1Title: Assessment of Physical demands and Fluid balance in elite female handballplayers during a 6-day competitive tournament

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25Title: Assessment of Physical demands and Fluid balance in Elite Female Handball Players During a 6-Day Competitive Tournament

28Abstract

29Little data exists on drinking behaviour, sweat loss and exercise intensity across a competitive 30handball tournament in elite female athletes. Heart rate (HR), fluid balance and sweat electrolyte 31content were assessed on 17 international players across a 6-day tournament involving 5 games 32and 2 training sessions played indoors ($23 \pm 2^{\circ}$ C, $30 \pm 2\%$ relative humidity). Active play 33(effective) mean HR was 155 ± 14 bpm ($80 \pm 7.5\%$ HR_{max}) with the majority of time (64%) spent 34exercising at intensities >80% HR_{max}. Mean (*SD*) sweat rates during games was 1.02 ± 0.07 L·h⁻¹ 35and on 56% of occasions fluid intake matched or exceeded sweat loss. A significant relationship 36was observed between estimated sweat loss and fluid intake during exercise ($r^2 = 0.121$, P = 370.001). Mean sweat sodium concentration was 38 ± 10 mmol·L⁻¹, with significant associations 38observed between player sweat rates and time spent exercising at intensities >90% HR_{max} ($r^2 =$ 390.181, P = 0.001). Fluid and electrolyte loss appear to be work rate dependent in elite female 40handball players, whom appear well capable of replacing fluids lost within a tournament 41environment. Due to large between-athlete variations, a targeted approach may be warranted for 42certain players only.

43Introduction

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45Handball is a fast paced body-contact Olympic sport played by two competing teams of 7 players 46(including a goalkeeper) on an indoor court (40m x 20m) over two 30 minute periods. It is 47generally recognised that games are played at high-intensity due to the nature of rolling 48substitutions and recent rule changes on starting the game from the centre after a goal is scored. 49Despite its popularity, a paucity of data exits to describe the games physical demands. Recent 52interspersed by a short duration of very high-intensity anaerobic actions (Póvoas et al., 2012). In 53the latter study, players were observed to exercise at a mean intensity of $82 \pm 9.3\%$ of HR_{max}. To 54our knowledge, only one study describes the physical demands within the elite women's game 55with players observed to cover distances of $4002 \pm 551m$ and exercise at high mean relative 56workloads (79.4±6.4% VO₂-max) during match-play (Michalsik et al., 2013).

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58Handball is played indoors, and although most playing halls have temperature controlled 59environments, high-intensity exercise may result in thermal stress and subsequent disruption of 60body water and electrolyte balance. Team sport players have been shown to lose variable amounts 61of electrolytes in sweat (Kurdak et al., 2010; Maughan et al., 2007b) and the magnitude of solute 62loss appears dependant on both sweat rate and composition. In soccer, large individual variations 63in player sweat response are known to occur as a result of differences in environmental 64temperature and humidity, work-rates, state of acclimation and individual fitness (Maughan, 65Watson, Evans, Broad & Shirreffs, 2007). When exercising in the heat, a 1-3% loss in BM due to 66sweat losses has been reported to result in elevated HR, core temperature (Tc), rating of perceived 67exertion, and plasma osmolality (Montain & Coyle, 1992; Buono & Wall, 2000). More recently, it 68has been observed that muscle glycogenolysis significantly increases early in exercise with a BM 69 loss of < 2% and this appears related to a rise in core and muscle temperature (Logan-Sprenger et 70al., 2013). However, available data has shown that values in excess of 2% BM are needed before 71 aspects of team sport physical and cognitive performance begin to be negatively affected (Baker et 72al., 2007; McGregor et al., 1999). While many studies have investigated fluid balance in team 73sport athletes during practice or competition in the field (Maughan et al., 2005; Kurdak et al., 742010; Maughan et al., 2007) few, if any, have explored changes in these variables across a

75competitive tournament. In particular, there are no data available on women's handball. Olympic 76handball teams may play up to 5 games in 9 days during group stages of a major competition. 77Since prior hypohydration will amplify the effects of any fluid deficit incurred during exercise, 78added emphasis on fluid-electrolyte replacement may be required during such time periods to 79ensure adequate recovery between games. For this reason, understanding the pattern of fluid and 80electrolytes losses during a tournament may assist with the provision of targeted interventions in 81players with high sweat rates or where failure to replace lost fluids might raise concerns of an 82ergolytic effect.

