Determinants of last lap speed in paced and maximal 1500 m time trials

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ABSTRACT

Purpose: The present study identified the physiological and performance characteristics that are deterministic during a maximal 1500-m time trial and in paced 1500-m time trials, with an all-out last lap. **Methods:** Thirty-two trained middle-distance runners ($n=21$ male, $\text{VO}_{2\text{peak}}$: 72.1 ± 3.2 ; $n=11$, female, $\text{VO}_{2\text{peak}}$: 61.2 ± 3.7 $\text{mL} \cdot \text{kg}^{-1} \cdot \text{min}^{-1}$) completed a 1500-m time trial in the fastest time possible (1500_{FAST}) as well as a 1500_{MOD} and 1500_{SLOW} trial whereby mean speed was reduced during the 0–1100-m by 5% and 10%, respectively. Anaerobic speed reserve (ASR), running economy (RE), the velocity corresponding with $\text{VO}_{2\text{peak}}$ ($v\text{VO}_{2\text{peak}}$), maximal sprint speed (MSS) and maximal accumulated oxygen deficit (MAOD) were determined during additional testing. Carnosine content was quantified by proton magnetic resonance spectroscopy in the gastrocnemius and expressed as a Z-score to estimate muscle fibre typology. **Results:** 1500_{FAST} time was best explained by RE and $v\text{VO}_{2\text{peak}}$ in female runners (adjusted $r^2=0.80$, $P<0.001$), in addition to the 0–1100-m speed relative to $v\text{VO}_{2\text{peak}}$ in male runners (adjusted $r^2=0.72$, $P<0.001$). Runners with a higher gastrocnemius carnosine Z-score (i.e., higher estimated percentage of type II fibres) and greater MAOD, reduced their last lap time to a greater extent in the paced 1500-m trials. Neither ASR nor MSS were associated with last lap time in the paced trials. **Conclusion:** These findings suggest that $v\text{VO}_{2\text{peak}}$ and RE are key determinants of 1500-m running performance with a sustained pace from the start, while a higher carnosine Z-score and MAOD are more important for last lap speed in tactical 1500-m races.

Keywords: TACTICAL RACING; ANAEROBIC SPEED RESERVE; MUSCLE FIBRE TYPE COMPOSITION; CRITICAL SPEED; D' ; CARNOSINE; PACING

Abbreviations: RE; running economy, $v\text{VO}_{2\text{peak}}$ minimal velocity that elicits $\text{VO}_{2\text{peak}}$, ASR; anaerobic speed reserve, MSS; maximal sprinting speed, CS; critical speed, D' ; curvature constant, $\text{VO}_{2\text{peak}}$; Peak oxygen uptake, 1500_{FAST} ; maximal 1500 m time trial, 1500_{MOD} ; moderately paced 1500 m time trial, 1500_{SLOW} ; slow paced 1500 m time trial, MAOD; maximal accumulated oxygen deficit, d ; distance run, t ; time, CV; coefficient of variation, VO_2 ; oxygen uptake, GET; gas exchange threshold, TTE; time to exhaustion, $^1\text{H-MRS}$; proton magnetic resonance spectroscopy, MRI; magnetic resonance imaging, SD; standard deviation.

INTRODUCTION

In elite 1500 m running competitions, championship races (i.e., World Athletics Championships and Olympic Games) can be won in relatively fast times ($\leq 3:34.00$), moderately fast ($> 3:34.00 - \leq 3:41.99$) or even slow times ($\geq 3:42.00$) (Sandford et al. 2019a), but with an extremely fast last lap (e.g., $< 0:51.00$). The tactical racing strategy that produces fast race times resides from a relatively even pacing profile with a sustained pace from the start (Tucker et al. 2006). Conversely, slower race times result from slow initial laps and a greater number of race surges, that precede a ‘last lap kick’ (Sandford et al. 2019a). Indeed, the ‘last lap kick’ may differentiate between successful and unsuccessful competitors in 1500 m championship races (Sandford et al. 2019a; Casado and Renfree 2018; Hanley et al. 2019); however, the physiological and performance characteristics that underpin last lap speed in tactical 1500 m races have not been fully elucidated.

Many attempts have been made to characterise the key physiological determinants of 1500 m running performance (Iwaoka et al. 1988; Ingham et al. 2008; Ferri et al. 2012; Billat et al. 1996; Billat et al. 2009). The prevailing conclusion from these studies is that aerobic characteristics, such as VO_{2peak} (Ingham et al. 2008), the minimal running velocity that elicits VO_{2peak} (vVO_{2peak}) (Billat et al. 1996) and running economy (RE) (Ingham et al. 2008; Billat et al. 1996) are the most deterministic characteristics relating to 1500 m performance. Other studies that have profiled the energy system contribution to simulated (Spencer and Gastin 2001) and actual track running (Duffield et al. 2005) 1500 m time trials report that the predominant energy system contribution is derived from aerobic metabolism (77 – 85%). However, one caveat of this literature is that 1500 m running performance has been assessed under conditions whereby the subjects are likely to be attempting to run the fastest time possible, such as a maximal time trial (Ferri et al. 2012; Duffield et al. 2005), simulated maximal time trial on a treadmill (Spencer and Gastin 2001), a season best (Billat et al. 1996; Iwaoka et al. 1988) or recent 1500 m track performance (Ingham et al. 2008; Billat et al. 2009). During these scenarios, the key physiological and performance characteristics may not align with the key attributes that are required to perform a tactical race, characterized by an extremely fast last lap, but slower initial pace.

