Changes of Selected Haematological Parameters in a Female Team during the 25th African Handball Winners’ Cup Played at Cotonou (Benin)

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The performance in team sports partly depends on the oxygen transport capacity that is also associated with the erythrocyte values, which should be regularly assessed throughout the sports season. This controlled observational study aims at measuring the haematologic modifications induced by the physical load associated with a series of matches in a Benin Division 1 female handball team. It was carried out at Cotonou during the 25th African Handball Winners’ Cup, with 22 female handball players of the Beninese champion team, divided into an experimental group (EG: n = 16) and a control one (CG: n = 6). The red blood cell count (RBC), the rates of Hb (Hb) and haematocrit (Ht), the Wintrobe’s constants and four iron parameters were assessed prior to and after a series of four matches played in six days. At the end of the competition, RBC increased significantly (p < 0.05), but the increases in Hb and Ht were non-significant (p > 0.05). Serum iron, serum transferrin and total iron binding capacity decreased significantly (p < 0.05). The results suggest an improvement in the players’ diet by highly protein enriched foods and erythropoiesis raw materials. They also suggest that the girls may adopt adequate hydric practices, in order to compensate for water losses during matches and training sessions.

Keywords: Erythrocytes; Iron; Competition; Handball; Tropical Climate

Introduction

Sports performance strongly depends on the oxygen transport capacity to supply exercising muscles. This capacity is associated with the erythrocyte values, which may thus be regularly assessed throughout the sports season (Fallon, 2004; Lesesve et al., 2000) to allow trainers and medical staff members to collect useful fitness and health related information on players. Many factors (environment, seasons, physiological state, nutritional profile, alcoholism, etc.) tend to modify haematological parameters, which sometimes leads to anaemia (Malcovati et al., 2003; Rietejen, 2002). In a sportsman practicing an intense long duration activity, water losses are accompanied by a decrease in iron store. In case this situation persists, it may lead to anaemia, as the sport considered requires important energy expenditure. More than one quarter of the male and the three quarters of the female long distance race specialists would be affected by iron deficiency (Clement & Sawchuck, 1984). The decrease in iron stores is indeed deeper in sporting women, because of the periodic menstruation which represents a source of blood loss. A decrease in Hb and haematocrit rates and red blood cell count was also recorded among runners and swimmers, in spite of a sufficient iron dietary intake (Pizza et al., 1997). The deficiencies may also appear in an environment like that of Benin where the prevalence of malnutrition and enteric parasites is high and malaria endemic (INSAE/Benin, 2012; Ministère de la Santé Publique du Bénin, 2009; Kindé-Gasard et al., 2000). On the first hand, the haemolysis and enteric bleedings that these last affections are likely to cause may induce a decrease in the Hb rate (Dillon, 2000). On the other hand, the higher an individual’s Hb rate, the stronger his/her oxygen transportation capacities (Gledhill et al., 1999). Many studies reported the effects of a training and competition period on the erythrocyte values and the iron status in long distance runners, swimmers, cyclists and soccer players (Santhiago et al., 2009; Silva et al., 2008; Zapico et al., 2007). In most of these studies, RBC, Hb and serum transferrin increase at the end of a high-level intensity exercise (Ashenden et al., 1999; Cordova & Escanero, 1992). The impact of the physical load associated with a tournament in team sports like handball on the haematological parameters is not well known, mainly in an environment like that of Benin, where it is likely to find a high prevalence of anaemia. When they exist, studies concern more male sportmen (Santhiago et al., 2009; Silva et

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Faecal samples were collected in individual test tubes to seek the presence of enteric worms by a coprology test. Height (near to a cm) and weight (near to 100 g) were then measured, using respectively a stadiometer and a digital bio-impedancemeter BG 22 (Beurer, Germany). A multifunction station (Meteo-Star) installed near the playing ground made it possible to record the ambient temperature and the relative humidity during the matches and training sessions. Resting heart rate (HRr) was taken by the Physician of the team on an electronic monitor UA-767 plus 30 (AND A&D Medical, Japan) in the players, seated for 10 min at least in a quiet room. A blood sample of 5 mL was drawn out after a 12 hour-fasting by venepuncture at the fold of the left elbow, to assess erythrocyte values and evaluate their malarial and iron status.

The erythrocyte values (red blood cell count or RBC, Ha and haematocrit rates, Wintrobe’ constants) were assessed using an automat counter M-Series (Medonic, Sweden). The iron parameters i.e. serum iron (Fe), serum transferrin (TR), ferritin (FERI), the rate of transferrin saturation (RTS) and the total iron binding capacity (TBC), protidemia (PTD) and plasma albumin (ALB) were assessed by the enzymatic and colorimetric method, using a spectrophotometer RT-9200 (Rayto, Germany). The same operations were repeated under the same conditions 36 hours after the last match (Figure 1).