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84The main aim of this investigation was to observe normal fluid balance behaviour and quantify 85electrolyte losses across a competitive tournament in a team preparing for the Olympic Games. 86Secondary aims were to describe the physical demands of elite women handball players and 87investigate whether or not fluid-electrolyte losses could be explained from exercise intensity and 88effort distribution.

89Methods

90*Participants.* Data were collected on an international team (n = 17) of female handball players ($26 \pm$ 915 yr, 1.72 ± 0.06 m, 70.7 ± 8.5 kg: mean \pm SD) during a tournament scenario over six days.

92*Experimental design*.

93The team participated in two training sessions and five competitive friendly games against top club 94opposition whom were normally playing at the highest standard in European handball competition 95(Figure 1). Data collection took place at two training venues seven months before the 2012 Olympic 96Games which the team under investigation subsequently participated. Games were played according 97to international handball federation rules and with the exception of game 4 (lasted 90 minutes due to 98a prior agreement between coaching teams), consisted of two periods of 30 minutes. The team 99under investigation recorded three wins and two losses during the investigation. Training sessions 100lasted 104 ± 2.1 minutes, consisting of warm-up, calisthenics, position-specific drills, small-sided 101games and technical drills. Environmental conditions during the games/training sessions are shown 102in Table 1 and all measurements were recorded with a portable thermohygrometer (Higbo, Oregon 103Scientific, Berkshire, UK).

104 Study approval was obtained from the national governing body involved and collection of data 105arose as part of the service provision supplied to the team in preparation for competition (Winter & 106Maughan, 2009). All athletes agreed to take part and signed consent was obtained. To ensure player 107confidentiality, data were anonymized before analysis.

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109*Testing procedures* Players wore shorts and short sleeve T-shirts throughout testing. On the 110morning of training or games, players provided a urine sample (first void) into a pre-labelled urine 111collection pot (60 ml, Greiner Bio-One Ltd, Stroud, UK). Urine refractive index (Pocket 112Osmocheck, Vitech Scientific Ltd, Sussex, UK) were subsequently determined to provide an index 113of pre-exercise hydration status (Shirreffs, 2003; Sparks and Close, 2012). Players were provided 114with dietary plans by the affiliated nutritionist in the lead up to training/games. As per norm, food 115was self-prepared in large player groups within nearby shared accommodation to the team home 116venue and this was not altered during the investigation. Neither dinner nor breakfast was recorded 117or standardized so as to avoid interference with player's normal dietary/fluid habits. Player 118menstrual state was not recorded during the 6-day tournament.

119Over a 15-min period (-45 to -30 min) before the start of each game/training, players were weighed 120(nude) to the nearest 20 g on a calibrated electronic scale (Marsden W/M, Oxford, UK). After entry 121to the playing court, players were fitted with a heart rate (HR) strap and individual HR data was 122subsequently collected wirelessly at one-second intervals (Polar Team System, Polar Electro,

123Kempele, Finland). Following team warm-down, players were re-weighed (towelled dry nude) on 124the same calibrated scale. Body mass changes were used as a proxy measure for sweat losses. While 125this represents a limitation of the study, previous work has indicated that this represents a realistic 126field measurement to estimate hypohydration (Maughan et al., 2007a). Players were asked to refrain 127from eating between weighing points but were free to drink fluids *ad libitum* and whenever breaks 128in play were permitted (1 minute time-out per-half is allowed by handball rules). Players were asked 129to consume drink products which they would normally consume during everyday practice and 130games. This included either tap water or tap water mixed with sugar free cordial for the current 131player group. Observed checks confirmed that no player consumed commercial drink products or 132electrolyte sachets during exercise. Therefore electrolyte intake during sessions/games was 133considered negligible.