One concept that may underpin last lap speed is the anaerobic speed reserve (ASR), which is the difference between a runner's maximum sprinting speed (MSS) and vVO_{2peak} (Bundle et al. 2003). While the ASR is negatively associated with 1500 m running speed (i.e., larger ASR equates to a slower mean 1500 m speed; $r = -0.68 \pm 0.54$) (Sandford et al. 2019c), it has been suggested that a greater ASR may be an important determinant of a fast last lap in a tactical, slow 1500 m race (Sandford et al. 2019b; Mytton et al. 2015), but this is yet to be

tested experimentally. Alternatively, another concept that may underpin performance in tactical and maximal approaches to 1500 m racing arises from the parameters of the hyperbolic relationship between running speed and the tolerable duration, which is defined by the speed asymptote (critical speed; CS) and the curvature constant (D') (Monod and Scherrer 1965). It could be purported that a 1500 m runner with a higher CS (but lower D') would benefit from a higher sustained pace from the start of the race as another competitor with a lower CS (but greater D') would be required to rely on a larger portion of their D' to maintain pace. In contrast, this competitor would likely be more suited to a reduced pace in the initial laps, in order to take advantage of a superior D' for the 'last lap kick' (Pettitt 2016). However, despite the attractiveness of this proposal, further interrogation of the relationship between these parameters and 1500 m running performance with different pacing strategies is warranted. Lastly, it is conceivable that muscle fibre typology may be an important characteristic that could relate to preferential pacing strategies in middle-distance runners. Classical work (Costill et al. 1976) has shown that international level middle-distance runners possess a much greater between-athlete variation in muscle fibre typology (percentage type I fibres) in the gastrocnemius (mean; range: 52%; 41-69%) compared to long-distance runners (69%; 63-74%) and sprinters (24%; 21-27%). Given the superior capacity for anaerobic metabolism (Pette 1985), greater buffering capacity (Parkhouse et al. 1985) and superior mechanical properties (Harber et al. 2004) in type II compared to type I fibres, possessing a greater proportion of type II fibres relative to another competitor may be a key attribute for last lap speed when the initial laps are slow, but this proposal needs further examination. Therefore, the purpose of this study was to identify the physiological and performance characteristics that are deterministic during a maximal 1500 m time trial (i.e., fastest time possible) and in paced 1500 m time trials with either moderate or slow-paced initial laps, but an all-out last lap.

METHODS

Subjects

Thirty-two trained middle-distance runners ($n = 21$ men, VO_{2peak} : 72.1 ± 3.2 mL·kg⁻¹·min⁻¹; $n = 11$ women, VO_{2peak} : 61.2 ± 3.7 mL·kg⁻¹·min⁻¹) participated in this study who had a consistent training history of at least 2 years in the 1500 m event and were without major injury interruption for the previous 3 months. The male runners had season best performance times during outdoor 1500 m track competition of $3:57.50 \pm 0:09.70$ (3:40.00 – 4:14.08 s), while the female runners achieved times of $4:39.20 \pm 0:14.52$ (4:17.44 – 5:08.10). At the time of the study, the male runners had a mean running training volume of 81.6 ± 16.2 km·week⁻¹, while the female runners

ran 64.6 ± 12.1 km·week⁻¹. All runners provided written informed consent prior to participating in this study which was approved by the Griffith University Human Research Ethics Committee.

Study design

Subjects completed a 1500 m time trial in the fastest time possible (1500_{FAST}) with lap split times recorded, and two pacing trials on an outdoor athletics track. In the pacing trials, runners were paced for the 0 – 1100 m, whereby speed was reduced on average by 5% (1500_{MOD}) and 10% (1500_{SLOW}) relative to 1500_{FAST}, while the last 400 m was completed in the fastest time possible. In addition, subjects also completed laboratory treadmill running tests to determine RE, $\dot{V}O_{2peak}$, $\dot{V}VO_{2peak}$ and the maximal accumulated oxygen deficit (MAOD). Subjects also completed additional outdoor time trials (2200 and 3000 m) for the quantification of CS and D' and a MSS assessment (flying 50 m sprint) for the determination of the ASR. All trials were conducted on separate days across a 5-wk period. Carnosine content was quantified by proton magnetic resonance spectroscopy in the gastrocnemius medialis muscle and was expressed as a sex-specific Z-score to estimate muscle fibre typology (Baguet et al. 2011).

Running time trials

The running time trials were conducted on an outdoor 400 m synthetic athletics track. All running trials were preceded by a standardized warmup that consisted of a 10 min self-paced jog which was followed by a 5-min bout of submaximal running that increased to a rating of perceived exertion of 5 (CR-10 scale) and six repetitions of 10-s strides, with a walk-back recovery (~60 s), whereby strides were defined as bouts of fast running which were to be completed at each subjects perceived 1500 m race pace. The warm-up procedures were followed by 10 min of recovery. During 1500_{FAST}, subjects ran individually and were not permitted to view their race split times and were instructed to complete the trial in the fastest time possible. Electronic timing gates (Smartspeed, Fusion Sport, Australia) were placed at the start and at the 300 m mark of the first lap in order to obtain each lap split time (0 – 400 m, 400 – 800 m, 800 – 1100 m and 1100 – 1500 m). In the paced trials (1500_{MOD} and 1500_{SLOW}), runners were paced for the 0 – 1100 m, whereby the 0 – 400 m lap speed was reduced by 10% (1500_{MOD}) and 15% (1500_{SLOW}), the 400 – 800 m split speed was reduced by 5% (1500_{MOD}) and 15% (1500_{SLOW}), while the 800 – 1100 m split speed remained equivalent to 1500_{FAST} in both trials. Subjects were instructed to complete the final 400 m in the fastest time possible. These lap paces were chosen given that they reflected the relative speed decrement of moderately fast ($>3:34.00$ - $\leq 3:41.99$) and slow ($\geq 3:42.00$) races relative to fast races ($\leq 3:34.00$), as classified as in the study of Sandford et al. (2019a). While these are clearly world class standard races, we

