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1	Daily Changes of Resting Metabolic Rate in Elite Rugby Union Players.
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#### 21 Abstract

INTRODUCTION: Preparation for competitive contact sport has been extensively
researched. There are, however, limited data to guide players as to how the demands of their
sport affect the energy requirements of recovery. We aimed to provide novel data on changes
in resting metabolic rate (RMR) in contact sport athletes and relate these to the physical
demands of training and competition.

27 **METHODS:** 22 Elite professional Premiership Rugby Union players were recruited to the 28 study. Indirect calorimetry (Vyntus CPX canopy, CareFusion) was used to measure RMR 29 each morning of the competitive game week, in a fasted, rested state. External loads for 30 training and game play were monitored and recorded using global positioning systems 31 (Catapult Innovations, Australia), whilst internal loads were tracked using rate of perceived 32 exertion scales. Collisions were reviewed and recorded by expert video analysts for contacts 33 in general play (breakdown and tackle area) or the set piece (scrum or maul). 34 **RESULTS:** There were significant (p=0.005) mean increases in RMR of ~231kcal the morning after (GD+1) and 3 days after the game (GD+3), compared with the day before the 35 36 game (GD-1). The players were exposed to internal and external loads during the training 37 week comparable to that of a match day, however, despite the equivocal loads between 38 training and game play, there were no significant increases in RMR following training. 39 **CONCLUSION:** The collisions experienced in rugby match play are likely to be responsible 40 for the significant increases in RMR at GD+1 and GD+3. Consequently, the measurement of

41 RMR via indirect calorimetry may provide a novel non-invasive measure of the effects of

42 collisions. This study provides a novel insight to the energy requirements of recovering from

43 contact sport.

**Keywords:** Damage, Team Sport, Contact, Injury, DOMS, Soreness

#### 46 Introduction.

47 Rugby Union is a dynamic and combative team sport participated in globally (1). Two teams of 15 players, broadly categorised as forwards (n=8) and backs (n=7) contest a match for 80 48 49 minutes (1). The sport is comprised of intermittent, high intensity activities incorporating high speed running, sprinting, accelerations and decelerations (2-4). Rugby Union also involves 50 51 collision-based activities at the tackle area (tackle and breakdown contest) and the set piece 52 (scrum and maul). Time motion analysis and global positioning systems (GPS) studies report that forwards experience  $\sim 60\%$  more high level impacts during contact situations than backs 53 (4). However, there are significant limitations of using GPS technology to determine contact 54 55 occurrence and quantitative measurement of force, rendering it unreliable to determine the 56 physical strain placed on the players (5).

57 Whilst the technology to accurately quantify physical collisions in rugby is currently lacking, 58 the recognition of their impact made upon the athlete is not (6). The forces and mechanical stress in rugby can cause exercise induced muscle damage (EIMD) and impact induced muscle 59 damage (IIMD) which may be distinct in their symptomology and recovery time course (7). 60 61 These physical collisions have been shown to increase indirect markers of muscle damage (8, 62 9), reduce neuromuscular function (10, 11), and increase perception of muscle soreness (11). 63 Sport scientists have examined a wide array of modalities to enhance recovery from the 64 damaging collisions of rugby match play, some of which may mildly alleviate symptoms (12). 65 However, despite multiple interventions being implemented, we have reported that elite rugby players are in pain every day throughout a competitive rugby season (13). It is therefore crucial 66 67 that accurate and quantitative markers are developed to assess the extent of the IIMD to allow 68 more targeted interventions to be developed. One potential candidate is assessing the energy 69 expenditure of players given that the total energy expenditure (TEE) of young rugby league 70 players was 5% higher when training weeks involved collisions (14).

71 Resting metabolic rate (RMR) is the primary component of TEE and is the energy expended to 72 maintain homeostasis at rest. Indirect calorimetry (IC) requiring both oxygen ( $\dot{V}O_2$ ) and carbon 73 dioxide ( $\dot{V}CO_2$ ) to be measured is the most accurate method of assessing RMR (15). Large 74 variations in the estimation of RMR using prediction equations have been noted in a variety of 75 sports (16), especially athletes with a high fat free mass (17) such as rugby (18-20). It is 76 therefore imperative that RMR is accurately measured rather than predicted using equations. 77 Importantly, much of the existing understanding around effectively calculating an athlete's energy requirements are based upon studies which primarily utilise recreational or youth 78 athletes and are thus limited by lower training ages and exposures to lower absolute intensities 79 80 of work. To our knowledge there are no data on the daily variations in RMR across an entire 81 competitive match week in any sport, including positional differences. It is therefore crucial that potential changes in RMR are explored in highly trained professional athletes with indirect 82 83 calorimetry performed prior to and the days following a competitive fixture.

