 1st 3 to 5 days are highly vulnerable and neonates are highly prone to neonatal hyperglycemia in this period.⁴ Hyperglycemia is known to have an indirect linkage with birth weight and gestational age.⁵ Newborns weighing less than 1000 gm are estimated to have 72% prevalence of hyperglycemia in comparison to less than 5% among newborns with birth weight of more than 2500 gm.⁶ Prematurity, intrauterine growth retardation (IUGR), stress, intravenous (IV) lipid infusion, birth asphyxia, high rates of infusion of glucose, nonexistence of enteral feeding and some drugs like theophylline, dopamine and steroids are some of the other known risk factors for hyperglycemia among neonates.⁷,⁸

Pathophysiology of hyperglycemia among neonates involves increased endogenous glucose production, hypo-insulinemia, insensitivity/resistance of insulin receptors, catecholamine and some anti-insulin hormones.¹ Hyperglycemia may present with or without symptoms. Common symptoms of hyperglycemia include dehydration, fever and inability to thrive. Many a times, hyperglycemia is the 1st presentation indicating diabetes mellitus among neonates.⁹

Local data lacks about aspects of neonatal hyperglycemia while international researchers have shown variable relationship between hyperglycemia and commonly found morbidities.
among these neonates like intra-ventricular hemorrhage (IVH), infections, necrotizing enterocolitis (NEC), increased length of hospital stay.\textsuperscript{10,11} The current study was aimed to find out morbidity and mortality associated with hyperglycemia among neonates at our setting. The results of this study will help us designing tools and techniques to handle commonly associated aspects of hyperglycemia in neonates presenting at our institution.

MATERIAL & METHODS
Department of Neonatology, Civil Hospital, Bahawalpur was the venue for this observational case-control study from 1\textsuperscript{st} July 2019 to 31\textsuperscript{st} December 2019. Approval from Institutional Ethical Committee was sought for this research. Informed consent was taken from parents/guardians of all the study participants.

For cases having hyperglycemia the sample size was calculated according to the formula: 
\[ n = \frac{z^2 \times p \times (1 - p)}{e^2} \]

Where: 
- \( z = 1.96 \) for a confidence level (a) of 95%,
- \( p = 50\% \),
- \( e = \) margin of error as 10%.

Sample size for cases having hyperglycemia turned out to be 97. Another 97 neonates who were euglycemic matched for age, weight, gestational age and clinical status served as controls. So, the study included 194 neonates (97 each for both groups). Hyperglycemia was labeled as glucose levels above 125 mg/dl (6.9 mmol/L) or serum glucose concentration of more than or equal to 150 mg/dl (8.3 mmol/L). Neonates having any kind of congenital malformations or whose parents/guardians did not give permission for inclusion in this study were excluded from his study.

Data like age (days), gender and weight were noted for all while presence of morbidities like IVH, NEC or infections were also recorded. Outcome among study participants of both groups was noted in terms of length of hospital stay and discharged/expired.

All the study information was noted on a pre-designed proforma. SPSS version 21.0 was used for data handling and analysis. Chi square test was used to note association of study variables with study groups and p value less than or equal to 0.05 was taken as statistically significant.

RESULTS
Out of a total of 194 study participants, 114 (58.8\%) were male and 80 (41.2\%) female. Overall, majority of the neonates, 101 (52.1\%) had age between 1 to 7 days while 84 (43.3\%) had birth weight below 1.5 kilograms. There was no statistical difference between gender, age and birth weight between cases and controls (p value > 0.05).

In terms of morbidity, significantly more neonates had morbidity among cases as compared to controls (p value < 0.001). Among cases, 48 (49.5\%) had IVH in comparison to 23 (23.7\%) in controls (p value < 0.001) representing OR of 3.152 (95\% CI: 1.705-5.826). Among cases, NEC was seen in 72 (74.2\%) as compared to 20 (20.6\%) among controls (p value < 0.001) representing OR of 11.088 (95\% CI: 5.674-21.670). Regarding infections, 55 (56.7\%) neonates had infections in comparison to 24 (24.7\%) controls (p value < 0.001) representing OR of 3.983 (95\% CI: 2.161-7.342).