employed the relative difference in lap times from these races to prescribe the split speeds of the paced trials in the present study. To assist with pacing from the 0 – 1100 m mark in the paced trials, each runner had a pacer who ran 4 m in front of the subject and had access to target split times and a wristwatch (Garmin Forerunner 235, Canton of Schaffhausen, Switzerland). Members of the research team also assisted with pacing by providing splits verbally at each 200 m mark. All subjects completed a familiarization of the paced trials on a separate day prior to the actual recorded paced trials. During the paced trials, if a subject recorded a lap split time within the 0 - 1100 m that deviated from the prescribed split by ≥ 1.00 s, the trial was discarded and performed on another day. This occurred on two occasions (both 1500_{SLOW} trials).

Subjects also completed additional outdoor track time trials of 2200 and 3000 m in order to determine CS and D' . These distances were chosen in order to yield finishing times between 2 and 12 min (Hughson et al. 1984). These trials were performed with the same instructions as 1500_{FAST} and CS and D' were determined using a linear distance-time model. Subjects also performed a 50 m sprint for the determination of MSS. Linear sprint speed was evaluated over 50 m using electronic timing gates positioned at the start line and at 40 and 50m. In addition to the standardized warmup procedure, subjects also performed two 50 m running efforts at 75% and 90% of each subject's perceived maximal sprinting pace. MSS was determined as the mean speed derived from the 40 – 50 m split time. ASR was defined as the difference between the MSS and the $v\text{VO}_{2\text{peak}}$, estimated from the submaximal and maximal incremental running test (see below). Air temperature, relative humidity and wind speed/direction were recorded using a thermal environment monitor (Questemp-15 Area Heat Stress Monitor, Quest Technologies, WI). Testing was conducted at an identical time of day (6:00 – 8:00 am) for all subjects which assisted in achieving relatively consistent atmospheric conditions (air temperature range 23.8 – 25.9 °C; relative humidity 66.0 – 71.0%; wind speed 4.6 – 7.6 km·h⁻¹). Eight subjects (n = 4 male and female runners) from this study performed duplicate trials on separate days one week apart in order to determine the test-retest reliability for the 1500 m time trials. The coefficient of variation (CV) for 1500_{FAST} performance time and 1500_{MOD} and 1500_{SLOW} last lap times were 1.8%, 2.2% and 2.1%, respectively.

Maximal incremental running test

Following a warm-up (10 min at 8 – 14 km·h⁻¹) and a 5-min rest period, each subject performed an incremental treadmill (HP cosmos Saturn, Traunstein, Germany) run to volitional exhaustion starting at 10 km·h⁻¹ (women) or 12 km·h⁻¹ (men) and 1% gradient with the speed increasing by 1 km·h⁻¹ each min until volitional exhaustion. Respiratory variables were measured using a Cosmed Quark b² (Rome, Italy), which had been calibrated according

to the manufacturer's instructions (Quinton Instruments, USA). Exhaled air was analysed to determine VO_2 and VCO_2 . The gas exchange threshold (GET) was determined using the simplified V-slope method previously described (Schneider et al. 1993). $\text{VO}_{2\text{peak}}$ was determined as the highest VO_2 value using a rolling 1-min average of breath-by-breath data. The inter-day CV for GET and $\text{VO}_{2\text{peak}}$ measurements in our lab is 2.00% and 3.13%, respectively ($n = 26$; measurements taken three weeks apart).

Submaximal laboratory running test

On a separate day, subjects completed six, 4-min submaximal incremental stages on a motorised treadmill which were completed in 5% incremental speeds ranging from 85 - 110% of the GET which was determined from the maximal incremental running test. The treadmill belt was also set at 1% gradient (Jones and Doust 1996). Pulmonary (breath-by-breath) gas exchange was monitored throughout each stage, whereby the highest VO_2 value was obtained using a rolling 1-min average of breath-by-breath data. Immediately following the final stage, an earlobe blood sample was taken for the determination of blood lactate concentration ($[\text{La}]_b$) with a Lactate Pro 2 device (Arkray inc. Japan).

Calculation of running economy and $v\text{VO}_{2\text{peak}}$

Oxygen uptake (VO_2) and carbon dioxide production (VCO_2) during the final minute of each submaximal stage was used to determine RE ($\text{kcal} \cdot \text{kg}^{-1} \cdot \text{km}^{-1}$). RE was expressed as an energy cost, rather than an oxygen cost, as this accounts for differences in substrate oxidation within and between subjects and is more sensitive to changes in running speed (Fletcher et al. 2009). Updated non-protein respiratory quotient equations (Peronnet and Massicotte 1991) were used to estimate substrate utilisation ($\text{g} \cdot \text{min}^{-1}$) during the final minute. The energy derived from each substrate was then calculated by multiplying fat and carbohydrate usage by 9.75 kcal and 4.07 kcal, respectively, reflecting the mean energy content of the metabolised substrates during moderate to high-intensity exercise. The $v\text{VO}_{2\text{peak}}$ was calculated by solving the regression equation describing the individual VO_2 -running speed relationship based on the mean VO_2 values and during the final min of each 4-min submaximal incremental stage and $\text{VO}_{2\text{peak}}$ measured from the maximal incremental running test using linear regression. The inter-day CV% for RE and $v\text{VO}_{2\text{peak}}$ in our lab is 2.3% and 3.1%, respectively ($n = 26$; measurements taken three weeks apart).