134All bottles were individually numbered and players drank from bottles assigned to them by team 135staff. All players were left to behave as they wished with regard to fluid consumption. During the 136time period from weighing players before and after exercise, each player's fluid intake was 137measured to the nearest 0.001kg on a calibrated scale (Adam Equipment Ltd., Milton Keynes, UK). 138A volume of 1 mL ingested fluid was assumed to weigh 1 g. In the event of players wishing to 139urinate during the analysis period, they were accompanied by research staff to the toilet and 140provided with a pre-weighed urine collection bag with funnel (P-Bag, Medipost Ltd, Dorset, UK). 141Filled urine collection bags were subsequently weighed to aid in determination of fluid balance. All 142bottles/collection devices were weighed out of view of the players at all times.

143

144Sweat collection and analysis

145Before three of the five games, players were prepared for sweat sample collection using methods 146outlined previously (Maughan et al., 2007b; Maughan & Shirreffs, 2008). Briefly, players had 147absorbent patches (Tegaderm + pads, 3M, Loughborough, UK) placed on four body locations 148(chest, upper forearm, back and thigh). Before attaching patches to the skin, the locations were 149washed thoroughly with deionized water and dried with electrolyte-free gauze. At the end of each 150game, patches were removed with sterile forceps and each patch was subsequently placed into pre-151weighed screw-top containers (20 ml, Sterlin, Cambridge, UK) until analysis. No patch showed 152signs of undue saturation (e.g. dripping) across the period of investigation. Following collection, 153samples were stored at 4°C before analysis (within one week of collection).

154To determine electrolyte concentrations, each sweat patch and container were weighed to the 155nearest 4 decimal places before 1 mL of deionized water was added. After mixing, the sample was 156analysed for sodium and potassium concentration using flame photometry (Sherwood 410, 157Sherwood Scientific, Cambridge, UK) and for chloride concentration using the mercuric thiocynate 158method (Randox, Crumlin, UK). All samples were analysed in duplicate and intra-run coefficients 159of variations were <5%.

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161Heart rate data

162Player heart rate max data (HR_{max}) was obtained in the weeks leading up to experimentation using 163an incremental running test to exhaustion (Yo-Yo IE2 test) conducted by strength and conditioning 164staff. Confirmatory HR_{max} analysis was achieved by comparing these results to those achieved 165previously (-4 months) using a laboratory incremental test to exhaustion. Previous work has shown 166that no systematic differences occur between HR_{max} achieved using the Yo-Yo IE2 test and 167incremental treadmill test in intermittent sport (Bradley et al., 2011).

168Heart-rate readings were classified into 5 intensity zones, ranging from 50-59%, 60-69%, 70-79%, 16980-89% and 90-100% HR_{max} for assessment of physiological load in team handball (Póvoas et al.,

1702012). The percentage of time spent exercising in each zone was determined to provide an 171 indication of exercise intensity for each player and all HR values were analysed during the first and 172 second halves. Given that team handball rules allow for unlimited substitutions, it is unusual that 173 one athlete plays an entire game. Furthermore, a one-minute time-out period is allowed for each 174 team per-half. Therefore, HR during the games were analysed (a) as total HR (i.e. HR during the 175 total game time) to provide an index of global cardiovascular load and (b) HR during active play 176 time (effective HR) in order to classify game demands during time when a player was actually on 177 the playing court (Póvoas et al., 2012). The time period for the half-time break was excluded from 178 HR analysis.

179Calculations

180Sweat rate (SwtR: $L \cdot h^{-1}$) was estimated as net body mass loss (kg) during the training session/game 181plus the total fluid intake divided by the exercise time (min). Corrections for any individual player 182urine loss were also made.

 $183SwtR = \{ [pre-exercise mass (kg) - post-exercise mass (kg)] - urine volume (L) + fluids consumed (L) \} / time (h)].$

184Fluid intake $(L \cdot h^{-1})$ was calculated as total fluid intake during the recorded exercise session divided 185by the exercise time (per hour). Changes in BM before and after exercise were to assess acute 186changes in hydration status (% fluid deficit). Net balance of respiratory and metabolic contributions 187to BM loss was considered negligible as observed previously under similar environmental 188conditions for indoor team sport (Hamouti et al. 2010). Sweat electrolyte losses (in grams) were 189calculated from the sweat electrolyte concentration, the molecular weight of the electrolyte, and the 190total sweat loss of the individual (Maughan et al., 2007a). It was assumed that the mean sodium 191concentration (from 4 collected sites) represented mean whole-body sodium concentration although 192it is acknowledged that this may be an overestimate (Shirreffs & Maughan, 1997).