Significantly more cases, 57 (58.8\%) expired in comparison to 38 (39.2\%) controls (p value 0.006). Length of hospital stay was also significantly more prolonged among cases as compared to controls (p value < 0.001).

DISCUSSION
Mechanisms like iatrogenic causes, insufficiency to suppress hepatic glucose production and insulin resistance/intolerance.\textsuperscript{1} Present was the 1\textsuperscript{st} of its kind at our institution to find out impact of hyperglycemia in terms of morbidity and mortality among neonates. In the present study, male predominance was noted among study participants. Around the world, it is a common finding that more male are brought to healthcare facilities to find medical care.
Sabzehei MK\textsuperscript{6} et al and Alexandrou G et al\textsuperscript{9} also noted more males to have hyperglycemia in their researches. Likewise, a local study from Lahore\textsuperscript{12} noted male to female ratio of 1.8 times which is close to what we noted in the present study.

Rozance PJ and Hay WW\textsuperscript{4} concluded that neonates with birth weight of less than or equal to 1 kg have 18 times more chance of having hyperglycemia in comparison to those who are between 1 to 2 kg. In the present study, 44.3\% of the neonates with hyperglycemia had birth weight less than 1.5 kg. Being low weight given enhancement to inheriting complex mechanisms like increased endogenous glucose production, hypoinsulinemia, insulin receptor insensitivity or resistance.\textsuperscript{13,14} Very similar to what we noted, Gul R et al\textsuperscript{12} found 48\% of the cases having hyperglycemia to be having birth weight of less than or equal to 1.5 kg. Yoo HS et al\textsuperscript{15} while observing extremely low birth weight newborns noted 85\% of the cases to have hyperglycemia. Beardsall K et al\textsuperscript{16} also observed 80\% of their very low birth weight cases to have hyperglycemia.

In the present study, 74.2\% of the neonates were reported within 1\textsuperscript{st} week of life. Neonates are more prone to have hyperglycemia during 1\textsuperscript{st} of life because of stress aiding to alteration in glucose metabolism resulting in hyperglycemia.\textsuperscript{1,4} Many researchers from around the world have recorded that 60\% to 85\% of the neonates with hyperglycemia are reported within 1\textsuperscript{st} week of life\textsuperscript{2,5} which is exactly what we noted in the present study. Another local study from Lahore\textsuperscript{12} noted 74\% of their neonates having hyperglycemia to be in 1\textsuperscript{st} week of life.

Presence of NEC among cases having hyperglycemia have been reported with varying results. We noted 74.2\% of the neonates having

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Cases (n=97)</th>
<th>Controls (n-97)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>58</td>
<td>56</td>
<td>0.9592</td>
</tr>
<tr>
<td>Female</td>
<td>41</td>
<td>39</td>
<td></td>
</tr>
<tr>
<td>Age (days)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;1</td>
<td>21 (21.6%)</td>
<td>17 (17.5%)</td>
<td>0.825</td>
</tr>
<tr>
<td>1-7</td>
<td>51 (52.6%)</td>
<td>50 (51.5%)</td>
<td></td>
</tr>
<tr>
<td>8-14</td>
<td>11 (11.3%)</td>
<td>16 (16.5%)</td>
<td></td>
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<tr>
<td>15-21</td>
<td>9 (9.3%)</td>
<td>8 (8.2%)</td>
<td></td>
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<tr>
<td>22-28</td>
<td>5 (5.2%)</td>
<td>6 (6.2%)</td>
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<tr>
<td>Birth Weight (kg)</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>&lt;1.5</td>
<td>43 (44.3%)</td>
<td>41 (42.3%)</td>
<td>0.947</td>
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<tr>
<td>1.6 - 2.5</td>
<td>32 (33.0%)</td>
<td>34 (35.1%)</td>
<td></td>
</tr>
<tr>
<td>&gt; 2.5</td>
<td>22 (22.7%)</td>
<td>22 (22.7%)</td>
<td></td>
</tr>
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</table>

Table-I. Characteristics of Study Participants (n=194).