Supramaximal laboratory running test

The supramaximal running test was performed at a running speed equivalent to 110% of $v\text{VO}_{2\text{peak}}$. Subjects performed a 15-min warm-up consisting of 5 min at 8 – 14 $\text{km}\cdot\text{h}^{-1}$, 5 min at 140% of the GET and 2 x 6 s running bouts at 90% of $v\text{VO}_{2\text{peak}}$, followed by 10 min of rest. Each subject wore a safety harness during the supramaximal effort which was attached to their chest to ensure maximal effort. Time to exhaustion was measured to the nearest second and blood lactate concentration was determined from earlobe samples taken at 1, 3, 5, and 7 min after the completion of the test, with the highest value obtained considered the peak blood lactate concentration.

Maximal accumulated oxygen deficit

As originally described (Medbo et al. 1988), the individual VO_2 -running speed relationship was extrapolated to estimate the VO_2 demand corresponding to 110% of $v\text{VO}_{2\text{peak}}$. The accumulated VO_2 demand was estimated by multiplying the time to exhaustion by the VO_2 demand and the MAOD was calculated by subtracting the accumulated VO_2 , measured during the exercise bout, from the estimated accumulated VO_2 demand. Absolute MAOD values were reduced by 9% to correct for reductions in the oxygen stores of the body (Medbo et al. 1988). The CV for supramaximal TTE and MAOD calculations in our lab is 4.0% and 7.1%, respectively ($n = 7$; 4 men; measurements taken one week apart).

Carnosine quantification via ^1H -MRS

Muscle carnosine content was measured by ^1H -MRS in the gastrocnemius medialis muscle of each subject's right limb in order to estimate muscle fibre typology. We chose to estimate the muscle fiber typology of the gastrocnemius medialis because; i) we can measure carnosine reliably in this muscle, ii) carnosine content in the gastrocnemius medialis muscle has been positively correlated with the percentage area occupied by type II muscle fibers (Baguet et al. 2011), and; iii) the gastrocnemius medialis is an extremely active muscle during running, whereby the mean and maximal electromyographic activities relative to a maximal voluntary contraction is the highest compared to other prominent lower limb muscles (Tsuji et al. 2015). ^1H -MRS measurements were performed on a 3-T whole body MRI scanner (Philips Medical Systems Best, The Netherlands). Subjects were lying in a supine position, while their lower leg was fixed in a spherical knee-coil. All the spectra were acquired using single voxel point-resolved spectroscopy (PRESS) with the following parameters; repetition time of 2000 ms, echo time of ~40 ms, number of excitations was 128 (carnosine) and 16 (water), spectral bandwidth was 2048 Hz, and an acquisition time of 4 min 16 s (carnosine) and 32 s (water). The voxel size was 40 mm x 15 mm x 20 mm. Spectral data analysis was carried out using jMRUI (version 6.0) with carnosine peaks fitted and expressed relative to the internal water signal.

Carnosine content (mM) was calculated using following formula:

$$C_m = \frac{(CS)}{(H_2O_s)} \cdot \frac{(H_2O_{T1r})}{(CT_{1r})} \cdot \frac{(H_2O_{T2r})}{(CT_{2r})} \cdot H_2O_{\text{muscle}} \cdot H_2O_{\text{protons}} \quad [2]$$

where C_m is the carnosine concentration, C_s is the carnosine signal, H_2O_s is the water signal, C_{T1r} , C_{T2r} , H_2O_{T1r} , H_2O_{T2r} are the relaxation correction factors for carnosine (earlier described by Baguet et al. (Baguet et al. 2011)) and water (earlier described by MacMillan et al. (MacMillan et al. 2016)), H_2O_{muscle} is the concentration of water in muscle, which was deducted from the molar concentration of water (55,000 mM) and the approximate water content of skeletal muscle tissue (0.7 L/kg wet weight of tissue) and H_2O_{proton} is the number of protons in water. The CV for test-retest inter-day carnosine measurements in our laboratory is 4.3% (gastrocnemius; $n = 15$ subjects). In the present study, the carnosine concentration was converted to a Z-score based on the normal distribution of our reference population. The absolute value of Z-score represents the distance between the individual value of a participant and the reference population mean value in units of the standard deviation. The reference sample population consisted of 40 recreationally active male and 33 female subjects who were scanned in our laboratory in the past 12 months.

Statistical analysis

Results are expressed as mean $\pm$ SD. Multiple stepwise regression was performed to identify the physiological and performance characteristics for which the majority of the variance in the overall time and last lap time of 1500_{FAST} could be attributed to as well as the last lap time and improvement in last lap time in 1500_{MOD} and 1500_{SLOW}, relative to 1500_{FAST}. Coefficients of determination were identified for the physiological and performance characteristics using univariate analyses and were included in the regression model if a significant r^2 value ($p < 0.05$) was reported. The magnitude of correlations was classified as small: 0.10 - 0.29, moderate: 0.30 - 0.49, large: 0.50 - 0.69, very large: 0.70 - 0.89 and extremely large: 0.90 – 1.00. A two-way analysis of variance with Tukey's post-hoc tests were conducted to compare segment speed times (i.e., 0 – 400, 400 – 800, 800 – 1100 and 1100 – 1500 m) across trials (i.e., 1500_{FAST}, 1500_{MOD} and 1500_{SLOW}).

