

Physiological and training characteristics of recreational marathon runners

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Purpose: The aim of this study was to examine the physical and training characteristics of recreational marathon runners within finish time bandings (2.5–3 h, 3–3.5 h, 3.5–4 h, 4–4.5 h and >4.5 h).

Materials and methods: A total of 97 recreational marathon runners (age 42.4 ± 9.9 years; mass 69.2 ± 11.3 kg; stature 172.8 ± 9.1 cm), with a marathon finish time of 229.1 ± 48.7 min, of whom $n = 34$ were female and $n = 63$ were male, completed an incremental treadmill test for the determination of lactate threshold (LT1), lactate turn point (LT2) and running economy (RE). Following a 7-min recovery, they completed a test to volitional exhaustion starting at LT2 for the assessment of $\dot{V}O_{2max}$. In addition, all participants completed a questionnaire gathering information on their current training regimes exploring weekly distances, training frequencies, types of sessions, longest run in a week, with estimations of training speed, and load and volume derived from these data.

Results: Training frequency was shown to be significantly greater for the 2.5–3 h group compared to the 3.5–4 h runners ($P < 0.001$) and >4.5 h group ($P = 0.004$), while distance per session ($\text{km} \cdot \text{session}^{-1}$) was significantly greater for the 2.5–3 h group (16.1 ± 4.2) compared to the 3.5–4 h group (15.5 ± 5.2 ; $P = 0.01$) and >4.5 h group (10.3 ± 2.6 ; $P = 0.001$). Race speed correlated with LT1 ($r = 0.791$), LT2 ($r = 0.721$) and distance per session ($r = 0.563$).

Conclusion: The data highlight profound differences for key components of marathon running ($\dot{V}O_{2max}$, LT1, LT2, RE and $\% \dot{V}O_{2max}$) within a group of recreational runners with the discriminating training variables being training frequency and the absolute training speed.

Keywords: endurance running, nonelite, workout structures, maximal oxygen uptake, running economy, aerobic capacity

Introduction

Marathon running is one of the largest mass participation sports offering opportunities in big city races for elite, nonelite and recreational runners to pit themselves over the 42.195 km distance. Concerning marathon performance, it is well recognized that running speed is regulated through aerobic metabolic pathways in the engaged muscle mass and economic conversion of the derived energy to muscle actions.¹ Indeed, the ability to sustain race speed across the marathon is dependent on running economy (RE) reflecting the O_2 cost of running at submaximal speeds,^{2,3} maximal oxygen uptake ($\dot{V}O_{2max}$),^{2,4} fractional utilization of $\dot{V}O_{2max}$,^{2,5,6} the size of the aerobic capacity as reflected by the submaximal blood lactate response to exercise and the speed associated with lactate threshold (LT1) and lactate turn point (LT2).^{2,5,7}

$\dot{V}O_{2\max}$ is an oft-cited variable in relation to the marathon; the relevance is exemplified by the notion that $\dot{V}O_{2\max}$ represents the integration of the cardiovascular, respiratory and muscular systems to utilize O_2 and is reflected through the Fick principle, where $\dot{V}O_{2\max} = \text{maximal cardiac output } (\dot{Q}_{\max}) \cdot \text{maximal arteriovenous oxygen difference } (a-vO_{2\text{difmax}})$. Typical $\dot{V}O_{2\max}$ values for elite male and female runners have been reported in the order of 67–85 mL·kg⁻¹·min⁻¹ with runners referred to as “good” (finishing times of 150–180 min) exhibiting a value of 65.5 ± 1.2 mL·kg⁻¹·min⁻¹, while those classified as “slow runners,” that is finishing time >180 min, showing a $\dot{V}O_{2\max}$ of 58.7 ± 1.9 mL·kg⁻¹·min⁻¹.^{1,2,6,7} Of significance to the marathon runner is the fractional utilization of $\dot{V}O_{2\max}$ (% $\dot{V}O_{2\max}$) that can be sustained as reflected by the manifestation of the LT2 response, reflecting the inability of fatty acid metabolism to sustain oxidative phosphorylation to meet the requisite exercise intensity.⁵ Indeed, it has been reported that, in elite marathon runners, this point occurs between 85% and 90% $\dot{V}O_{2\max}$,^{2,6,7} while for “slower” runners with finishing time >180 min LT2 has also been reported to occur at 85% $\dot{V}O_{2\max}$.⁶ Associated with this point is the fractional utilization at LT1 representing the balance between lactate efflux from the muscle and disappearance from the blood^{2,8} characteristically occurring at 50–80% $\dot{V}O_{2\max}$, although in highly trained marathon runners (<150 min for the marathon) it has been shown to occur at 65–80% $\dot{V}O_{2\max}$. The submaximal cost of running (RE) reflects the $\dot{V}O_2$ for any given speed and has previously been shown to account for significant variations in distance running performance among athletes with a similar $\dot{V}O_{2\max}$,^{5,9} with RE being shown to be influenced by a myriad of factors including $\dot{V}O_{2\max}$, training volume and training history.⁹

