Study of bio-clinical parameters in anemic and non-anemic individuals of selected age groups

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ABSTRACT

The word Anemia is a combination of two words, A standing for NO and Nemia standing for BLOOD. The decrease in the percentage of Hb, the decrease in the percentage of total count of RBC, and also the decrease in oxygen carrying capacity of blood is called Anemia. It is one of the most common of disorders seen more in the female population, than in the male. Anemias could be of both general and hereditary type. The present work aims at identifying different age group individuals (male and female), to carry a survey concerning their present and past case history with reference to their habits and style of living. A series of blood samples from selected diseased and undiseased population of either sex would be screened with reference to parameters like determination of Hemoglobin percentages, TC and DC counts, ESR, PCV, BT, CT, Blood Glucose, Cholesterol, Bilirubin, Urea and Creatinine levels will be assessed, correlated and finally a conclusion would be drawn so as to identify the major cause of the disease and what happens to the various biological parameters in the body.

Keywords: Anemia, Analysis, Correlation, RBC, WBC, Hemoglobin.

INTRODUCTION

Anemia AmE or Anemia (BrE) from the Greek (iā́čiśá) meaning “without blood”, refers to a deficiency of red blood cells (RBCs) and/or hemoglobin. This results in a reduced ability of blood to transfer oxygen to the tissues, causing hypoxia; since all human cells depend on oxygen for survival, varying degrees of anemia can have a wide range of clinical consequences. Hemoglobin (the oxygen-carrying protein in the red blood cells) has to be present to ensure adequate oxygenation of all body tissues and organs. Different clinicians approach anemia in different ways; two major approaches of classifying anemia’s include the “kinetic” approach which involves evaluating production, destruction and loss, and the “morphologic” approach which groups anemia by red blood cell size. The morphologic approach uses a quickly available and cheap lab test as its starting point (the MCV). Regardless of one’s philosophy about the classification of anemia, however, any methodical clinical evaluation should yield equally good results.

MATERIALS

Blood samples, Glucose, urea, Cholesterol, Bilirubin standards, RBC Diluting fluid, WBC diluting fluid. Working GOD/POD reagent for glucose estimation, N/10 HCl, 10% Sodium tungstate, Sulphuric acid, 2/3 N.Diacetyl monoxime (2 %) for Urea estimation, Diazo reagent and methanol for bilirubin estimation, Haemometer, Haemocytometer, Micropipettes, Elico SL-159 Spectrophotometer, Compound Microscopes, Table top centrifuge.
METHODS

Estimation of Hemoglobin

N/10 HCl was taken in the Hb tube up to lowest mark 20. The finger was pricked with sterile needle and measured amount of blood was placed into Hb tube. The Hb tube was placed in the computer or haemometer and drop by drop distilled water was added into it until the colour of solution in the tube coincides with the glass plate of the computer or haemometer. Based on color comparison in the tube and meter, the percentage of Hb was calculated.

Estimation of RBC

Blood was drawn in RBC pipette and filled up to the mark with diluting fluid. A cover slip was placed on the counting chamber and a small drop of diluted blood was allowed, to slip into the counting chamber by capillary action. The counting chamber was kept aside for three mins for the cells to settle. The cells were observed by placing them in neubaur chamber and the cells were counted from upper left corner. The total RBC was calculated using the formula:

\[
\text{Total RBC} = \frac{\text{No of cells} \times \text{Dilution Factor} \times \text{Depth Factor}}{\text{Area Count} \times \text{Total Ruled Area}}
\]

Estimation of WBC

Blood was drawn in WBC pipette and filled up to the mark with diluting fluid. A cover slip was placed on the counting chamber and a small drop of diluted blood was allowed, to slip into the counting chamber by capillary action. The counting chamber was kept aside for few mins for the cells to settle. The cells were observed by placing them in neubaur chamber and the cells were counted in four corners of the neubaur chamber. The total WBC was calculated using the formula:

\[
\text{Total WBC} = \frac{\text{No of cells} \times \text{Dilution Factor} \times \text{Depth Factor}}{\text{Area Count}}
\]

Estimation of blood glucose

This method was based on the principle of glucose oxidase and peroxidase method. Three clean and dry Test tubes are taken and 1 ml of working GOD & POD reagent was pipetted in all the three tubes. 0.01 ml of plasma was added in tube labeled test and 0.01 ml of working glucose STD in tube labeled standard the tubes are kept aside for 10 minutes at 37°C or 15 minutes at room temperature. The absorbance of the solution was measured against reagent blank at 505 nm.

Estimation of urea

This method was based on the principle of reaction of urea with diacetyl monoxime to produce a colored chromogen that is measured colorimetrically or spectrophotometrically. 0.1 ml of blood was washed into 3.3 ml of water and 0.3 ml of 10 per cent sodium tungstate was added along with 0.3 ml of 2/3 N sulphuric acid. The tubes are mixed well and centrifuged. 1 ml of supernatant fluid is taken and 1 ml of water, 0.4 ml diacetyl monoxime and 1.6 ml of the sulphuric acid-phosphoric acid mixture are added. Tubes are placed in a boiling water bath for thirty minutes, cooled, and read against reagent blank at 480 nm. At the same time the colour is developed from 1 ml of the two standards.

Estimation of Bilirubin

0.2 ml of serum is washed into 5.4 ml of water and mixed well. 2.8 ml of this was pipetted into a second tube to be used as blank. To the test 0.7 ml of diazo-reagent and to blank 0.7 ml of sulphanilic acid solution was added. Tubes are thoroughly mixed and kept aside for five minutes. The absorbance of the resulting solution is read at 540 nm or using a green filter. This gives the conjugated bilirubin. To obtain the total bilirubin 3.5 ml of methanol was added to each tube and read again after standing for five minutes. 0.2 ml of bilirubin standard to 3.5 ml of methanol was added, to this 0.7 ml of diazo-reagent was added and after mixing was read against a water blank after keeping the tubes aside for five minutes.