193Statistical analysis

194Data sets were initially tested for normality of distribution using the Shapiro–Wilk test. 195Comparisons between data were then made using paired t-tests and a one-way repeated measures 196analysis of variance (ANOVA). Differences in regional sweat electrolyte concentration were 197determined by 1-way ANOVA and with Tukey's test of honestly significant difference. 198Relationships between parameters were performed using a least-squares regression model, and the 199coefficient of determination (r^2) is reported (Maughan et al, 2005). All statistical analyses were 200performed using SPSS v. 20. Data are presented as mean \pm standard deviation (SD) with range of 201data given in parentheses. Statistical significance was set at P < 0.05.

202

203 204**Results**

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206*Heart rate*

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208Player HR_{max} achieved during incremental running test was 191 ± 7 bpm⁻¹. Peak HR over the 5 209games (182 ± 9 bpm⁻¹; 93 ± 5% of HR_{max}) was significantly higher than values recorded in training 210(169 ± 13 bpm⁻¹; 86 ± 5% of HR_{max}); P < 0.05. Player effective mean HR (155 ± 14 bpm⁻¹; 80 ± 8% 211of HR_{max}) was also significantly higher than values observed in training (129 ± 10 bpm⁻¹; 65 ± 5 % 212of HR_{max}) and total mean game HR (131 ± 14 bpm⁻¹) (P < 0.05). Average time spent by players in 213"active" play during games was 33 ± 15 minutes.

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215No between game-half differences were observed for either total or effective mean player HR 216(Figure 2). Furthermore, no significant differences were observed between game halves at different 217interval percentages of players' HR_{max} (Figure 3). The majority of effective game time (64%) was 218spent exercising at intensity >80% HR_{max} , with 23% of game time spent exercising >90% HR_{max} . No 219between game differences were observed for percentage of effective or total game time spent any of 220the designated HR zones.

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222223Morning hydration status224

225On the mornings of the games, 47% of players awoke with a urine refractive index \geq 700 226mOsmol/kg. No relationship was observed between previous days fluid intake during exercise and 227the following mornings urine refractive index (P = 0.116). Furthermore, no relationship was 228observed between morning urine refractive index and subsequent fluid intake during exercise, (r² = 2290.014; P = 0.307; Figure 4a). A significant relationship was observed between morning urine 230refractive index and percentage change in player BM during training/games (r² = 0.159; P = 0.001), 231Figure 4b. No between game differences in morning measures of hydration status were observed 232(Figure 6).

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234

235Fluid balance

236Across the games, mean team fluid intake was 1.05 ± 0.16 L. After correction for amount of fluid 237ingested, estimated mean team fluid loss was 1.08 ± 0.18 L. A significant relationship was observed 238between fluid intake during exercise and estimated fluid (sweat) loss (r² = 0.0121; P = 0.001; Figure 2395). Fluid intake (L) was significantly lower during training sessions in comparison to games 2, 3 240and 4 (P < 0.001) (Figure 6). Between game comparisons indicated that players consumed 241significantly more fluid during game 4 than games 2 and 5. Larger volumes of fluids were also 242consumed during game 3 in comparison to game 5 (P < 0.001).

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244Mean player sweat rates during games $(1.02 \pm 0.07 \text{ L}\cdot\text{h}^{-1})$ were significantly higher than those 245observed during training $(0.56 \pm 0.15 \text{ L}\cdot\text{h}^{-1})$ despite similar environmental temperatures (P < 0.05; 246Table 1). No between game differences were observed. A relationship was observed between 247individual player sweat rates (L·h⁻¹) and time spent exercising at intensities >90% HR_{max} (r² = 0.181; 248P = 0.001). During games, 51% of players had a sweat rate < 1 L·h⁻¹ and 47% had a sweat rate 249between 1 and 2 L·h⁻¹. Only on two occasions did player sweat rates exceed 2 L·h⁻¹ during games. 250In contrast, all players had sweat rates < 1 L·h⁻¹ during training. A large inter-individual variability 251was observed in sweating response between players (Figure 5).