There is little doubt that the nature and accumulation of training play a pivotal role in developing the underlying physiological characteristics and thus race speed. Indeed, across a series of works,^{6,7,10,11} training compositions of marathon runners have been cataloged highlighting typical distances covered per week of 50.5 ± 9.1 km for runners with a marathon time ~204 min, coupled with an average training speed of 11.0 ± 1.4 km·h⁻¹. Furthermore, in the elite runner with marathon times of 129 ± 2 min and 149 ± 3 min for males and females, respectively, training histories showed weekly distances of 206 ± 26 km and 166 ± 11 km coupled with a training frequency of 13.0 ± 0.7 session·week⁻¹ and 12.2 ± 0.4 session·week⁻¹. However, when reflecting on average finish times in mass participation marathons, the data reported for both training and underlying system physiology seem at odds, with a recent study.⁴ This study showcased in

a sample of 91,929 that the greatest proportion of male runners (66.6%) had a finish time of >240 min and >270 min (51.1%) for females, with data from the 2015 edition of the London marathon, excluding the registered elite runners who showed a finish time of 262 ± 53 min ranging from 138 min to 459 min. Therefore, given the apparent disparities between reported training and physiological characteristics of marathon runners and typical finish times for the majority of runners, this study explores the physical and training-orientated characteristics of nonelite marathon runners with an average finish time of ~3.5 h.

Materials and methods

Following local institutional ethics approval (Faculty Research and Ethics Panel, Anglia Ruskin University) and having provided written informed consent, n = 97 marathon runners volunteered and agreed to participate (age 42.4 ± 9.9 years; mass 69.2 ± 11.3 kg; stature 172.8 ± 9.1 cm; body mass index [BMI] 20.2 ± 2.5 kg·m⁻²), with a marathon finish time of 229.1 ± 48.7 min, of whom n = 34 were female and n = 63 were male. Participants were recruited through an online UK-based running website and word of mouth, with the primary inclusion criteria being that they must be completing an International Athletics Federation (IAAF) or UK Athletics (UKA) sanctioned marathon between March and May 2016. All laboratory testing was completed at least 8 weeks prior to the subsequent spring marathon, and all training data were collected at this same time point.

Submaximal treadmill test

For the determination of LT1, LT2 and RE, each participant completed an incremental test, where running speed was increased 1 kph·3 min⁻¹ until LT2 was reached, upon which the test was terminated; throughout all stages the treadmill gradient was held at 1%.¹² During all trials, gas exchange responses were ascertained on a breath-by-breath basis via a pre-calibrated metabolic cart (Metalyzer 3B; Cortex, Leipzig, Germany). Upon the completion of each 3-min stage, the participant stood astride of the treadmill to facilitate the collection of capillary blood sample (20 µL) for the determination of blood lactate. Each stage was separated by 1-min recovery. The initial running speed was selected to coincide with that which the athlete's normally warm-up at, so as to enable them to ease into the protocol.

For each participant, the blood lactate responses (mM) were plotted against exercise intensity (km·h⁻¹), with LT2 being determined through a visual inspection of the curve and validated independently by two physiologists. Quantification

of LT1 was based on the criteria of the first initial rise beyond baseline, and again this was verified by two physiologists.

$\dot{V}O_{2max}$

Following a 7-min active recovery, upon completion of the submaximal treadmill component, treadmill speed was adjusted to that which coincided with LT2, with speed remaining constant and gradient increasing by 1%·min⁻¹ until volitional exhaustion, or when the participant could not maintain a predetermined position at the front of the treadmill. Again, expired air was recorded on a breath-by-breath basis, and cardiovascular responses were determined. Upon completion of the test, a capillary blood sample (20 μ L) was attained for immediate determination of postexercise blood lactate and glucose concentration. $\dot{V}O_{2max}$ was confirmed according to previously established criteria.