To facilitate recovery, it is essential that rugby players are provided with the correct nutrition 84 85 in terms of both the total energy intake and the provision of recovery promoting foods. The 86 majority of nutrition research in rugby has focussed upon preparation for match play, ensuring 87 muscle glycogen concentrations are optimal for performance (21). It appears elite players now 88 have a good understanding of this (22), however, the nutritional intakes in the days following 89 a match are much more variable (22) with many players decreasing total energy intake the day 90 after a game. If muscle damage arising from match play causes an increase in energy 91 requirements in recovery, current guidelines could be underestimating player's needs post 92 competition.

93 To this end, the objectives of the present study were twofold. 1) To assess, for the first-time, 94 changes in RMR in an elite group of professional rugby union players measured throughout a 95 competitive week, including the days before and after a professional game using indirect 96 calorimetry. 2) To explore the relationship between game day factors, (e.g. the number of 97 physical collisions), and changes in RMR. These data would provide more accurate 98 information into the energy requirements of players in the days after a game, which could help 99 recovery strategy, as well as providing a novel non-invasive assessment of the effects of the 910 physical collisions upon the players.

101 Methods

#### 102 **Participants**

103 A convenience sample of twenty-two healthy elite rugby union players, all members of an 104 English Premiership squad, were recruited for this study. The participants included six 105 internationals, and many established Premiership or Super 15 players (mean  $\pm$ SD, age; 25.7 106  $\pm 4.1$  years, body mass; 104.6  $\pm 12.6$  kg). Five participants were excluded from the analysis 107 having sustained an injury during games which prevented them from completing all aspects of 108 the study. All playing positions were covered in the remaining 17 players who were eligible 109 for the full study analysis. All participants gave written informed consent prior to commencing 110 the study. Ethical approval (18/SPS/004) was granted by the university research ethics committee at Liverpool John Moores University, UK. 111

#### 112 Research Design

The study was designed to allow RMR to be measured within the training schedules of elite rugby players during a complete microcycle. Timepoints throughout the study are described relative to game day (GD) using +/- symbols for days before (-) and days after (+) GD. Due to the timing of team selection defining when recruitment could occur, the first measurement was taken at GD-2. Measurements were then repeated every day, apart from the game day itself, as this was deemed too disruptive to the players' habitual routine. Table 1 details the training schedule for the match week. Seven microcycles were used to attain the total data set, with all games played on the Saturday afternoon (Game Day). This ensured that the training schedules throughout the microcycle were the same and there were no conflicting kick-off times, which would alter the time relative to match play of the subsequent measures. Internal and external loads for training and match play were recorded throughout the week. The weeks chosen were throughout the middle of the season (weeks 13-30) so the players were accustomed to the training load and rigours of match play.

## 126 Resting Metabolic Rate

127 The RMR of participants was assessed 6 times in total. All measures were completed at the 128 same time between 7-9am and players arrived after an overnight fast, with their last meal at 129 least 8 hours prior to measurement. Players awoke and came straight to the training ground as 130 per reliable outpatient protocol (23). To ensure best practice, a private room was established 131 at the training facility away from the main building where temperature was maintained at 21-132 23 °C, the room was dimly lit, and quiet (15). Players lay in a comfortable supine position and 133 were reminded to stay awake. A twenty minute resting period was prescribed, as the minimum 134 sufficient time to achieve rest (24). A ventilated hood was employed rather than mouth piece 135 and nose clip to reduce day-to-day variance (25). The coefficient of variance for our protocol 136 was measured at 1.13% for RMR and 1.62% for RER. The ventilated hood was placed over 137 the head of the athlete and expired gas was analysed using the dilution canopy method (Vyntus 138 CPX canopy, CareFusion, Hoechberg, Germany). The gas analyser was calibrated every day 139 using the manufacturer's automated flow and digital volume transducer calibration (15.92% 140  $O_2$  and 5.03%  $CO_2$ ). The first 5 minutes of measurements were discarded following best 141 practice guidelines (15). Measurements were subsequently recorded for 15 minutes continuously at 10 second intervals for VO2 and VCO2. Data were exported into Microsoft 142 143 Excel (2018, Seattle, USA), and mean respiratory exchange ratio (RER) across the

measurement period generated, with the calorific value, carbohydrate and fat oxidation ratesdetermined according to the table of Zuntz (26).