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254On 56% of occasions, player fluid intake matched or exceeded fluid (sweat) loss. Sweat loss was 255significantly higher during game 4 than training sessions and game 2 (P < 0.05), game 3 and 5 (P < 2560.01). Across the five games, 43% of players lost body mass (BM) as a result of exercise with most 257(36%) of the group recording losses of <1% BM. In turn, 7% of the player group recorded a loss of 258>1% BM and no player was dehydrated >2% of BM. Conversely, 6% of the players gained more 259than 1% in BM across the games. On average, $0.0 \pm 0.1\%$ and $0.1 \pm 0.1\%$ net fluid deficits were 260recorded following games and training respectively.

261262263 *Electrolyte balance*264

265The mean (4 site) sweat sodium, chloride and potassium concentrations were $38 \pm 10 \text{ mmol}\cdot\text{L}^{-1}$, 27 266± 11 mmol·L⁻¹ and 5 ± 1 mmol·L⁻¹, respectively (Table 2). No significant effects for time (between 267games) were observed for sweat sodium, chloride or potassium content (P > 0.05). Mean sodium 268content of sweat in game 4 was significantly higher than game 5 (P < 0.01) while higher sweat 269chloride content was also observed in game 4 compared to game 2 and game 5 (P = 0.03). Sodium 270concentrations in the sweat patches of the 'back' torso were lower compared to all other sites (P < 2710.001). Higher 'arm' chloride concentration was observed in comparison to values recorded from 272players 'back' and 'thigh' (P < 0.001). In turn, chest chloride concentrations were higher than both 273'back' and 'thigh' samples (P < 0.001). Arm potassium concentrations were higher than back, chest 274and thigh concentrations while thigh concentrations were higher than back and chest (P < 0.001; 275Table 2). No association was observed between mean player sweat sodium concentration and sweat 276rate (r² = 0.001; P = 0.57).

278 A significant correlation was observed for total sodium loss $(1.03 \pm 0.43 \text{ g})$ and percentage of time 279spent by players exercising between 90-100% HR_{max} ($r^2 = 0.210$; P = 0.016).

280 281**Discussion** 282

283The aim of this investigation was to assess fluid balance behaviour and determine electrolyte losses 284across a competitive tournament in elite female handball players. Secondary aims were to assess 285physical demands and whether or not observed changes in the above variables could be explained 286from player exercise intensity data.

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288While a limited number of studies have documented intensity of exercise in elite men's handball, to 289date, only one study has been carried out in the elite women's game (Michalsik et al., 2013). 290Present data shows that female players exercise at a similar mean intensity (155 \pm 14 bpm⁻¹; 80 \pm 2918% of HR_{max}) to their elite male counterparts (Chelly et al., 2011; Póvoas et al., 2012) and that the 292majority of time (64%) is spent performing movements at high >80% HR_{max} exercise intensities. As 293expected, mean exercise HR was considerably higher during competition in comparison to those 294observed in training sessions. This was due to the fact that the coaches used the training sessions 295with a tactical focus and structured lower intensity than normal to ensure players were not fatigued 296during the tournament. However, team HR values did not decrease during the second-half of games 297as observed previously in the elite men's game (Póvoas et al., 2012). In the latter study, modest 298average relative body weight losses (0.9 \pm 0.3%) were recorded. This, combined with data from the 299current study (0.0 \pm 0.1% fluid deficit), suggests that factors other than dehydration may have be 300responsible for any observed drop off in exercise intensity in Handball. Although dehydration due 301to sweat loss is a known factor which can impair exercise performance (Coyle, 2004); other factors 302such as game rotation strategy, score differential, physiological or physical factors may all 303contribute to game fatigue. It should also be noted that HR data only provides an indication of 304cardiovascular demands and time-motion characteristics should be studied to better understand how 305fatigue affects movement patterns.