Pulmonary gas exchange responses

Using a low-resistance mouthpiece and turbine, assembly volumes and flow rate were determined. For the determination of expired gas concentration, O₂ and CO₂ were analyzed at a rate of 60 mL·min⁻¹ while being drawn off directly from the mouthpiece. Using custom metabolic cart software, the gas concentrations and respiratory responses were aligned to reflect breath-by-breath gas exchange variables ($\dot{V}O_2$, $\dot{V}CO_2$ [$\dot{V}CO_2$ = volume of carbon dioxide], minute ventilation [VE] and respiratory exchange ratio [RER]). Prior to all trials, the metabolic cart was calibrated for both volume/flow and gas concentration according to the manufacturer's specifications.

Cardiovascular responses

During both the submaximal stages and the $\dot{V}O_{2max}$ trial, heart rate (HR) responses were recorded with a 5 s sampling frequency using a Polar 810s telemetric system (Polar, Kempele, Finland).

Blood chemistry

Prior to the commencement of all trials, baseline capillary blood samples (150 μ L) were collected for the automated analysis of key hematological and biochemical markers (Opti CCA-TS; Una Health, Cardiff, UK). A resting blood lactate/glucose sample (20 μ L) was also recorded (Biosen C; EKF, Stoke on Trent, UK). All equipment was calibrated as per the manufacturer's instructions.

Training characteristics and history

All participants completed the training history questionnaire post laboratory testing. The questionnaire was designed in collaboration with physiologists, psychologists and running

coaches as well as taking into account work that had been conducted previously in this field. The questionnaire included questions pertaining to the athletes' age, racing experience, predicted finish time for the marathon, race number, use of pacing devices and personal best times across different race distances. Questions regarding training focused on the number of sessions per week (defined as the typical training week), days training per week, weekly distance covered and longest run in a week and long runs per week (>10 km), with weekly distance defined as the typical distance completed in the preparation for the marathon. From these data, the following computations were possible: average training speed (km·h⁻¹), average training duration per session (h·session⁻¹), training volume and training load (AU).

Statistical analyses

Analysis of the data was completed using Statistical Package for Social Sciences (SPSS, v.21; IBM Corporation, Armonk, NY, USA) for Windows and Graphpad Prism v.7 (GraphPad Software, Inc., La Jolla, CA, USA). All data are expressed as mean \pm SD. Data were screened for normality of distribution and homogeneity of variance through a Shapiro–Wilk normality test. One-way analysis of variance (ANOVA) was performed to compare the physical and training characteristics between each of the groups, while post hoc pairwise comparisons were made using Tukey's adjustment. Additional analysis of association between training and physical characteristics was made using a Pearson product-moment correlation. Statistical significance was set at $P < 0.05$.

Results

Group characteristics

Of the original $n = 97$ athletes, only 82 completed a spring marathon; thus, all data are presented on these $n = 82$ runners. The runners were subdivided, based on their performance in a sanctioned spring marathon, into five groups of which the basic anthropometric and physiological data are presented in Table 1, and the training characteristics in Table 2. Those in the >4.5 h group (274.7–409.4 min) had a mean completion time of 305.0 \pm 39.2 min ($n = 17$) of whom $n = 12$ were female and $n = 5$ were male. The 4–4.5 h group had a finish time of 253.9 \pm 9.1 min ($n = 7$) with $n = 2$ females and $n = 5$ males (240.6–263.4 min), while the 3.5–4 h group had a marathon completion time of 225.3 \pm 9.2 min (210.4–239.0; $n = 24$) with $n = 9$ females and $n = 15$ males. The 3–3.5 h group ($n = 23$), $n = 3$ females and $n = 12$ males, exhibited a mean completion time of 197.6 \pm 6.9 min (186.5–209.5 min). The fastest group of runners (2.5–3 h; $n = 11$) had a completion time of 170.6 \pm 7.0 min (158.9–179.8 min) of whom there was $n = 1$ female and $n = 10$ males.