## 146 Measurement of lean body mass

Lean body mass was measured using a dual-energy-X-ray absorptiometry (DXA) fan beam
scanner (Hologic Horizon W, Hologic, Bedford, MA), with scanning and analysis performed
by the same trained individual using Apex software version 13.5.3.1(Hologic, Bedford, MA).
Players were scanned twice during the period of data collection for this study and the scan
corresponding closest to their week of participation used, which was no longer than 4 weeks.
Measurements were taken first thing in the morning prior to eating, drinking, or exercise and
protocols implemented to maximise reliability of positioning (27).

## 154 *Training and Match loads*

155 Internal loads for each training day and the game day were assessed by the session rating of 156 perceived exertion (sRPE) using a modified Borg scale (28). This RPE of the training session 157 was multiplied by the training duration to calculate a player load in arbitrary units (sRPE; AU) 158 (28). External demands of all rugby training sessions and match play were recorded using 159 micro-technological units worn by players containing GPS (10Hz) and accelerometer (100Hz) 160 (Catapult Innovations, Melbourne, Australia). Data were downloaded and analysed using 161 Catapult Sprint software (Catapult Innovations, Melbourne, Australia). The total distance 162 covered, number of high-speed efforts (>60% positional average) and the number of very high-163 speed efforts (>80% individual average) were recorded (29, 30). The GPS sampling frequency 164 of 10Hz is the most reliable in team sports measuring high speed running activities (31).

165 Contacts were analysed in match play by a professional rugby union analyst with over five 166 years' experience working in English domestic and European rugby using NacSport (Analysis 167 Pro, UK). The potential collisions were then further reviewed by an expert ex-professional 168 player with 15 years and over 250 matches played in English domestic, European and International Rugby Union. The games were reviewed to ensure contacts recorded involved an actual collision. For example, a scrum may be analysed as a single contact but there may have been more than one engagement process involving a full collision before the match restarted. Some players may also be analysed as having been involved at a breakdown to keep possession, but they may not necessarily have endured a collision as part of this. The nature of these were then also accounted for as either set piece based (scrum or maul), or general phase play (breakdown and tackle area).

#### 176 Data analysis

177 All data are presented as mean ( $\pm$  SD). All statistical analyses were completed using SPSS 178 (Version 24 for Windows, SPSS Inc., Chicago, IL). A one-way repeated measures ANOVA 179 was used to compare all gas exchange measures and the work completed by players throughout 180 training days and during the competitive game day. The tests of within subjects' effects provided values for Mauchly's test for sphericity. If this was violated, then a Greenhouse-181 182 Geisser correction was used. The difference between means were tested at a significance level of p<0.05. The least significant difference (LSD) was used post hoc to compare specific time 183 184 points when the ANOVA revealed a significant difference between measures over the week. 185 This was examined in the whole group (n=17), sub-groups forwards (n=11) and backs (n=6). 186 A Spearman's correlation was run to assess any associations between changes in RMR 187 throughout the microcycle, with the metrics of physical load and collision data gathered from 188 the competitive match play (n=17). A Spearman's rank-order correlation coefficient value  $(r_s)$ was generated and this was tested at p < 0.05 to test the significance of any relationships found 189 190 (32).

191

## \*\*TABLE 1 ABOUT HERE\*\*

193	Results
194	Training and match demands
195	The training schedule and structure of sessions can be seen in Table 1 with the internal and
196	external demands of the week in Table 2. It should be noted that data are presented as n=14 for
197	these analyses due to faults with GPS data collection, resulting in lost running metrics for some
198	training sessions in three of the participants.
199	**TABLE 2 ABOUT HERE**
200	Player Load
201	There was no significant difference in player load on GD+3 compared with GD. This was also
202	true for the sub-groups of forwards and backs. The player load on all other days of the training
203	week were significantly lower than the game day in the whole group and when subdivided into
204	forwards, and backs.
205	High Speed Running Distance
206	In the whole group, there was no significant difference in high speed running distance covered
207	on GD+3 compared with GD. In the forwards sub-group, there was only significantly less HSR
208	distance covered on GD-1(p=0.001) and GD+2(p=0.013) compared with GD. In the backs sub-
209	group, there was significantly less HSR distance covered on GD-2 (p=0.005), GD-1
210	(p<0.0005), GD+2 (p<0.0005), and GD+3(p=0.019) compared with GD.
211	Number of High Speed Running Efforts
212	In the whole group, there were significantly fewer HSR efforts on GD-2 (p=0.002), GD-1
213	(p<0.0005), GD+2 (p<0.0005), and GD+3 (0.031) compared with GD. In the forwards sub-