Estimation of Cholesterol

0.1 ml of serum was added to 10 ml of ferric chloride-acetic acid reagent in a glass-stoppered centrifuge tube. Tube was mixed well and allowed to stand for 10-15 minutes (or over night) for the proteins to flocculate. 5 ml of Centrifuge was transferred from the clear supernatant fluid to a glass-stoppered centrifuge tube. For standard 0.1 ml of physiological saline and 10 ml of the cholesterol standard were mixed for use and transferred 5 ml to a second-stoppered centrifuge tube. For blank
5ml of the ferric chloride-acetic acid reagent was taken in a third tube. Add 3 ml of sulphuric acid from a burette to all three tubes. The tubes are stoppered tightly and mixed by repeated inversion. The stopper is carefully loosened and allowed to stand for 20-30 minutes. Unknown and standard are read against reagent blank using a yellow filter or at 560 millimicrons.

RESULTS

After a systematic survey of population in different age groups is carried out and after pooling up data of the experiments and examination, the results with reference to various biological and clinical parameters were tabulated and are presented in table 1

Table 1: Clinical parameters in different age group individuals (Anemic and Non Anemic)

<table>
<thead>
<tr>
<th>Age group</th>
<th>Haemoglobin content gm(%)</th>
<th>Total RBC 3.5-5.0 (millic- oncells/mm³)</th>
<th>Total WBC 4000-11000 cells/mm³</th>
<th>Blood Glucose 80-140 mg/dl</th>
<th>Serum Urea 24-45 mg/dl</th>
<th>Serum Cholesterol 150-250 mg/dl</th>
<th>Serum Bilirubin 0.2-1.0 mg/dl</th>
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<tbody>
<tr>
<td>20-40</td>
<td>11</td>
<td>3.8</td>
<td>4200</td>
<td>107.812</td>
<td>36.6</td>
<td>184.615</td>
<td>0.97</td>
</tr>
<tr>
<td>40-60</td>
<td>10.5</td>
<td>3.6</td>
<td>8000</td>
<td>92.187</td>
<td>26.5</td>
<td>107.692</td>
<td>1.15</td>
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<tr>
<td>20-40</td>
<td>8</td>
<td>2.8</td>
<td>3400</td>
<td>93.75</td>
<td>27.61</td>
<td>115.2</td>
<td>0.8</td>
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<tr>
<td>60 and above</td>
<td>8</td>
<td>2.8</td>
<td>6200</td>
<td>90.625</td>
<td>25.81</td>
<td>84.615</td>
<td>0.77</td>
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<tr>
<td>20-40</td>
<td>10</td>
<td>3.5</td>
<td>8000</td>
<td>123.437</td>
<td>37.91</td>
<td>123.19</td>
<td>0.66</td>
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<tr>
<td>40-60</td>
<td>9.8</td>
<td>3.4</td>
<td>10400</td>
<td>140.625</td>
<td>45.1</td>
<td>369.23</td>
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<td>20-40</td>
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<td>4.2</td>
<td>7600</td>
<td>96.875</td>
<td>33.84</td>
<td>76.923</td>
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<td>4.6</td>
<td>2800</td>
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<td>49.01</td>
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<td>7.5</td>
<td>2.6</td>
<td>7400</td>
<td>90.625</td>
<td>25.81</td>
<td>92.06</td>
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<td>3.5</td>
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<td>37.17</td>
<td>76.92</td>
<td>0.65</td>
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<tr>
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<td>2.4</td>
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<td>87.5</td>
<td>25.51</td>
<td>123.19</td>
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<tr>
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<td>7200</td>
<td>121.875</td>
<td>36</td>
<td>107.69</td>
<td>0.84</td>
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<tr>
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<td>4600</td>
<td>103.125</td>
<td>35.91</td>
<td>107.69</td>
<td>0.95</td>
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</table>

DISCUSSION

After the results are tabulated and patterns of various parameters are studied and keeping in view the other health particulars of the subjects obtained from the survey forms it is found that subject who are anemic are usually found to possess very low hemoglobin values as is an evident feature of many anemic individuals. The variation in the count of RBC was evident but there was no subsequent change in the WBC content of the blood. The normal RBC count is usually between 3.5 to 5.0 million-cells/mm³; however in the present study the values only in normal cases were in the mentioned range and in anemic individuals they were sub-normal. In anemic individuals the levels of glucose were found to be slightly abnormal, whereas there was no significant change in the urea and bilirubin levels. The levels of cholesterol in few subjects who were suffering from anemia and cardiac diseases were found to be very high especially above 300 mg/dl.

CONCLUSION

It can be concluded that in female individuals and individuals suffering from respiratory disorders the concurrence of anemia is very high.
and the survey depicts the relationship between low Hemoglobin and RBC levels that are ultimately the root cause of the disease. However it was not very clear that, anemia could be associated with other diseases, however the survey revealed a few interesting facts, indicating a variation in glucose and cholesterol levels but normal urea and bilirubin levels, indicating that kidneys and liver do not undergo much damage and the carbohydrate and lipid metabolism undergoes considerable changes.

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REFERENCES

1. Practical Biochemistry by Jayaraman
2. Practical Biochemistry by David Plummer.
3. Textbook of Medical Lab Technology by Ramnik and Sood.
7. http://www.clinchem.org/cgi/content/abstract/19/2/253.
13. Textbook of Medical Biochemistry by Chaterjee and Shinde.