Table 1 Physiological characteristics of the n = 97 marathon runners

Characteristics	2.5–3 h	3–3.5 h	3.5–4 h	4–4.5 h	>4.5 h
Age (years)	40.0 ± 7.3	43.6 ± 9.9	42.4 ± 11.6	43.6 ± 9.3	42.1 ± 10.3
Mass (kg)	65.6 ± 7.4	71.2 ± 9.0	67.8 ± 11.0	72.4 ± 17.5	69.6 ± 13.7
Height (cm)	174.9 ± 8.1	175.6 ± 7.8	171.9 ± 8.0	171.9 ± 10.8	169.1 ± 11.3
BMI (kg·m ⁻²)	18.7 ± 1.5	20.2 ± 2.1	19.6 ± 2.5	20.9 ± 4.1	20.4 ± 2.8
$\dot{V}O_{2max}$ (mL·kg ⁻¹ ·min ⁻¹)	63.3 ± 7.7	55.7 ± 4.8 ^a	53.2 ± 4.6 ^a	53.0 ± 8.6 ^a	46.5 ± 5.2 ^a
LT1 (km·h ⁻¹)	12.4 ± 0.7	11.0 ± 0.8 ^a	10.3 ± 1.1 ^b	10.1 ± 1.5 ^a	8.6 ± 0.9 ^{a-d}
LT2 (km·h ⁻¹)	15.5 ± 0.7	13.8 ± 0.7 ^a	13.1 ± 1.6 ^{ab}	12.7 ± 2.1 ^{ab}	10.9 ± 1.2 ^{ab}
LT1 (mM)	1.3 ± 0.4	1.6 ± 0.5 ^a	1.5 ± 0.6 ^{ab}	1.5 ± 0.4 ^{ab}	1.9 ± 0.8 ^{ab}
LT2 (mM)	2.8 ± 0.5	2.8 ± 0.5	2.7 ± 0.6	2.5 ± 0.5	3.0 ± 0.7
LT1 (% $\dot{V}O_{2max}$)	68.7 ± 7.5	70.7 ± 6.4	70.5 ± 5.7	73.3 ± 7.5	71.6 ± 6.9
LT2 (% $\dot{V}O_{2max}$)	84.1 ± 5.2	84.4 ± 4.3	84.1 ± 4.2	85.1 ± 2.8	83.6 ± 4.6
$\dot{V}O_2$ -LT1 (mL·kg ⁻¹ ·min ⁻¹)	43.2 ± 4.2	39.4 ± 2.3	37.5 ± 4.5	35.9 ± 5.2	33.0 ± 2.3
$\dot{V}O_2$ -LT2 (mL·kg ⁻¹ ·min ⁻¹)	52.8 ± 5.6	47.0 ± 3.7	44.7 ± 4.3	44.1 ± 7.2	38.7 ± 3.4
$\dot{V}O_2$ -LT1 (mL·kg ⁻¹ ·km ⁻¹)	209.5 ± 15.2	215.4 ± 14.6	219.2 ± 20.1	214.0 ± 15.2	230.4 ± 18.3
$\dot{V}O_2$ -LT2 (mL·kg ⁻¹ ·km ⁻¹)	204.2 ± 17.3	205.2 ± 10.0	208.0 ± 14.0	204.4 ± 7.7	214.3 ± 13.7
HR1 (b·min ⁻¹)	137.5 ± 7.6	139.5 ± 14.7	141.0 ± 15.2	131.9 ± 13.7	139.0 ± 11.3
HR2 (b·min ⁻¹)	160.8 ± 8.1	159.1 ± 13.7	161.0 ± 10.6	157.0 ± 18.3	161.9 ± 14.7
HR _{max} (b·min ⁻¹)	176.4 ± 9.5	178.2 ± 13.9	176.7 ± 18.3	174.1 ± 18.0	179.3 ± 14.0
VE _{max} (l·min ⁻¹)	149.4 ± 21.4	141.6 ± 21.0 ^a	132.9 ± 30.8	129.3 ± 42.9	114.4 ± 24.4 ^{ab}
PBLa (mM)	8.9 ± 1.0	9.0 ± 1.9	9.0 ± 2.8	8.8 ± 2.1	9.1 ± 2.9

Notes: Data presented as mean ± standard deviation. LT1, lactate threshold; LT2, lactate turn point; HR1, HR at LT1; HR2, HR at LT2; PBLa, peak blood lactate concentration. ^aSignificant difference to the 2.5–3 h group. ^bSignificant difference to the 3–3.5 h group. ^cSignificant difference to the 3.5–4 h group. ^dSignificant difference to the 4–4.5 h group.