group, significantly fewer HSR efforts were completed on GD-1 (p=0.001) and GD+2 

215 (p=0.014) compared with GD. In the backs sub-group, significantly fewer HSR efforts were

completed on GD-2 (p=0.003), GD-1 (p=<0.0005), GD+2 (p=0.001), and GD+3 (p=0.001)

compared with GD.

- 218 Very High Speed Running Distance
- In the whole group, VHSR distance was only significantly lower on GD-1 (p=0.002) and GD+2
- 220 (p=0.002) compared with GD. Within the forwards sub-group, there was no significant
- 221 difference in VHSR distances covered on any day compared with GD. The backs covered
- significantly fewer VHSR metres on GD-1 (p=0.005) and GD+2 (p=0.006).
- 223 Very High Speed Running Efforts

In the whole group, the number of VHSR efforts completed was only significantly lower on GD-1 (p=0.003), and GD+2 (p=0.013) compared with GD. In the forwards sub-group, there was no significant difference in VHSR efforts on all training days compared with GD. In the backs sub-group, there were only significantly less VHSR efforts on GD-1 (p=0.001), and GD+2 (p=0.013) compared with GD.

- \*\*FIGURE 1. a-f ABOUT HERE\*\*
- 230 \*\*TABLE 3 ABOUT HERE\*\*
- 231 Changes in resting metabolic rate

Changes in RMR adjusted for lean body mass across the microcycle can be seen in Figure 1a whilst the absolute (kcal·day<sup>-1</sup>) and relative (kcal·kg·day<sup>-1</sup>) RMR measures are displayed in Table 3. Lean body mass (measured by DXA) was 74.8 $\pm$ 7.4kg for the whole group, 78.2 $\pm$ 5.6kg for the forwards, and 68.6 $\pm$ 6.0kg for the backs. In the whole group, there was a significant increase in RMR from GD-1 to GD+1 (p=0.005) and GD-1 to GD+3 (p=0.04). In the forwards sub-group, there was a significant increase in RMR between GD-1 to GD+1 (p=0.017) and 238 GD-1 to GD+3 (p=0.045). However, in the backs sub-group, there was no significant difference

in RMR at any time point across the week.

## 240 Changes in respiratory exchange ratio

Changes in RER across the microcycle can be seen in Figure 1b. In the whole group, there were significant increases at GD+2 (p=0.030) and GD+3 (p=0.006) compared with GD-1. In the positional subgroups there were no significant differences across the microcycle p=0.065 and p=0.177 for forwards and backs respectively.

#### 245 Changes in VO<sub>2</sub> and VCO<sub>2</sub>

246 Figures 1c and 1d show the measures of VO<sub>2</sub> and VCO<sub>2</sub>. There were significant increases in 247  $VO_2$  in the whole group at GD+1 (p=0.008) and GD+3 (p=0.041) compared with GD-1. These 248 significant increases were also observed in the forwards at GD+1 (p=0.025) and GD+3249 (p=0.027) compared with GD-1. There were no significant differences for  $VO_2$  in the backs 250 subgroup across the week. There were significant increases in  $VCO_2$  in the whole group at 251 GD+1 (p=0.008), GD+2 (p=0.01), and GD+3 (p=0.001) compared to GD-1. These significant 252 increases were also observed in the forwards at GD+1 (p=0.037) and GD+3 (p<0.001) 253 compared to GD-1. There were no significant differences across the week in measures of  $VCO_2$ 254 in the backs.

## 255 Changes in carbohydrate and fat oxidation

Measures of carbohydrate and fat oxidation are displayed in Figures 1e and 1f. Carbohydrate oxidation significantly increased at GD+2 (p=0.044) and GD+3 (p=0.003) compared with GD-1 in the whole group. In the forwards a significant increase was measured at GD+3 (p=0.003) compared with GD-1, whilst there were no significant differences across the microcycle in the backs for carbohydrate oxidation. Fat oxidation decreased significantly at GD+3 (p=0.029) in the whole group and at the same time point in the forwards (p=0.028) compared with GD-1. 262 There were no significant differences measured for fat oxidation across the microcycle in the263 backs.