RESULTS

1500 m trials

Table 1 displays the segment split times for each 1500 m trial. Given the manipulation in pacing, the 0 – 400 m and 400 – 800 m split times were faster in 1500_{FAST} compared to 1500_{MOD} and 1500_{MSLOW}, while the 800 – 1100

m split was similar. However, the last lap was significantly faster in both 1500_{MOD} and 1500_{MSLOW} compared to 1500_{FAST} .

1500_{FAST}

The mean speed of 1500_{FAST} relative to vVO_{2peak} and CS was $101.2\% \pm 3.2\%$ and $120.1\% \pm 7.0$, respectively. Multiple stepwise regression analysis revealed that vVO_{2peak} , RE and the speed that the 0 – 1100 m was completed relative to vVO_{2peak} (adjusted $r^2 = 0.72$, $P < 0.0001$) explained the most variation in 1500_{FAST} performance time in males. In female runners, vVO_{2peak} and RE best explained the variance in 1500_{FAST} performance time (adjusted $r^2 = 0.80$, $P < 0.001$). The ASR was associated with slower 1500_{FAST} performance times in male runners ($r^2 = 0.25$, $P = 0.02$) and female runners ($r^2 = 0.89$, $P < 0.001$), while MSS was also associated with slower times in females ($r^2 = 0.76$, $P < 0.001$). In male runners, the variance in 1500_{FAST} last lap time was best explained by a two-parameter model including vVO_{2peak} and RE (adjusted $r^2 = 0.63$, $P < 0.0001$). In female runners, vVO_{2peak} and RE also provided the best model (adjusted $r^2 = 0.53$, $P = 0.02$), while ASR was associated with slower times.

1500_{MOD}

The mean speed of the 0 – 1100 m of 1500_{MOD} relative to vVO_{2peak} and CS was $97.8\% \pm 3.5\%$ and $116.1\% \pm 7.0$, respectively. Multiple stepwise regression analysis for the determinants of last lap time in 1500_{MOD} in male runners revealed that vVO_{2peak} , MAOD and gastrocnemius carnosine Z-score provided the best model (adjusted $r^2 = 0.56$, $P < 0.0001$; Figure 2 A-D), while in female runners vVO_{2peak} and MAOD were associated with the best model (adjusted $r^2 = 0.75$, $P < 0.001$), but gastrocnemius carnosine Z-score did not further improve the model ($P = 0.184$). The ASR and the mean speed of the 0 – 1100 m relative to vVO_{2peak} (both $P > 0.05$) were associated with a slower 1500_{MOD} last lap in female runners. When the improvement in 1500_{MOD} last lap time was expressed relative to 1500_{FAST} last lap time, gastrocnemius carnosine Z-score and MAOD provided the best model in male runners (adjusted $r^2 = 0.60$, $P = 0.017$), whereas in female runners gastrocnemius carnosine Z-score and supramaximal TTE provided the best model (adjusted $r^2 = 0.66$, $P = 0.026$).

1500_{SLOW}

The mean speed of the 0 – 1100 m of 1500_{SLOW} mean speed relative to vVO_{2peak} and CS was $92.7\% \pm 3.3\%$ and $110.1\% \pm 6.7$, respectively. Multiple stepwise regression analysis for the determinants of last lap time in 1500_{SLOW} in male runners revealed that vVO_{2peak} , gastrocnemius carnosine Z-score and MAOD provided the best model (adjusted $r^2 = 0.63$, $P < 0.001$; Figure 3 A-D) which was also similar in female runners (adjusted $r^2 = 0.80$, $P <$

0.0001). When the improvement in 1500_{SLOW} last lap time was expressed relative to 1500_{FAST} last lap time, gastrocnemius carnosine Z-score and MAOD provided the best model in male runners (adjusted $r^2 = 0.76$, $P = 0.017$), whereas in female runners gastrocnemius carnosine Z-score and MAOD provided the best model (adjusted $r^2 = 0.81$, $P < 0.001$).

Muscle fibre typology

The middle-distance runners included in the present study predominantly had negative gastrocnemius carnosine Z-score values (28/32 runners; figure 4). Figure 5 shows the association between gastrocnemius carnosine Z-score and RE at 110% GET in both male ($r^2 = 0.39$, $P < 0.01$) and female runners ($r^2 = 0.46$, $P = 0.02$). This association was comparable for all running speeds that were tested (85 - 110% GET), whereby r^2 values ranged from 0.35 – 0.47). All runners had a blood lactate concentration of ≤ 2.1 mM suggesting that energy contribution of anaerobic metabolism was minimal. Gastrocnemius carnosine Z-score was also associated with MSS in both male ($r^2 = 0.23$, $P = 0.02$) and female runners ($r^2 = 0.35$, $P = 0.05$), CS in male runners ($r^2 = 0.24$, $P = 0.02$), as well as supramaximal TTE ($r^2 = 0.22$, $P < 0.01$; male runners and female runners combined). There was no significant association between gastrocnemius carnosine Z-score and vVO_{2peak} .

DISCUSSION

Novel findings from the present study show that the improvements in last lap time in the paced trials were largely explained by MAOD and muscle fibre typology in both male and female runners. That is, runners with a higher gastrocnemius carnosine Z-score (i.e., higher estimated percentage of type II fibres) and greater MAOD, reduced their last lap time to a greater extent in the paced 1500 m trials. Furthermore, a lower carnosine Z-score (i.e., estimated higher percentage of type I fibres) was associated with superior RE (lower energy cost). We also confirm that RE and vVO_{2peak} largely explain the variation in maximal 1500 m time trial time, while vVO_{2peak} is also a key determinant of last lap time in the paced trials. These findings suggest that middle-distance runners who possess diverse physiological profiles may be best suited to adopting polarized racing strategies in order to have the best chance of winning. The extent to which the diversity in the athlete profiles relate to differences in training characteristics and subsequent adaptations or inherent differences in genetic predisposition requires further examination.