During the matches and the two training sessions, the energy expenditure was evaluated with pedometers F700 (Geonaute, China). The meals consumed were weighed and the samples analysed in bomb calorimeters to determine the daily energy, vitamin C and iron consumption in each player. The water drunk was measured using individual bottles of 1.5 L given to each player at the beginning of each match and training session.

Statistical Analysis

The data were processed with the software Statistica of Stat Soft Inc. (version 5.5). The normality of the distribution of the variables was checked using the test of Kolmogorov Smirnov. The mean value (M) and the standard error of the mean (SEM) were calculated for each variable. The distribution of the variables not being normal, the Wilcoxon test was used for comparing measures within the same group, and the Mann-Whitney U test, for inter-group comparisons. Otherwise, the parametric tests were used. The level of significance of the statistical tests was settled at $p < 0.05$.

Results

Biometric Characteristics and the Results of the Matches

There was no significant difference between the EG and CG
groups ($p > 0.05$), so far as the biometric characteristics were concerned (Table 1). The ASO Baobab team was eliminated after having lost their three matches of the preliminary round against Heritage of Congo (23 versus 25 goals), TKC of Cameroon (24 versus 30 goals), and Electro of Angola (15 versus 29 goals), as well as their last match against Rombo of Côte d’Ivoire (30 versus 31 goals).

**General Data of the Matches, Dietary Patterns, Oestrian and Parasitic Status**

As it is shown in Table 2, the EG and CG groups were different with regard to the data collected during the matches and/or training sessions, i.e. the mean distance covered, the mean energy expenditure and the average quantities of water consumption were all higher in EG than in CG ($p < 0.05$). No significant difference ($p > 0.05$) was observed between the two groups regarding the daily energy, iron intake and C vitamin intake (Table 3).

At the beginning of the competition, 14 players of EG were in the follicular phase of their menstrual cycle versus two in the luteal phase. In CG, they were four in the follicular phase versus two in the luteal phase. Protidemia was reduced by $8.7 \pm 8.5 \text{ g/L}$ versus $78.8 \pm 5.2 \text{ g/L}$, $p = 0.001$) in EG and by $2.0 \pm 5.7 \text{ g/L}$ versus $79.0 \pm 2.9 \text{ g/L}$, $p = 0.500$) in CG. Plasma albumin was reduced by $4.8 \text{ g/L}$ versus $87.6 \pm 7.4 \text{ g/L}$, $p = 0.001$) in EG against $1.1 \text{ g/L}$ versus $81.0 \pm 2.9 \text{ g/L}$, $p = 0.396$). In EG, three positive cases of enteric parasites and five of infestation by *plasmodium falciparum* were found versus respectively zero and four cases of enteric parasites, then of infestation by *plasmodium* in CG.

**Table 1.** Biometric characteristics of the studied female handball players, prior to and after competition.

<table>
<thead>
<tr>
<th></th>
<th>EG (n = 16)</th>
<th>CG (n = 6)</th>
<th>$\Delta$ (EG)</th>
<th>Prior to competition</th>
<th>After competition</th>
<th>$\Delta$ (CG)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>23.81 ± 4.02</td>
<td>-</td>
<td>22.83 ± 3.26</td>
<td>-</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Height (cm)</td>
<td>167.31 ± 4.93</td>
<td>-</td>
<td>169.67 ± 4.03</td>
<td>-</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>60.56 ± 4.60</td>
<td>60.76 ± 5.12</td>
<td>74.08 ± 8.73</td>
<td>74.28 ± 7.56</td>
<td>0.20 ± 1.68</td>
<td></td>
</tr>
<tr>
<td>BMI (kg·m$^{-2}$)</td>
<td>21.59 ± 1.50</td>
<td>21.87 ± 1.51</td>
<td>25.82 ± 3.35</td>
<td>25.90 ± 3.09</td>
<td>0.8 ± 0.58</td>
<td></td>
</tr>
<tr>
<td>Resting HR (bpm)</td>
<td>54 ± 1</td>
<td>52 ± 2</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>-2 ± 1</td>
<td></td>
</tr>
</tbody>
</table>

Note: n: sample size; EG: experimental group; CG: control group; M: mean value; SEM: standard error of the mean; BMI: body mass index; Resting HR: resting heart rate; bpm: beats per minute. There is no significant difference between EG and CG groups for these values.