264 \*\*TABLE 4 ABOUT HERE\*\*

## 265 Associations of match demands with changes in metabolic measurements

266 Table 3 displays the Spearman's coefficient associations between the physical match demands,

and changes in RMR. In the whole group, there were no significant associations found between

268 phase contacts, total contacts, player load, HSR meters, HSR efforts, VHSR meters, VHSR

- efforts and the change in RMR observed between GD-1 to GD+1. This was also true when the
- 270 positional sub-groups of forwards and backs were analysed.

#### 271 Discussion

272 The aim of the present study was to assess changes in RMR in an elite group of professional 273 RU players measured throughout a competitive week and explore the impact of game day 274 factors on changes in RMR. To this end, we monitored RMR using indirect calorimetry 275 alongside game day and training demands in 22 Premiership RU players throughout a game 276 week. We report, for the first-time, that RMR increased significantly following elite rugby 277 union match play, a change that was not observed following intense training with the same 278 training loads. These data therefore illustrate that changes in RMR following match days exist, 279 reflecting a yet unreported increased energy demand in the days after a game of elite rugby and 280 allows the development of individualised nutritional strategies to help facilitate recovery. 281 Furthermore, increased RMR may also represent the physical collisions of match play and 282 indeed could suggest that RMR may be used as a non-invasive marker of muscle damage.

283 We have reported a mean increase in RMR following match play of  $\sim 231$  kcal per day at GD+1, 284 a 10% increase from GD-1. We are confident this represents a truly significant increase given that it is greater than the suggested 6% required as meaningful change using the canopy method 285 286 (25). The rigour in our protocol also resulted in a lower coefficient of variance than reported 287 previously (25). Importantly, these increases in RMR were due to significant increases in VO2 288 and VCO2 and are not merely EPOC being measured as increased VO2. The range of increased 289 RMR was large, with individual responses between 240-1000kcal. The greatest increases in 290 RMR were seen in the forwards, who underwent more physical collisions during a game at the 291 scrum, maul and tackle area vs. backs (4). The whole group, and forwards positional group, 292 also experienced increased RMR which remained elevated 3 days post-game. This sustained 293 increase at GD+3 may be a result of the lower limb resistance training session on GD+2 given 294 that resistance training, especially with an eccentric component, has been shown to increase 295 RMR (33). It is possible that this sustained increase in RMR, as a result of the resistance

training session, negatively affected the recovery from match play, therefore extending the period during which RMR remained elevated, although this suggestion remains speculative and requires further investigation.

299 Along with changes in RMR in the days after the game we also report significant changes in 300 RER. The increased RER at GD+2 and GD+3 corresponds with significant increases in resting 301 carbohydrate oxidation coupled with a significant reduction in fat oxidation at GD+3. These 302 significant changes in carbohydrate oxidation are occurring at a time where markers of muscle 303 damage and soreness typically peak following match play (8, 9, 11). Muscle damage induced 304 reductions in glucose transport may result in a decreased whole-body glucose tolerance which 305 has been reported after a laboratory based muscle damage protocol (34). It should also be 306 recognised that the inflammatory cytokine activity associated with muscle damaging exercise, 307 together with the presence of various cell types such as neutrophils and macrophages (35), may 308 alter substrate oxidation in the recovery period (36). Taken together, we have demonstrated 309 increased RMR and altered carbohydrate oxidation, following match play, which suggests that post-exercise nutrition should be specifically tailored to the unique metabolic demands of this 310 311 time period. Moreover, we have shown highly individual responses with some players 312 increasing their RMR by 1000kcal. It is crucial to identify such players and tailor their dietary plans and recovery strategies accordingly. 313