Abbreviations: BMI, body mass index; HR, heart rate; PBLa, peak blood lactate concentration; VE, minute ventilation.

Table 2 Training and racing characteristics of the n = 82 marathon runners

Training characteristics	2.5–3 h	3–3.5 h	3.5–4 h	4–4.5 h	>4.5 h
h·week ⁻¹	8.1 ± 2.5	7.9 ± 3.2	6.5 ± 2.7	7.7 ± 2.4	7.3 ± 5.4
Runs·week ⁻¹	5.7 ± 1.0	5.0 ± 1.0	4.1 ± 1.3 ^{ab}	4.9 ± 1.0 ^b	4.4 ± 1.1 ^a
km·week ⁻¹	91.7 ± 31.6	81.5 ± 26.0	62.4 ± 27.3 ^{ab}	56.2 ± 14.8 ^{a-c}	43.8 ± 9.5 ^{a-d}
h·session ⁻¹ ·*	1.5 ± 0.4	1.6 ± 0.5	1.7 ± 1.1	1.6 ± 0.2	1.7 ± 1.0
km·session ⁻¹ ·*	16.1 ± 4.2	16.4 ± 3.0	15.5 ± 5.2 ^a	12.2 ± 2.1 ^{bc}	10.3 ± 2.6 ^{a-c}
Longest run (km)	37.3 ± 5.8	31.1 ± 7.2	31.8 ± 5.5	32.2 ± 3.6	29.1 ± 8.2
Speed (km·h ⁻¹)·*	11.4 ± 2.0	11.1 ± 3.6	10.1 ± 3.8	8.1 ± 1.6 ^{ab}	8.0 ± 4.0 ^a
Volume (AU)·*	537.7 ± 266.1	429.7 ± 230.5	267.3 ± 169.9 ^b	371.1 ± 296.5	201.7 ± 87.0 ^{ab}
Load (AU)·*	34892 ± 16307	26888 ± 15360	14960 ± 7455 ^{ab}	22646 ± 18598	11909 ± 5259 ^{ab}
Years training	14.0 ± 7.6	11.0 ± 6.7	11.4 ± 11.5	12.6 ± 13.6	7.2 ± 8.4
Race speed (km·h ⁻¹)	14.9 ± 0.6	12.8 ± 0.5	11.3 ± 0.6	10.0 ± 0.4	8.4 ± 8.9

Notes: Data presented as mean ± standard deviation. *Aggregated scores: h·session⁻¹ = h·week⁻¹/sessions·week⁻¹, km·session⁻¹ = km·week⁻¹/sessions·week⁻¹; speed = km·session⁻¹/h·session⁻¹; volume = sessions·week⁻¹ × km·week⁻¹; load = % $\dot{V}O_{2max}$ × volume. ^aSignificant difference to the 2.5–3 h group. ^bSignificant difference to the 3–3.5 h group. ^cSignificant difference to the 3.5–4 h group. ^dSignificant difference to the 4–4.5 h group.

$\dot{V}O_{2max}$
Significant differences were observed for $\dot{V}O_{2max}$ between the 2.5–3 h group and 3–3.5 h group ($P = 0.004$), 3.5–4 h group ($P < 0.001$), 4–4.5 h runners ($P = 0.01$) and with the >4.5 h runners ($P < 0.001$), with further differences observed between 4–4.5 h and >4.5 h finishers ($P = 0.000$) and between the 3–3.5 h and >4.5 h groups ($P < 0.001$). These findings were coupled with those for VE_{max} which showed significant differences between the 2.5–3 h and 3–3.5 h groups ($P = 0.03$) and against the >4.5 h group ($P < 0.001$), while

additional differences were observed between the 3–3.5 h and >4.5 h groups ($P < 0.001$) with the >4.5 h group also showing a significant difference when compared to 3.5–4 h runners ($P = 0.02$). There were no significant differences for HR_{max} or peak blood lactate concentration (PBLa) ($P > 0.05$) between any of the groups.