**Table 2.** General data collected during the matches and the training sessions.

<table>
<thead>
<tr>
<th></th>
<th>EG (n = 16)</th>
<th>CG (n = 6)</th>
<th>$\Delta$ (EG)</th>
<th>Prior to competition</th>
<th>After competition</th>
<th>$\Delta$ (CG)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean distance covered during the matches and the training sessions (m)</td>
<td>3062.7 ± 746.3</td>
<td>1763.0 ± 311.4</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>55 ± 0</td>
<td>-2 ± 1</td>
</tr>
<tr>
<td>Mean energy expenditure during the matches and the training sessions (kcal)</td>
<td>439.17 ± 142.81</td>
<td>174.78 ± 46.96</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>55 ± 0</td>
<td>-2 ± 1</td>
</tr>
<tr>
<td>Amount of water drunk during the matches and the training sessions (L)</td>
<td>1.38 ± 0.64</td>
<td>0.76 ± 0.38</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>55 ± 0</td>
<td>-2 ± 1</td>
</tr>
</tbody>
</table>

Note: The data are reported by player; n: sample size; M: mean value; SEM: standard error of the mean; EG: experimental group; CG: control group; $\Delta$: difference with the experimental group significant at $p < 0.05$.

Though opposite in both EG and CG (respectively $-0.8\% \pm 3.3\%$ versus $+0.5\% \pm 3.0\%$), the variation in plasma volume does not differ significantly between them ($p = 0.396$). In EG, three positive cases of enteric parasites and five of infestation by *plasmodium falciparum* were found versus respectively zero and four cases of enteric parasites, then of infestation by *plasmodium* in CG.

**Table 3.** Dietary intake data of the female handball players surveyed during the competition period.

<table>
<thead>
<tr>
<th></th>
<th>EG (n = 16)</th>
<th>CG (n = 6)</th>
<th>$\Delta$ (EG)</th>
<th>Prior to competition</th>
<th>After competition</th>
<th>$\Delta$ (CG)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Daily energy supply (kcal)</td>
<td>2185.95 ± 126.92</td>
<td>2176.55 ± 189.09</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>55 ± 0</td>
<td>-2 ± 1</td>
</tr>
<tr>
<td>Dietary iron (mg/day)</td>
<td>65.89 ± 1.64</td>
<td>65.99 ± 1.21</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>55 ± 0</td>
<td>-2 ± 1</td>
</tr>
<tr>
<td>Dietary C vitamin (mg/day)</td>
<td>42.98 ± 1.90</td>
<td>43.60 ± 1.92</td>
<td>-1 ± 1</td>
<td>54 ± 1</td>
<td>55 ± 0</td>
<td>-2 ± 1</td>
</tr>
</tbody>
</table>

Note: n: sample size; EG: experimental group; CG: control group; M: mean value; SEM: standard error of the mean.
Table 4.
Erythrocyte values in the studied female handball players, prior to and after the competition.

<table>
<thead>
<tr>
<th></th>
<th>EG (n = 16)</th>
<th></th>
<th></th>
<th></th>
<th>CG (n = 6)</th>
<th></th>
<th></th>
<th></th>
<th>RV</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Prior to competition</td>
<td>M ± SEM</td>
<td>After competition</td>
<td>M ± SEM</td>
<td>∆ (EG)</td>
<td>M ± SEM</td>
<td>Prior to competition</td>
<td>M ± SEM</td>
<td>After competition</td>
</tr>
<tr>
<td>RBC (10¹²·L⁻¹)</td>
<td>4.79 ± 0.44</td>
<td>4.85 ± 0.44</td>
<td>0.06 ± 0.12</td>
<td>4.53 ± 0.43</td>
<td>4.65 ± 0.44</td>
<td>0.12 ± 0.44</td>
<td>4.17</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hb (g·dL⁻¹)</td>
<td>13.00 ± 0.88</td>
<td>1.03 ± 1.01</td>
<td>0.03 ± 0.28</td>
<td>12.35 ± 1.15</td>
<td>12.30 ± 1.29</td>
<td>−0.05 ± 0.35</td>
<td>11.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ht (%)</td>
<td>40.44 ± 3.18</td>
<td>40.48 ± 3.18</td>
<td>0.44 ± 1.03</td>
<td>38.67 ± 3.07</td>
<td>38.67 ± 3.26</td>
<td>0.0 ± 0.63</td>
<td>37</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MCV (fl)</td>
<td>83.69 ± 2.77</td>
<td>83.25 ± 2.35</td>
<td>−0.44 ± 6.71</td>
<td>85.67 ± 6.50</td>
<td>82.67 ± 2.73</td>
<td>−3.00 ± 7.34</td>
<td>88</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MCHC (g·dL⁻¹)</td>
<td>32.44 ± 1.09</td>
<td>32.06 ± 0.92</td>
<td>−0.38 ± 0.80</td>
<td>32.17 ± 1.72</td>
<td>32.00 ± 1.09</td>
<td>−0.17 ± 0.75</td>
<td>32</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MCH (pg)</td>
<td>27.00 ± 1.26</td>
<td>26.63 ± 1.02</td>
<td>−0.38 ± 0.80</td>
<td>27.33 ± 2.65</td>
<td>26.17 ± 0.75</td>
<td>−1.17 ± 2.48</td>
<td>29</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: n: sample size; EG: experimental group; CG: control group; RV: reference values for female adults, used in the National Laboratory in the Republic of Benin; M: mean value; SEM: standard error of the mean; ∆ (EG): difference between the data collected prior to and after competition in EG; ∆ (CG): difference between the data collected prior to and after competition in CG; RBC: red blood count; Ht: haematocrit; Hb: haemoglobin; MCV: mean corpuscular volume; MCH: mean corpuscular haemoglobin; MCHC: mean corpuscular haemoglobin concentration; *p < 0.05.