Given that the participants in the present study were full-time professional players, in the middle of a competitive playing season, it was not possible to either control or record dietary intake. It is possible that some of the differences in RMR and RER seen between the forwards and backs could have been a result of differing diets of the 2 sub-groups. However, whilst there is evidence that the thermic effect of food and the total energy content of a meal may alter resting metabolic rate measures (15) we do not believe that the player to player variations in diet would have any meaningful effects on RMR or RER in the present study. Previous research 321 has reported that a large meal containing 1300kcal had negligible effects upon measuring RMR 322 and RER when measured 7 hours later, and in lean male subjects both measures had returned to baseline at 8 hours following this meal (37). Given that both the forwards and backs in the 323 324 present study had undergone a minimum of an 8 hour fast prior to having their RMR and RER 325 assessed, it is unlikely that differences in diet would be a primary contributor to the observed 326 changes. Moreover, we believe that this group of players consumed a more than adequate 327 energy availability as indicated by no major changes in body mass over the testing periods. 328 This group are unlikely to be in low energy availability, however, future studies should attempt to measure or control dietary intake to fully explore this hypothesis. 329

330 We propose that the muscle damage as a result of elite rugby union match play could be a key 331 factor in accounting for the changes in metabolism we have witnessed. By carefully monitoring 332 the internal and external demands of the competitive week we have shown that when contact 333 sport athletes are exposed to comparable player load (including HSR and VHSR metrics) to 334 that of a match day but without the physical collisions, there is no change in RMR in the following days. We therefore speculate that the collisions encountered on a game day could be 335 336 responsible for the significant changes in RMR reported at GD+1. This may account for the 337 increases in TEE previously observed in youth players when a training session contained 338 collisions similar to that of match play (14).

When we investigated the positional groups of forwards and backs there were differences in how they reacted to match play. The backs sub-group did not show any significant changes in RMR or RER post-match, albeit they did show a similar pattern across the week as seen in the forwards sub-group. The backs did not experience as many contact incidents as the forwards as has previously been shown (4), and they were not involved in the static exertions of the scrum and maul which are potentially damaging. These positional differences may further substantiate our hypothesis that the contact-based activities are responsible for the metabolicchanges reported here.

347 The total number of contacts were rigorously evaluated; however, the Spearman's correlations 348 did not show any significant correlations of changes in RMR with the match demands or 349 collisions experienced. There was one back who exhibited a large increase of  $\approx$ 796kcal in 350 RMR. Although the actual number of contacts performed by this player were not significantly 351 different to the mean of the backs group, subjective analysis of these collisions (by experienced 352 rugby staff) classified the magnitude and intensity of these as being much greater than typical. 353 Examples like this, coupled with the current inability to accurately quantify collision activities, 354 emphasises the need for a practical measure of the impact contact sports have upon these 355 athletes to be developed.

## 356 *Practical implications.*

357 From an applied perspective the periodisation of nutrition throughout microcycles to optimise 358 adaptation and ultimately performance is well established under the 'Fuel for the work 359 required' paradigm (38). The novel data presented here could enhance the application of this 360 in team sports, especially those involving muscle damage due to collision-based activities. 361 Even using a modest physical activity level (PAL) of 1.3-1.4 for a GD+1 rest day, would 362 translate these findings into a required increase in energy intake of >300kcal. This, on a day 363 where the continued restoration of muscle glycogen is a primary concern, in a population 364 who habitually appear to consume lower than the recommended carbohydrate intakes, may 365 require a conscious intervention (22). Carbohydrate intake as part of an in-season week in 366 elite rugby union players appears to be 3g/kg on GD+1 (22), therefore an extra ~70g 367 carbohydrate could be an increase of  $\sim 20\%$  required on that day. We speculate the timing of 368 carbohydrate feeding may also require further investigation though, if indeed substrate 369 oxidation is altered until the muscle damage due to match play is resolved (34, 39).

370 Given that the true definition of resting metabolic rate involves 'strict and steady resting 371 conditions' it could be argued that the present study did not actually measure RMR at any 372 time point where in fact Morning Metabolic Rate (MMR) was actually measured. Indeed, it 373 could be argued that rugby players (and indeed many athletes) during a competitive season 374 are never truly at 'rest' bringing about methodological questions over when during a training 375 period RMR should be measured to accurately predict energy requirements. A protocol 376 according to best practice and adhering strictly to a minimum rest time, fasted measurement and proper outpatient protocols as per resting metabolic rate are crucial for reliability but this 377 378 measure may need to be categorised differently (15, 23). In the applied world, the term 379 'Morning Metabolic Rate' may be a more accurate description of what is actually being 380 measured and future studies may choose to adopt this terminology.