<table>
<thead>
<tr>
<th>Morbidity</th>
<th>Cases (n=97)</th>
<th>Controls (n=97)</th>
<th>P Value</th>
<th>Odds Ratio</th>
<th>OR at 95% Confidence Interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intra-Ventricular Hemorrhage</td>
<td>48 (49.5%)</td>
<td>23 (23.7%)</td>
<td>&lt;0.001</td>
<td>3.152</td>
<td>1.705 - 5.826</td>
</tr>
<tr>
<td>Necrotizing Enterocolitis</td>
<td>72 (74.2%)</td>
<td>20 (20.6%)</td>
<td>&lt;0.001</td>
<td>11.088</td>
<td>5.674 - 21.670</td>
</tr>
<tr>
<td>Infections</td>
<td>55 (56.7%)</td>
<td>24 (24.7%)</td>
<td>&lt;0.001</td>
<td>3.983</td>
<td>2.161 - 7.342</td>
</tr>
</tbody>
</table>

Table-II. Morbidity comparisons between cases and controls (n=194).

Figure-1. Outcome comparison between cases and controls.
Hyperglycemia had NEC. Very similar to our findings, Gul R and Colleagues\textsuperscript{12} found 70.8\% of their neonates having hyperglycemia to have NEC. Mohamed S et al\textsuperscript{17} noted prevalence of NEC to be 4.1\% among neonates having hyperglycemia while Lugt NM et al\textsuperscript{18} and Sabzehei MK et al\textsuperscript{6} noted this to 2\% and 33\% respectively. The reason for this low prevalence in other studies could be because those researchers only included cases that met stage II and III of Modified Bell’s criteria while we included stage 1 cases along with those as well. Our results were well in accordance to local findings where similar diagnostic criteria for NEC was employed.\textsuperscript{12}

Hyperglycemia is known to impact white blood cells so this could be an important reason why increased chances of having infections lie with neonates having hyperglycemia.\textsuperscript{12} Bekhof J et al\textsuperscript{19} found infections to be present among 47.2\% of their neonates having hyperglycemia which is well in accordance to our results where we noted 56.7\% of our neonates having hyperglycemia to accompany infections. Other local researchers\textsuperscript{12} have noted 50\% of their neonates to have infections which are very close to present findings. Contrary to our results, studies done from other parts of the world have seen infections to be present among 29-36.4\% of their neonates with hyperglycemia.\textsuperscript{6,16} The reason for those authors to find infections less commonly affecting neonates with hyperglycemia could be because they only labeled infections on the basis of blood culture reports while we used clinical and laboratory investigations for this purpose.

Hyperglycemia is thought to cause hyperosmolarity which could be the reason for brain cell dehydration, capillary dilatation as well as cerebral bleeding.\textsuperscript{12} In the present work, we noted almost half of our cases to have IVH which is very close to what has been found around the globe where IVH has been noted to be ranging between 36-45.8\% of the neonates with hyperglycemia.\textsuperscript{11-16} USG cranial is a common way of diagnosing IVH among neonates\textsuperscript{12} and same method was applied in the present study.

Neonates having hyperglycemia are noted to have more chances of having complications following hyperglycemia that could prolong their length of hospital stay and adverse outcomes in comparison to other neonates.\textsuperscript{6} In the current study, we noted significantly more neonates to have expired. Very similar to our findings, a study done in Lahore\textsuperscript{12}, found the neonates with hyperglycemia to have significantly prolonged length of hospital stay (OR 0.821, 95\% CI:0.721-0.935, p=0.028), lower discharge rates (OR 0.739, 95\% CI:0.553-0.988, p=0.040) and higher mortality (OR 1.353, 95\% CI:1.012-1.809, p=0.040).

CONCLUSION
Neonates having hyperglycemia not only are at significantly increased risk of having morbidities like IVH, NEC and infections but they also have high risk of mortality.

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REFERENCES


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**AUTHORSHIP AND CONTRIBUTION DECLARATION**

<table>
<thead>
<tr>
<th>Sr. #</th>
<th>Author(s) Full Name</th>
<th>Contribution to the paper</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Muhammad Anwar</td>
<td>Manuscript writing, Data collection, Data entry, Analysis and proof reading.</td>
</tr>
<tr>
<td>2</td>
<td>Muhammad Asghar Ali</td>
<td>Data collection, data entry, Review research, Proof reading</td>
</tr>
<tr>
<td>3</td>
<td>Ali Hammad</td>
<td>Data collection, data entry, Review research proof reading</td>
</tr>
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</table>

**Author(s) Signature**

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[Signature]