RE and vVO_{2peak} were the most deterministic characteristics of 1500_{FAST} running performance which agrees with previous studies (Ingham et al. 2008; Sandford et al. 2019c). The mean speed of the 1500_{FAST} trial was $\sim 101\%$ of vVO_{2peak} , which is slightly lower than previous reports ($\sim 105 - 115\%$ vVO_{2peak}) (Billat et al. 2009; Sandford et al.

2019c; Billat 2001), but this can be explained by the faster times achieved in competition racing against competitors (Billat et al. 2009; Sandford et al. 2019c; Billat 2001) compared to completing individual time trials (i.e., 1500_{FAST}), as well as differences in the methodology used to determine vVO_{2peak} . We observed that a greater ASR was associated with slower 1500_{FAST} performance times. These findings also agree with previous research (Sandford et al. 2019c), which was conducted in senior and junior national and international level middle-distance runners, whereby the magnitude of ASR was associated with slower 1500 m running velocity ($r = -0.68 \pm 0.54$) (Sandford et al. 2019c). Given that vVO_{2peak} was a key determinant of 1500 m performance time in the present study and in previous research (Sandford et al. 2019c), it is likely that the association between ASR and slower 1500 m performance times manifests from a lower vVO_{2peak} , rather than greater MSS. For example, if one athlete has a larger ASR compared to another athlete, but both are matched for MSS, the athlete with the larger ASR would simply have a lower vVO_{2peak} without possessing superior mechanical speed capabilities.

In the present study, vVO_{2peak} was also a key determinant of the last lap time in the paced trials (i.e., 1500_{MOD} and 1500_{SLOW}), in addition to MAOD. These findings are consistent with the supposition that 1500 m racing is likely to elicit a maximum rate of aerobic metabolism and exhaust the finite capacity of the anaerobic energy systems (Billat et al. 2009; Ward-Smith 1999). Apart from 1500_{MOD} in female runners, gastrocnemius carnosine Z-score and MAOD significantly contributed to the regression models explaining some of the variation associated with last lap time and the improvement in last lap time in the paced trials. That is, runners with a higher carnosine Z-score (i.e., higher estimated percentage of type II fibres) and greater MAOD, reduced their last lap time to a greater extent in the paced 1500 m trials. This indicates an advantage for 1500 m runners who possesses a higher proportion of type II fibres, given that these fibres not only underpin speed and power performance (Tesch et al. 1989; Pette 1985; Harber et al. 2004), but also have the ability to adapt to high oxidative demands which are not necessarily subservient to type I fibres in well-trained endurance athletes (Jansson and Kaijser 1977; Ørtenblad et al. 2018). For example, succinate dehydrogenase activity levels have been shown to be similar between type I and II fibres in the gastrocnemius muscle of elite cross-country runners (Jansson and Kaijser 1977), while mitochondrial volume, average capillary-to-fibre ratio and maximal citrate synthase activity is comparable in the highly-trained type I and II fibres of elite cross-country skiers (Ørtenblad et al. 2018). In addition to a higher carnosine Z-score, a larger anaerobic capacity (i.e., MAOD) also significantly contributed to the best regression model explaining the variance in absolute last lap time and the relative improvement in the last lap time in the paced trials. The significant contribution of MAOD may have arisen due to the slower relative speed of the 0 – 1100 m in 1500_{MOD} and 1500_{SLOW} (~98% and ~93% of vVO_{2peak} and 116% and 110% of CS, respectively),

compared to 1500_{FAST} ($\sim 103\%$ vVO_{2peak} and 121% CS). Given the slower running speeds relative to vVO_{2peak} and CS, it could be expected that the capacity of anaerobic energy provision, as reflected by the remaining portion of the MAOD, could be preserved to enhance last lap running speed. Previous research in elite middle-distance runners has shown that at any point within a 1500 m time trial, running speed is controlled by the remaining anaerobic energy store (defined as the difference between end-race and instantaneous oxygen deficits) (Billat et al. 2009). Thus, in a group of athletes with a relatively homogenous vVO_{2peak} and CS, preservation of MAOD during the initial laps of a 1500 m race is likely to result in a greater last lap performance improvement for athletes with a larger MAOD.