#### 381 *Conclusions*.

382 In conclusion, the present study has for the first time assessed the resting metabolic rate of 383 elite rugby union players across a competitive match week using indirect calorimetry. We 384 report a significant increase in the RMR of these contact sport athletes in the days after match 385 play. There were also significant shifts in RER at two and three days after competition. We 386 propose these changes could be attributed to the collisions experienced in match play rather than the internal and external loads the athletes are exposed to throughout the microcycle. 387 The metabolites and markers of these muscle damaging actions need to be researched further 388 389 to help guide athletes as how best to feed their recovery after competition. This research is 390 the first step in working towards a novel non-invasive marker of muscle damage. Further 391 studies need to control factors of energy availability and exercise modalities responsible for 392 the forms of muscle damage. Protocols more readily useable in the 'real world' of applied 393 performance science then need to be designed to truly shift the paradigm of athlete 394 monitoring and optimise recovery from contact sports.

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## **Conflict of interest.**

The authors reported no potential conflict of interest. The results of the current study do not
constitute endorsement by ACSM. All results presented here are done so clearly, honestly, and
without fabrication, falsification, or inappropriate data manipulation.

- 404 Word Count 4444.

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563	Table 1. The training sessions throughout the competitive micro cycle. Game Day – GD.
564	Table 2. Comparison of metrics recorded for training and match play throughout the
565	competitive micro cycle.
566	*Denotes values significantly different (p<0.05) when compared with game day (GD) shown
567	in bold.
568	Figure 1. Gas exchange measurements across the microcycle. a. RMR (kcal·kg·day <sup>-1</sup> ). b.
569	RER. c. VO2 (L/min). d. VCO2 (L/min). e. Carbohydrate (CHO) oxidation (g/min). f.
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571	Measurements displayed as mean± S.D. with individual data points for all participants.
572	Forwards – (filled black triangle), Backs- O (empty circle).
573	*Denotes significant difference ( $p < 0.05$ ) for the whole group when compared to GD-1.
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575	Table 3. Absolute and adjusted measurements of RMR across the competitive
576	microcycle for all players (n=17).
577	Table 4. Spearman's coefficient (rs) associations derived from changes in RMR between
578	GD-1 and GD+1.
579	*denotes significant p<0.05 association

Time Point	GD-3	GD-2	GD-1	GD	GD+1	GD+2	GD+3	GD+4
Purpose	Rest & Recovery	Intensity	Team Run	Match Play	Rest & Recovery	Installation	Volume	Rest & Recovery
Resistance Training Content	None	Upper Limb Strength (30 min)	None	None	None	Lower Limb Strength (45 mins)	Upper Limb Strength (45min)	None
Rugby Content	None	Specific Game Prep (35 mins) Unit Split (15- 25mins)	Agility warm- up, Execution of specific game prep at a low- moderate intensity (35min)	Individual & Team Warm Ups. Rugby Match Play (80 mins).	None	Low-moderate intensity attack shapes and defensive systems. Running top- ups for some players. (60 mins)	High Intensity throughout rugby specific drills. Units Split- Forwards - Scrum/Maul Backs - Strike plays. (75 min)	None
Targets	Recovery	Execution of tactical game specifics at a high intensity. Rehearsal of set pieces in a unit split. Forwards – Lineouts Backs – Strike and skill execution.	Execution of specifics at a lower intensity, low intensity unit rehearsal of set pieces. Forwards – Lineouts Backs – Strike plays.	Full competitive rugby match play. Target physical performance and win.	Recovery	Learning of specifics for the following fixture and recovery.	High running volume, aiming to overload running volume relative to time.	Recovery

Table 1. The training sessions throughout the competitive micro cycle. Game Day – GD.