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307In the current study, player sweat rates $(1.02 \pm 0.07 \text{ L} \cdot \text{h}^{-1})$ were similar to the ones observed in 308 female basketball players (Brandenburg & Gaetz, 2012), indoor male handball $(1.1 \pm 0.3 \text{ L} \cdot \text{h}^{-1})$ and 309volleyball $(1.2 \pm 0.3 \text{ L.h}^{-1})$ players but considerably lower than indoor soccer $(1.8 \pm 0.7 \text{ L}^{-1})$ 310(Hamouti et al., 2010). However, a large inter-individual variability (range 0.26-2.1 L) was 311observed, as noted previously (Maughan et al., 2007b; Hamouti et al., 2010). This, taken together 312 with the large variability in fluid intake across the investigation (range 0.25-2.74 L) highlights the 313need for individualized assessment of fluid balance strategies in a team sport environment 314(Shirreffs, Sawka & Stone, 2006). With respect to the current athlete group, it was noted that 6% of 315 players gained more than 1% in body mass (BM) by the end of the games, suggesting that some 316 players might consume excessive fluids. Overall, 56% of players either maintained or increased 317their BM at the end of the game and this was particularly evident for goalkeepers and rolling 318substitutes, some of whom consistently consumed more fluids than were lost in sweat. This was 319most likely due to increased time available on the bench and ease of access to drinking bottles. 320During handball competition, BM has to be moved against gravity and theoretically an increase in 321BM from resultant fluid intake might have negative effects on performance in particular when 322 fatigue develops.

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324It has been reported that as little as a 2% loss in BM due to dehydration affects many physiological 325factors thought to indirectly affect performance. These include elevations in HR, core temperature 326(Tc), rating of perceived exertion (Montain & Coyle, 1992; Buono & Wall, 2000) and muscle 327glycogenolysis (Logan-Sprenger et al., 2012). In this study, no player was dehydrated >2% BM

328despite moderate to high sweat rates in some players. The most recent position stand from the 329American College of Sports Medicine (Sawka et al., 2007) has suggested that fluid intake during 330prolonged exercise should be sufficient to limit any BM loss to <2% of pre-exercise mass and that 331athletes should never drink so much that they gain BM during exercise. Therefore, the female 332handball players observed in this study appear to adequately replace fluids lost during exercise, and 333in some cases, education around the risks of overconsumption of fluids is warranted.

334

335Across the six-day tournament, 54% of players awoke with a urine refractive index of <700 336mOsmol/kg or a urine specific gravity <1.020, values previously suggestive of euhydration (Sawka 337et al. 2007). Research has reported that 91% of elite male athletes from a number of indoor team 338sports awoke in hypohydrated state before practice (Hamouti et al., 2010). Therefore, data is 339suggestive that fluid management following exercise appeared adequate in the current player group. 340Start time of training and games varied considerably between days and no difference in markers of 341hydration status was observed between sampling points. It is possible that given the elite nature of 342the current team and the tournament scenario, players were proactive in making sure to replace lost 343 fluids during exercise. Alternatively, it is possible that fluid replacement strategies were altered by 344 players as a result of been observed during the experimentation period. However, given that study 345 experimenters were familiar with players and training environment prior to experimentation, the 346 repeated observations and failure to identify trends in fluid intake across the study period; it does 347not appear that this was the case. It should be noted a small number of players (8%) consistently 348provided urine samples that were indicative of being very dehydrated (1.000-1.1290 mOsmol/kg; 349Armstrong et al., 2010) suggesting the need for targeted fluid replacement in such a player group. 350There is some evidence of a positive correlation between pre-training urine osmolality and the 351volume of fluid ingested during a training session where fluids are freely available (Maughan et al., 3522005). Therefore, athletes who begin training with higher urine osmolality/specific gravity may be

353likely to drink more due to a greater sensation of thirst. In the current study, no relationship was 354observed between morning urine measures and subsequent fluid intake. Such findings have been 355observed previously (Maughan et al., 2007b) and may be dependent on the time period from 356morning hydration assessment and exercise onset. In this study, some games started at 3 pm and so 357it is feasible that athletes had adequate time to correct any fluid imbalance through normal sensory 358means. Unlike previous findings (Maughan et al., 2007b; Hamouti et al., 2010), a modest but 359positive association was observed between the amount of sweat lost and volume of fluid consumed 360(Greenleaf, 1982), indicating that sensation of thirst and ample fluid breaks was capable of 361offsetting major disturbances in fluid balance. Further work should be carried out to determine the

362strength of this relationship in athlete groups whom display a wide range in sweat rates.