Blood lactate responses

When considering the blood lactate responses to exercise as shown in Figure 1, significant differences were observed for the appearance of LT2 when expressed as running speed (km·h⁻¹)

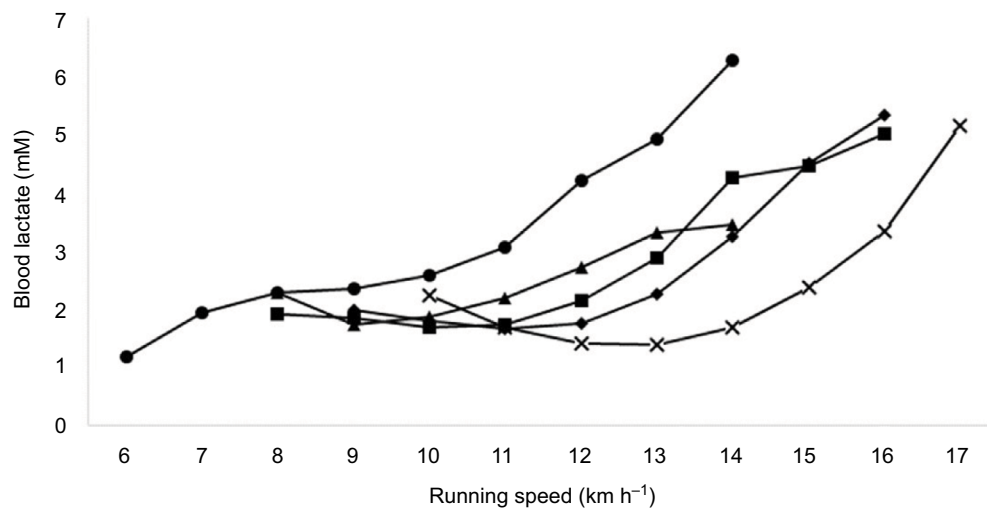


Figure 1 Blood lactate responses to incremental treadmill exercise.

Note: ●, >4.5 h group; ▲, 4-4.5 h group; ■, 3.5-4 h group; ◆, 3-3.5 h group; X, 2.5-3 h group.

between the 2.5-3 h, 3-3.5 h, 3.5-4 h, 4-4.5 h, and >4.5 h groups ($P < 0.001$, $P < 0.001$, $P = 0.005$, $P < 0.001$ and $P < 0.001$, respectively). Further differences were highlighted between the 3-3.5 h and 3.5-4 h groups ($P = 0.002$) and against the >4.5 h group ($P < 0.001$). For LT1, the >4.5 h runners were shown to be significantly different to the 4-4.5 h runners ($P = 0.01$), 3.5-4 h group ($P = 0.000$), 3-3.5 h runners ($P < 0.001$) and with the 2.5-3 h group ($P < 0.001$). The 4-4.5 h group was only shown to be significantly different to the 2.5-3 h runners ($P = 0.002$). Those in the 3.5-4 h group showed a significant difference of $0.8 \text{ km}\cdot\text{h}^{-1}$ against the 3-3.5 h group ($P = 0.005$) with a difference of $2.1 \text{ km}\cdot\text{h}^{-1}$ ($P < 0.001$) against the 2.5-3 h group. Those in the 3-3.5 h group showed a significantly slower running speed for LT1 of $2.3 \text{ km}\cdot\text{h}^{-1}$ ($P < 0.001$) compared to the 2.5-3 h runners. There were no significant differences ($P > 0.05$) for LT1 and LT2 as $\% \dot{V}O_{2\text{max}}$ or for the HR ($\text{b}\cdot\text{min}^{-1}$) associated with LT1 or LT2 between groups. Regarding the lactate concentrations (mM) associated with LT1 and LT2, differences were observed for LT1 between the 2.5-3 h group and the 3-3.5 h runners ($P < 0.001$), 3.5-4 h group ($P < 0.001$), 4-4.5 h group ($P = 0.002$) and against the >4.5 h runners ($P < 0.001$). Additional significant differences were observed between the 3-3.5 h group and the 3.5-4 h ($P = 0.005$) and >4.5 h groups ($P < 0.001$), while for LT2 no differences were observed between any of the groups ($P > 0.05$).