Time l	Point	GD -2	<b>GD -1</b>	GD	GD +1	<b>GD</b> +2	GD +3	GD +4
Player	Whole	$404.07 \pm 103.88*$	$27.00 \pm 34.52 *$	622.36± 98.70		$238.14 \pm 186.90 *$	$631.07 \pm 110.67$	
Load	group							_
(sRPE x	Forwards	$391.25 \pm 83.93*$	$31.25 \pm 35.65*$	595.13±106.45	_	243.13±212.10*	$654.37 \pm 100.19$	
Time)	Backs	421.17±132.60*	$21.33 \pm 35.39*$	658.67± 81.85	Re	231.50±166.58*	$600.00 \pm 125.57$	Re
High	Whole	$168.00\pm 60.12*$	6.21±10.15*	285.43±113.09	st	63.29± 95.43*	$254.64 \pm 214.15$	st
Speed	group				Ø			Ø
Running	Forwards	$132.88{\pm}45.89$	9.25±12.10*	$215.88 \pm 96.82$	R	72.00± 102.86*	$279.63 \pm 268.19$	- 72 72
Distance	Backs	$214.83 \pm 42.94 *$	2.17± 5.31*	378.17± 45.46	e	51.67±92.63*	221.33±127.33*	e
(m)					0			0
High	Whole	$11.43 \pm 3.34*$	$0.93 \pm 1.69*$	$20.29 \pm 7.23$	< e	$4.93 \pm 6.86*$	$14.79 \pm 8.91*$	<e></e>
Speed	group				Ĵ,			Ĵ,
Running	Forwards	$10.50 \pm 3.55$	$1.38 \pm 2.07*$	16.63± 6.99		5.63± 7.05*	$15.50 \pm 10.90$	
Efforts (n)	Backs	$12.67 \pm 2.88*$	$0.33 \pm 0.82*$	<b>25.17± 4.17</b>		4.00± 7.13*	$13.83 \pm 6.18*$	_
Very High-	Whole	$17.86 \pm 16.28$	0.00*	16.50±15.89		$0.36 \pm 0.93 *$	$23.29 \pm 31.10$	
Speed	group							
Running	Forwards	$9.88{\pm}~10.64$	0.00	10.75±16.46		$0.38 \pm 1.06$	$23.75{\pm}38.99$	-
Distance	Backs	$28.50 \pm 17.10$	0.00*	$24.17 \pm 12.42$		$0.33 \pm 0.82*$	$22.67 \pm 19.66$	
(m)								-
Very High-	Whole	$1.29 \pm 0.91$	0.00*	1.14± 1.17		$0.14 \pm 0.36*$	$1.29 \pm 1.14$	
Speed	group							-
Running	Forwards	$1.13 \pm 0.99$	0.00	1.00± 1.51		0.13± 0.35	$1.25 \pm 1.28$	-
Efforts (n)	Backs	$1.50 \pm 0.84$	0.00*	$1.33 \pm 0.52$		$0.17 \pm 0.41*$	$1.33{\pm}1.03$	

Table 2. Comparison of metrics recorded for training and match play throughout the competitive micro cycle.

\*Denotes values significantly different (p<0.05) when compared with game day (GD) shown in bold.

Time Point	GD-2	GD-1	GD+1	GD+2	GD+3	GD+4
Absolute RMR (kcal)	2318±182.1	2313±283.0	2544± 396.9	2391±274.2	2424± 312.0	2327± 305.3
Adjusted RMR (kcal·kg·day <sup>-1</sup> )	31.2± 3.0	31.1±4.7	34.1± 5.3	32.1±4.1	32.5± 4.2	31.3± 4.7

Table 3. Absolute and adjusted measurements of RMR across the competitive microcycle for all players (n=17).

Timepoints for comparison	Group	p Phase Contacts		Total Contacts P (Phase+set (s piece)		Player Load (sRPExTime)		HSR (m)		HSR (efforts)		VHSR (m)		VHSR (efforts)	
		rs	р	rs	р	rs	р	rs	р	rs	р	rs	р	rs	р
Change in	Whole	0.05	0.84	0.23	0.38	-0.17	0.95	-0.13	0.62	-0.26	0.31	-0.11	0.97	0.19	0.48
RMR GD-1 to	Group														
GD+1	Forwards	-0.10	0.77	0.16	0.63	-0.19	0.58	-0.28	0.40	-0.24	0.47	0.19	0.57	0.32	0.34
	Backs	0.09	0.87	0.09	0.87	0.34	0.51	0.37	0.47	-0.44	0.39	-0.09	0.87	0.00	1.00

Table 4. Spearman's coefficient (rs) associations derived from changes in RMR between GD-1 and GD+1.

\*denotes significant p<0.05 association.



Figure 1. Gas exchange measurements across the microcycle. a. RMR (kcal·kg·day<sup>-1</sup>). b. RER. c. VO2 (L/min). d. VCO2 (L/min). e. Carbohydrate (CHO) oxidation (g/min). f. Fat oxidation (g/min).

b

Measurements displayed as mean± S.D. with individual data points for all participants.

Forwards – (filled black triangle), Backs- O (empty circle).

\*Denotes significant difference (p < 0.05) for the whole group when compared to GD-1.

+Denotes significant difference (p < 0.05) for the forwards group when compared to GD-1.