363

364Data from tennis (Bergeron, 2003) and American football (Stofan et al., 2005; Horswill et al., 3652009), have suggested that athletes who sweat profusely and have a high sweat sodium 366concentration may be more likely to experience muscle cramps. However, few if any studies have 367investigated whether or not electrolyte losses vary throughout a condensed tournament environment 368when replacement of salts lost in sweat may be of added importance to the recovery process. In the 369current study, mean sweat electrolyte concentrations were on the lower end of those reported 370previously (Maughan et al., 2005; Hamouti et al., 2010). Very few studies have investigated 371electrolyte losses in elite female athletes (Kilding et al., 2009) and it is possible that gender 372differences may account for observed differences. Although speculative, given that players in this 373study voluntarily chose to consume plain water during exercise, it is also possible that this may, in 374part, account for the lower mean sweat sodium ($38 \pm 10 \text{ mmol}.L^{-1}$) concentrations observed (Sigal 375& Dobson, 1968). A large inter-individual variation was observed for all sweat electrolyte 376concentration (Table 2.) as noted elsewhere (Maughan et al., 2007b). Given that players were rather 377homogenous in terms of fitness level (had similar daily exercise programs), dietary intake (same

378food menus) and degree of acclimation, it's possible that genetic factors may underpin this finding. 379As observed previously, no relationships were found between whole body sweat rate and sweat 380concentration of any electrolyte (Maughan et al., 2007b; Kilding et al., 2009). This is in contrast to 381recent data in tennis which suggests sweat sodium concentration is related to an individual athletes 382sweat rate (Bergeron, 2014). Given that a significant correlation did exist between player sweat 383rates and time spent exercising at higher intensities, present data are suggestive that time exposure 384to intense exercise may, in part, have a role in overall electrolyte loss (Hamouti et al., 2011).

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386Regional differences in sweat electrolyte composition were observed as previously reported 387(Maughan et al., 2007b). In this study, lower back torso sodium concentrations were observed 388compared with other collection sites. These findings are in contrast to previous work carried out on 389male cyclists (Patterson, Galloway & Nimmo, 2000) and both male and female atheletes cycling in 390a heat chamber (Baker et al., 2009). Future studies should be carried out on female athletes 391undertaking differing exercise modes to explore further why such regional differences occur. While 392 previous studies which have primarily focused on either once off sweat composition measurements. 393the current study assessed electrolyte losses across three games in a six-day tournament. Results 394showed that with the exception of game 4, electrolyte concentrations in sweat were consistent 395between games. With respect to game 4, both teams agreed to play across 3 x 30 minute period (90 396minutes total) for technical development reasons. Although between game sweat rates did not differ 397in this study and stability of electrolyte concentration has been observed across varying exercise 398durations (Montain, Cheuvront, & Lukask., 2007), current data suggests that length of sweat patch 399sampling time does need to be taken into account when assessing electrolyte loss in athletes. A 400significant association was observed for total sodium deficit $(1.03 \pm 0.43 \text{ g})$ and per-cent change in 401BM. Although statistically significant, only a moderate correlation was observed ($r^2 = 0.367$) which, 402perhaps is not surprising given that salt losses through exercise are influenced by both sweat loss

403and electrolyte concentration. Although moderate sodium deficits were incurred, a large variability 404in sodium loss was observed (range: 0.3-2.37 g). Given that the current player group refrained from 405consuming electrolytes during exercise, recommendations to a few select individuals (whom 406experienced high sodium losses) on consumption of electrolyte containing beverages during 407exercise to may be required reduced the sodium deficit and to better retain ingested fluid. This may 408be particularly relevant during a tournament scenario where players have a game every other day.

409

410Present results indicate that moderate sweat and electrolyte losses occur during handball games 411played in temperate conditions and the magnitude of such losses appear related to time spent 412exercising at higher intensities. Furthermore, it seems evident that specific approaches need to be 413considered for certain individuals, not only to minimise fluids losses but also to avoid 414overconsumption of fluids which might have negative effects on performance. Lastly, our data 415seems to suggest that the cardiovascular demands of handball games in women are similar to those 416reported for men, albeit further studies are needed to quantify time-motion characteristics of elite 417women's handball.