Figure 2.
Modifications of serum iron in female handball players of Benin during the competition period (n = 22). *Difference between measures taken prior to and after competition in the experimental group, significant at p < 0.05.

Figure 3.
Modifications of serum transferrin in female handball players of Benin during the competition period (n = 22). *Difference between measures taken prior to and after competition in the experimental group, significant at p < 0.05.

Figure 4.
Modifications of total iron binding capacity in female handball players of Benin during the competition period (n = 22). *Difference between measurements taken prior to and after competition in the experimental group, significant at p < 0.005.

Figure 5.
Modifications of serum ferritin in female handball players of Benin during the competition period (n = 22). Differences between measures taken prior to and after competition in both groups were all non-significant (p > 0.05).
The main concern at the origin of this study was to measure the impact of a group form competition on the haematological parameters whose monitoring is necessary in athletes during the different periods of the season (Désidéri-Vaillant et al., 2003). This monitoring is all the more indicated as the players practise in a risky environment, i.e. an environment marked by strong heat, malnutrition and a high prevalence of transmissible diseases. As a hole, the characteristics of their daily environment contribute to increasing the risk of anaemia in the players (Ben Rayana et al., 2002). Therefore, in case the dietary survey revealed deficiencies, it was likely to expect an observation of modifications in the haematological parameters at the end of the competition.

This study was carried out with the players of the female champion handball team of Benin which took part in a continental-level competition. The studied team appeared relatively champion handball team of Benin which took part in a continental-level competition. The studied team appeared relatively

The Erythrocyte Values

Compared with the literature data (Aboussaleh et al., 2004; Moulessehoul et al., 2004), the mean erythrocyte data collected at the beginning of the competition in our female handball players are all within normal values. These results confirm those reported in French athletes practising various sports and whose biochemical and biological parameters were all within the normal limits (Désidéri-Vaillant et al., 2003). Similar results were previously recorded in female hockey and football players (Douglas, 1989).

The erythrocyte values do not differ considering the two groups of female handball players involved in the study sample. They were all trained players. Many studies revealed differences between trained athletes in various sports and untrained subjects (Boyadjiev & Taralov, 2000; Biancotti, 1992). The stability of the erythrocyte values observed in the study samples differs from data reported in Division 1 soccer players after five weeks of competition (Gouthon et al., 2011). Indeed, Gouthon and his contributors recorded an increase of RBC and Hb between test and retest in their study sample.

At the end of the competition, the erythrocyte values did not vary and remained within the reference ranges for female subjects, except for the RBC, which increased to a significant degree in the experimental group. Under these conditions, the lack of variation of the erythrocyte values in our series is probably due to the relatively short duration of the study, since repeated exercises at high intensity, are known to stimulate erythropoiesis (Szygula, 1990). The relatively low-level practice of the studied players could also explain the stability recorded in the erythrocyte values. A priori, one can also assert that the players’ dietary intake was deficient with regard to the nutrients constituting the raw materials for haematopoiesis, i.e. dietary iron, C and B12 vitamins. Since the erythrocyte parameters did not decrease and that C vitamin and iron intake are below the reference ranges and recommended values for athletes, i.e. respectively 110 mg per day and 16 mg per day (Martin, 2001; Agence Française de Sécurité Sanitaire des Aliments, 2000), one can suppose that the exercise-stimulating effect on the haematopoiesis intervened to compensate for the light nutritional deficiency (Szygula, 1990; Schumacher et al., 2002; Schoebersberger et al., 1990).