Running Economy

Group-based RE responses are shown in Figure 2 and reflect the relative $\% \dot{V}O_{2\text{max}}$ across the range of running speeds

employed during the treadmill test. Four running speeds were completed by runners from all five of the groups, which were then compared as a function of $\% \dot{V}O_{2\text{max}}$. At $10 \text{ km}\cdot\text{h}^{-1}$, mean responses were $75.3 \pm 6.9\%$, $75.7 \pm 6.7\%$, $68.9 \pm 6.3\%$, $66.9 \pm 6.4\%$ and $61.1 \pm 7.5\%$ for the >4.5 h, 4-4.5 h, 3.5-4 h, 3-3.5 h and 2.5-3 h groups, respectively. Significant differences were observed between the 2.5-3 h group and 4-4.5 h group ($P = 0.01$) and >4.5 h group ($P = 0.003$). Responses at $11 \text{ km}\cdot\text{h}^{-1}$ were $80.4 \pm 7.0\%$ (>4.5 h), $76.9 \pm 12.3\%$ (4-4.5 h), $74.6 \pm 7.2\%$ (3.5-4 h), $71.7 \pm 8.9\%$ (3-3.5 h) and $63.1 \pm 6.5\%$ (2.5-3 h). Significant differences were observed between the 2.5-3 h group and the 3-3.5 h group ($P = 0.003$), 3.5-4 h group ($P < 0.001$), 4-4.5 h runners ($P = 0.05$) and the >4.5 h grouping ($P < 0.001$). In addition, significant differences were observed between the 3-3.5 h runners and the >4.5 h runners ($P = 0.002$), 3.5-4 h group and >4.5 h group ($P = 0.01$). At $12 \text{ km}\cdot\text{h}^{-1}$, >4.5 h runners had a response of $84.8 \pm 5.7\%$ compared to $81.4 \pm 13.7\%$ (4-4.5 h), $79.0 \pm 6.3\%$ (3.5-4 h), $76.0 \pm 6.0\%$ (3-3.5 h) and $67.2 \pm 7.0\%$ (2.5-3 h). Once again, significant differences were observed between the 2.5-3 h and the 3-3.5 h runners ($P = 0.001$), 3.5-4 h group ($P < 0.001$), 4-4.5 h runners ($P = 0.03$) and >4.5 h runners ($P < 0.001$). There were also significant differences between the 3-3.5 h and >4.5 h runners ($P < 0.001$) and between the 3.5-4 h and >4.5 h groups ($P = 0.009$). At $13 \text{ km}\cdot\text{h}^{-1}$, responses of $87.0 \pm 3.1\%$, $79.4 \pm 16.3\%$, $84.2 \pm 6.3\%$, $80.2 \pm 6.0\%$ and $72.1 \pm 9.3\%$ were observed for >4.5 h, 4-4.5 h, 3.5-4 h, 3-3.5 h and 2.5-3 h, respectively. At this running speed, significant differences were highlighted between the 2.5-3 h group and

the 3–3.5 h group ($P = 0.01$), 3.5–4 h group ($P < 0.001$) and >4.5 h runners ($P < 0.001$). The 3–3.5 h runners were significantly more economical than the 3.5–4 h runners ($P = 0.01$) and >4.5 h runners ($P < 0.001$).

Blood chemistry

The results of blood-based variables are presented in Table 3 for which no significant differences were observed ($P > 0.05$) between any of the groups.

Training characteristics

The training indices for the athletes are presented in Table 2 reflecting the outcomes of intensity, frequency and duration of training. Significant differences were observed for training runs per week between the fastest runners, the 2.5–3 h group, and those in the 3.5–4 h group ($P < 0.001$) and those with the slowest finish time (>4.5 h; $P = 0.007$). Those in the 3–3.5 h group showed a significantly greater training frequency than

the 3.5–4 h group ($P = 0.004$), with those in the 3.5–4 h group accumulating significantly fewer runs than the 4–4.5 h group ($P = 0.046$). No other differences were observed for training frequency. Training distance per week was shown to be significantly greater in the 2.5–3 h group when compared to the 3.5–4 h group ($P = 0.01$) and also on comparison to both the 4–4.5 h ($P = 0.006$) and >4.5 h groups ($P < 0.001$). Those in the 3–3.5 h group accrued significantly greater training distance per week than the runners in the 3.5–4 h group ($P = 0.009$) as well as those in the 4–4.5 h ($P = 0.004$) and >4.5 h groups ($P < 0.001$). Further significant differences were also evident between the 3.5–4 h runners and those in the >4.5 h group ($P = 0.001$), while the latter were shown to accumulate significantly less weekly training than the 4–4.5 h group ($P = 0.04$). The average distance completed per session was calculated as a function of $d \cdot \text{week}^{-1}$ and $\text{km} \cdot \text{week}^{-1}$ and was shown to be significantly greater for the 2.5–3 h group compared to the 3.5–4 h ($P = 0.01$) and >4.5 h groups