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527Tables

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529Table 1. Environmental conditions: Temperature (°C) and relative humidity (%) taken at 15 min intervals during the 530games.

531

532Table 2. Between game comparison of sweat electrolyte concentrations (mmolxL⁻¹) per collection site. *Significantly 533different from game 4, P < 0.05. # Significantly different from game 5, P < 0.05.

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535Figures

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537Figure 1. Testing schedule across the tournament.

538

539Figure 2. Mean and maximal total and effective HR during the first and second halves and total match time. Data 540presented as absolute and relative to individual maximal HR values (means \pm SD). * Significantly different from total 541HR (P = 0.00). Inset showing effective and total HR data recorded during training.

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543Figure 3. Percentage of effective match time spent at different interval percentages of player maximal HR in the first 544and second halves. Values are means \pm SD.

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546Figure 4(a). Relationship between measured pre-exercise urine refractive index and measured fluid intake during 547training/games, $r^2 = 0.014$, P = 0.307.

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549Figure 4(b). Relationship between morning urine refractive index and percentage change in player body mass during 550training/games, $r^2 = 0.159$, P = 0.001.

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552Figure 5. Relationship between volume of fluid consumed during training/games and the amount of sweat loss (L), $r^2 = 5530.121$, P = 0.001.

555Figure 6. Rate of fluid intake and sweat loss in players across the tournament. * Significantly higher than training 556sessions (P<0.05). Morning urine refractive index values per game throughout tournament also provided. Values are 557mean \pm SD.

Table 1. Environmental conditions: Temperature (°C) and relative humidity (%) taken at 15 min intervals during games

	Game 1	Game 2	Game 3	Game 4	Game 5	Training*
Temperature (°C)	21±0	25 ± 0	24±0	22±2	21±1	23±2
Relative humidity (%)	30 ± 2	27 ± 1	30±1	33±2	27±3	31±0

Data are mean ± SD. *Averaged data from two training sessions at same venue.

Table 2. Between game comparison of sweat electrolyte concentrations (mmol·L-1) per collection site. (SD): standard deviation.

	Game 2		Game 4		Game 5				
	Mean <i>(SD)</i> Range		Mean <i>(SD)</i> Range		Mean (Mean <i>(SD)</i> Range		Mean <i>(SD)</i> Range	
Mean Sodium	36 <i>(13)</i>	19-70		40 <i>(8)</i>	27-54	*35 <i>(7</i>)	24-47	38 (10)	23-57
Arm	38 (14)	16-60		44 (12)	29-67	39 (13) 18-61	41 (13)	16-66
Chest	40 (14)	16-55		46 (11)	23-64	*40 (10)) 24-54	42 (12)	16-65
Back	31 (12)	14-60		33 (10)	17-46	*27 (7)	14-40	31 (10)	14-60
Thigh	36 (15)	19-75		42 (11)	26-70	38 (8)	24-48	39 (12)	19-75
Mean Chloride	*24(16)	2-54		31 (9)	16-47	*25 (8)	15-37	27 (11)	11-46
Arm	*26 (16)	2-50		33 (14)	12-58	31 (12)) 16-52	30 (15)	3-57
Chest	*29 (18)	2-55		38 (11)	19-53	*33 (10)) 21-46	34 (13)	3-55
Back	20 (13)	2-27		24 (10)	8-40	20 (9)	9-35	22 (11)	2-47
Thigh	*20 (15)	6-52		27 (11)	10-46	*21 (8)	6-33	23 (11)	6-52
Mean Potassium	5 (2)	3-7		6 (1)	4-7	*5 (1)	3-7	5 (1)	3-7
Arm	*6 (2)	3-9		8 (2)	5-12	7 (3)	3-12	7 (2)	3-12
Chest	5 (0)	4-5		5 (1)	3-6	*4(1)	2-5	4 (1)	2-6
Back	4 (1)	2-6		5 (1)	4-7	*5(1)	3-6	5 (1)	2-7
Thigh	#6 (1)	5-8		7 (3)	3-14	*5(1)	3-6	6 (2)	3-14

*Significantly different from game 4, P < 0.05. *Significantly different from game 5, P < 0.05.



578Figure 1



93%