It has been suggested that a greater ASR may be an important determinant of a fast last lap in a tactical, slow 1500 m race (Sandford et al. 2019b; Mytton et al. 2015). This supposition may be based on the premise that if the speed of the initial laps of a 1500 m race are slower, an athlete with a larger ASR may be able to utilize their superior speed reserve in order to run a faster last lap than their competitors (Sandford et al. 2019b; Mytton et al. 2015). Despite this, we found little evidence that ASR or MSS were associated with the last lap time in paced trials or the relative improvement in last lap time relative to 1500_{FAST} given that these values did not significantly contribute to the regression models explaining the variation in performance in the paced trials. While MSS has previously been associated with 800 m time trial performance (Bachero-Mena et al. 2017), it is unlikely that athletes would reach running speeds during 1500 m races that are close to their MSS, thus reducing the importance of MSS for 1500 m running performance. For example, the fastest mean 100 m sector for male 800 m championship medallists is $\sim 8.9 \text{ m}\cdot\text{s}^{-1}$, whereas 1500 m championship medallists only reach mean 100 m sector speeds of $\sim 7.0 \text{ m}\cdot\text{s}^{-1}$ (Hanley et al. 2019). In relation to ASR, it is unlikely that this construct represents an anaerobic energy reserve given that its upper boundary (i.e., MSS) is more likely to be limited from a mechanical standpoint rather than an energetics perspective (Bundle and Weyand 2012). The lower boundary of the ASR, demarcated by vVO_{2peak} , is thought to represent the minimal intensity that would elicit VO_{2peak} (Morgan et al. 1989) and certainly does not represent a threshold above which energy metabolism is solely accounted for by anaerobic metabolism. As such, the magnitude of ASR does not necessarily represent a surrogate measure of anaerobic capacity which has been previously proposed (Bundle et al. 2003). Indeed, we did not find a strong association between MAOD and ASR in the present study. It is plausible that the importance of ASR for the last lap speed in slower paced 1500 m trials may be related to the running speed relative to vVO_{2peak} (or CS) during the initial laps (i.e., 0 – 1100 m), but we found no such association. A classical study (Blondel et al. 2001) showed that time to exhaustion at 120% and 140% of vVO_{2peak} and the velocity expressed relative to the ASR had a high

correlation ($r = -0.83$, $P < 0.01$ and $r = -0.94$, both $P < 0.001$), suggesting that ASR may be a meaningful parameter when the intensity of exercise is supramaximal. However, the trials conducted in this study (Blondel et al. 2001) are in stark contrast to the 1500 m trials of the present study, whereby the supramaximal last lap speed is preceded by relatively high energetic requirements of the 0 – 1100 m. As such, in the context of our study, we did not find any substantial evidence that the ASR substantially contributes to last lap speed, or the improvement in last lap speed, in paced 1500 m trials in a group of trained 1500 m runners. It is likely that ASR may be a more important determinant of 800 m running performance, whereby the duration (122 ± 27 s and 65 ± 17 s, respectively) and running intensity (120% and 140% of $v\text{VO}_{2\text{peak}}$) of the supramaximal trials featured in Blondel et al. (2001) more closely reflect 800 m running duration and intensity (Billat et al. 2009; Sandford et al. 2019c; Billat 2001). This would also be in accordance with previous research showing a strong relationship between 800 m performance times and ASR in a group of ten elite 800 m runners (800 m personal best $\leq 1:47.50$) (Sandford et al. 2018).

Contrary to how D' may theoretically contribute to last lap performance during paced trials (Pettitt 2016), we did not find that D' was associated with the relative improvement or absolute last lap time during the paced trials. Furthermore, we did not find a strong association between MAOD and D' . This may seem surprising given that some investigators consider the magnitude of D' (Gaesser and Poole 1996; Hill and Smith 1993) to be synonymous with anaerobic capacity. However, there is still a great deal of uncertainty regarding the physiological determinants of D' (Vanhatalo et al. 2010). It has recently been demonstrated that W' during knee extension exercise, which is synonymous with D' during running, is reduced with the inspiration of hyperoxic gas (Vanhatalo et al. 2010), whereby the reduction in W' was inversely associated with the increase in critical power in hyperoxic conditions. These findings (Vanhatalo et al. 2010) suggest that W'/D' may not simply reflect an anaerobic energy store given that a particular measure of anaerobic capacity should be O_2 independent and insensitive to interventions which alter muscle O_2 delivery. Other research shows that the magnitude of the VO_2 slow component (Murgatroyd et al. 2011) and mental fatigue (Salam et al. 2018) may influence W' , indicating that the mechanistic basis of W'/D' is quite complex, and the compatibility of this parameter as a measure of anaerobic capacity seems unreasonable. Nonetheless, in the context of the present study D' did not relate to any aspect of performance in maximal or paced 1500 m trials.

An interesting finding from the present study was that gastrocnemius carnosine Z-score was positively correlated with RE, whereby subjects with a lower gastrocnemius carnosine Z-score (higher estimated proportion of type I fibres) were more economical (i.e., have a lower energetic cost of running) while running at speeds close to the GET (85 – 110%). This finding agrees with some (Bosco et al. 1987), but not all research (Kyrolainen et al. 2003)

that has investigated the relationship between muscle fibre typology and RE in highly-trained runners. However, the purported inverse relationship between the proportion of type II fibres and RE (Kyrolainen et al. 2003) likely manifested from the slower VO_2 on-kinetics associated with type II fibres (Pringle et al. 2003) given that the running speed ($\sim 25.2 \text{ km}\cdot\text{h}^{-1}$) and duration (1 min) employed were not suitable to achieve a steady state which is a necessary requirement for the valid measurement of RE. In contrast, we assessed subjects running at a relative running speed equivalent to 110% GET ($\sim 16.8 \text{ km}\cdot\text{h}^{-1}$), whereby all blood lactate concentration values were $\leq 2.0 \text{ mmol}\cdot\text{L}^{-1}$. While the relevance of this submaximal speed to 1500 m race pace could be questioned, the validity of RE measurements is dependent upon steady state submaximal running speeds. The superior RE in subjects with slow-twitch muscle typology in the present study could possibly be related to the higher muscular efficiency (i.e., ATP formation per oxygen used) of type I fibres, but is more likely related to the greater contractile efficiency (i.e., work per ATP usage) of type I fibres (Bottinelli and Reggiani 2000).