Few subjects (3 out of 22) were infested by the malaria parasite (Plasmodium falciparum) and enteric worms were present in only 40% of the girls. These data contrast with that of the overall Beninese population within which the prevalence of the enteric parasites is higher (INSAE, 2012). They are certainly the expression of the good quality of the counsels and the medical follow-up from which these female players of ASO Baobab profited and that make them a privileged group. In spite of the fact that an important number of the EG players were menstruating during the study period, the RBC increased. So, blood losses do not seem to have had a negative impact on blood volume in these players, but it would have been interesting to collect information relating to the duration of the menstruation. Physical exercise may indeed cause menstrual disorders in athletes, i.e. a reduction in the duration of the blood flow during the menses (Wilmore & Costill, 2006; Portal et al., 2003).

The reduction in plasma volume allows suspecting a lack of compensation of the hydric losses caused by match-related efforts, although the players of the experimental group drank an average of 1.38 L of water per match or training session. However, the low values of plasma albumin recorded before and after the competition, do not permit to retain this assumption. Indeed, only high values of this blood protein, i.e. those higher than 52 g/L (Janssens, 2009) are associated with water loss-related dehydration. It is why the weight gain (0.31% of the initial weight) at the end of the competition in EG players should be associated with an increased protein synthesis during recovery periods (Poortmans, 1984; Guezennec, 1989).

Iron Parameters

In this study, serum ferritin remained stable from the beginning to the end of the competition, thus testifying the fact that body iron stores were not depleted during this period. It was the same for the rate of transferrin saturation which constitutes one
of the iron bioavailability markers. TBC was on the other hand reduced, as well as RBC suggesting a decrease in the oxygen transportation capacity in the players of the experimental group.

These data suggest the following two observations.

The first one is about the dietary intake related to the raw materials for erythropoiesis in these female players. The data collected reveal weak iron and C vitamin intakes during the competition period. Under these conditions, a reduction of the erythropoiesis in the handball players could explain the decrease in TBC recorded at the end of the competition. Since physical exercise is regarded as a trigger of erythropoiesis (Szygula, 1990; Schobersberger et al., 1990; Schumacher et al., 2002), other factors may be taken into account to explain this result. A more important red cell destruction than red cell production which requires at least one week may be an explanatory hypothesis. Ultimately, the factor most likely to explain the reduction in TBC at the end of the competition remains the decrease in serum transferrin, which is one of the iron carrier proteins.

The second observation relates to the stability of iron stores in the players’ bodies, contemporary to the decrease in serum iron during this period. If there was a concomitant reduction of the bioavailable iron (RTS and TBC), (Haute Autorité de Santé, 2011) there would be the suspicion of an inflammatory process in the players as a result of the high intensity of the match-related physical loads. Since Hb has not decreased below the normal value (12 g/100 mL), such a hypothesis is given up.

Dietary deficit needs to be pronounced and chronic beyond a few days, to induce the depletion of iron stores in the body.

Though their average daily 2185 kCal intake near the minimum of 2200 kCal recommended for female athletes (Martin, 2001), these girls’ diet has to be improved quantitatively as well as qualitatively during competition periods.

Conclusion

This study belongs to the few ones addressing the modifications of the haematological parameters during a handball continental-level competition, played in a tropical environment. The four matches played in six days did not significantly modify the players’ erythrocyte values. Serum iron, TBC and serum transferrin were however reduced at the end of the matches of the preliminary round, without any case of anaemia. Thus, in the Beninese environment characterized by a high prevalence of transmissible diseases, it is possible to reduce the risk of anaemia in athletes, as long as the medical follow-up makes it possible to avoid parasites. The data of this study suggest associating good medical follow-up in these handball players, to an improvement of their diet with highly protein-enriched food and erythropoiesis raw materials such as iron, C and B12 vitamins.

The data also suggest that these girls adopt adequate hydration practices, in order to compensate for the effort-induced water losses during matches and training sessions.

Acknowledgements

The authors thank the President and Members of the ASO Baobab handball club of Lokossa (Republic of Benin) who agreed and helped for carrying out this research. We are also grateful to all the female players who accepted to take part in the study and to the Members of the Fitness Centre VITA FORM of Porto-Novo, for their technical assistance.

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