The aim of the present study was to provide more insight to the underpinning physiological and performance characteristics that dictate last lap speed in tactical 1500 m races. It should be acknowledged that all 1500 m races are somewhat unique, and it is impossible to truly replicate tactical 1500 m scenarios. In the present study, the 0 – 1100 m in 1500_{MOD} and 1500_{SLOW} were paced relative to 1500_{FAST} and did not include additional mid race surges that seem to be a characteristic of tactical 1500 m races (Sandford et al. 2019a). Furthermore, other factors such as drafting, re-positioning and running wide on bends or close to the rail would also influence the physiological requirements of a given race (Sandford et al. 2019a; Jones and Whipp 2002), and subsequently, last lap speed. Lastly, the results of the present study should be contextualised in light of the performance level of the subjects that participated. As is often the case in experimental studies, we recruited trained, but not elite 1500 m runners. Whether similar findings would be present in a group of elite 1500 m runners required further evaluation.

Perspective

The findings from the present study suggest that the diversity in the physiological profile of middle-distance runners could inform a preference for different racing strategies. While the majority of this diversity may be due to training characteristics and subsequent training adaptations, we highlight that muscle fibre typology explains a large degree of the variation in RE which is a key determinant of maximal 1500 m performance, but also underpins last lap speed in tactical 1500 m racing. Importantly, muscle fibre typology is largely dependent on genetic predisposition and is not thought to change substantially in response to training (Ingalls 2004). As such, there is a complex interplay between how much training induced adaptations and genetic predisposition contributes to the

diversity in the physiological profiles of middle-distance runners. Coaches should continue to maximise aerobic characteristics (i.e., RE and $\dot{V}O_{2\text{peak}}$) through training as these were important determinants of both maximal and tactical 1500 m racing. Once a given level of aerobic capability has been acquired, other characteristics such as anaerobic capacity and higher proportions of type II muscle fibres may be important for last lap speed, particularly in slower paced races. As such, developing anaerobic capacity and speed endurance should not be overlooked. While athletes with lower gastrocnemius carnosine Z-scores (i.e., higher estimated proportion of type I fibres) have the advantage of better RE and CS (at least in male runners), athletes with higher gastrocnemius carnosine Z-scores (i.e., higher estimated proportion of type II fibres) may possess the benefits of superior speed endurance and greater capacity for anaerobic energy provision. It could be suggested that runners with a lower gastrocnemius carnosine Z-score may be best suited to a racing strategy characterized by a sustained pace from the start in order to take advantage of superior RE, while runners with a higher gastrocnemius carnosine Z-score may be best suited to controlling the pace of the race in the initial laps, in order to take advantage of the superior last lap speed.

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TABLES

Table 1 – The mean (SD) lap split times for the maximal (1500_{FAST}), moderate (1500_{MOD}) and slow paced (1500_{SLOW}) 1500 m time trials.

^aindicates that 1500_{FAST} was significantly faster than 1500_{MOD} and 1500_{SLOW}

^bindicates that 1500_{MOD} was significantly faster than 1500_{SLOW}

^cindicates that 1500_{MOD} was significantly faster than 1500_{SLOW}

^dindicates that 1500_{SLOW} was significantly faster than 1500_{MOD} and 1500_{FAST}

Table 2 – The mean (range) physiological and performance characteristics of the female and male runners.

HR; heart rate, CS; critical speed, D'; curvature constant, TTE; time to exhaustion, MAOD; maximal accumulated oxygen deficit, ASR; anaerobic speed reserve, MSS; maximal sprint speed, RE; running economy

FIGURES

Figure 1 - Determinants of maximal 1500 m time trial performance time (1500_{FAST} ; panel A: male, panel B: female) and 1500_{FAST} last lap time (panel C: male, panel D: female). Bars indicate the magnitude of explained variance ($\% r^2$) obtained from Pearson's correlations. Black bars indicate determinants associated with faster times, while grey bars indicate variables that are associated with slower times. Closed bars indicate a significance ($P < 0.05$). Multiple stepwise regression analysis revealed a combination of predictors, which are presented at the top of each panel.

Figure 2 - Determinants of last lap time in the moderately paced trial (1500_{MOD} ; panel A: male, panel B: female) and the improvement in 1500_{MOD} lap time relative to 1500_{FAST} (panel C: male, panel D: female). Bars indicate the magnitude of explained variance ($\% r^2$) obtained from Pearson's correlations. Black bars indicate determinants associated with faster times, while grey bars indicate variables that are associated with slower times. Closed bars indicate a significance ($P < 0.05$). Multiple stepwise regression analysis revealed a combination of predictors, which are presented at the top of each panel.

Figure 3 - Determinants of last lap time in the slow-paced trial (1500_{SLOW} ; panel A: male, panel B: female) and the improvement in 1500_{SLOW} lap time relative to 1500_{FAST} (panel C: male, panel D: female). Bars indicate the magnitude of explained variance ($\% r^2$) obtained from Pearson's correlations. Black bars indicate determinants associated with faster times, while grey bars indicate variables that are associated with slower times. Closed bars indicate a significance ($P < 0.05$). Multiple stepwise regression analysis revealed a combination of predictors, which are presented at the top of each panel.

Figure 4 – Individual and mean (95%CI) carnosine Z-score values of the gastrocnemius of runners in the present study, as well as the non-athlete control population. The absolute carnosine concentration for the runners was converted to a sex- specific Z-score relative to an age-matched control population of active, healthy non-athletes (males: $n = 40$; females: $n = 33$).

Figure 5 – Relationship between muscle fibre typology (carnosine Z-score) and the energy cost of running at 110% of the gas exchange threshold in male (A) and female (B) subjects.