Table 3 Resting blood-based parameters of the $n = 82$ participants

Parameters	2.5–3 h	3–3.5 h	3.5–4 h	4–4.5 h	>4.5 h
Hemoglobin, $\text{g} \cdot \text{dL}^{-1}$	14.0 ± 1.1	13.9 ± 1.3	13.9 ± 0.9	14.6 ± 1.4	13.7 ± 1.7
Hematocrit, %	42.1 ± 3.1	41.7 ± 3.9	41.9 ± 2.8	43.9 ± 4.3	41.3 ± 5.4
Glucose, mM	5.1 ± 0.8	5.1 ± 0.7	5.2 ± 0.9	4.6 ± 0.5	4.9 ± 0.4
Lactate, mM	0.9 ± 0.4	1.0 ± 0.3	1.0 ± 0.3	1.1 ± 0.3	1.1 ± 0.5
pH	7.45 ± 0.02	7.45 ± 0.01	7.45 ± 0.02	7.44 ± 0.02	7.45 ± 0.01
HCO_3^- , mM	26.2 ± 1.5	26.8 ± 1.6	26.4 ± 1.5	27.4 ± 1.9	25.9 ± 1.7

Note: Data presented as mean \pm standard deviation.

Abbreviation: HCO_3^- , bicarbonate

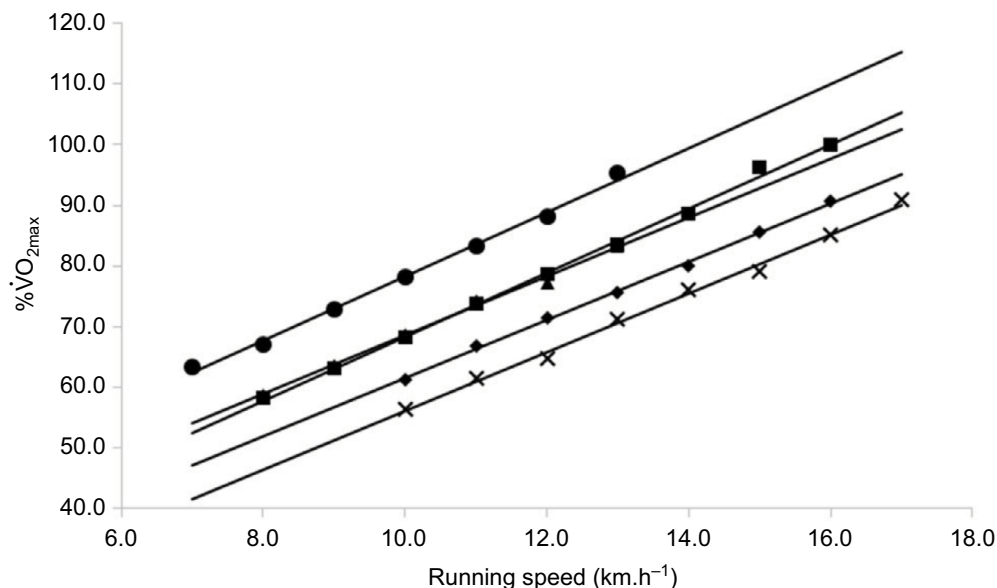


Figure 2 RE expressed as $\% \dot{V}O_{2\max}$ during incremental treadmill exercise.

Note: ●, >4.5 h group; ▲, 4–4.5 h group; ■, 3.5–4 h group; ◆, 3–3.5 h group; X, 2.5–3 h group.

Abbreviation: RE, running economy.

($P = 0.001$), while the 3–3.5 h runners completed a greater distance per session than those in the 4–4.5 h ($P < 0.001$) and >4.5 h groups ($P < 0.001$). Additional differences were observed between the 3.5–4 h group and those in the 4–4.5 h ($P = 0.009$) and >4.5 h groups ($P < 0.001$).

Average running speed during training was calculated from $\text{h}\cdot\text{session}^{-1}$ and $\text{distance}\cdot\text{session}^{-1}$ and highlighted a significantly faster estimated average training speed for those in the 2.5–3 h group compared to the 4–4.5 h group ($P = 0.002$) and >4.5 h runners ($P = 0.005$). In addition, those athletes in the 3–3.5 h pool of runners exhibited significantly faster average speeds during training than the 4–4.5 h group ($P = 0.004$). Training volume was significantly lower for the >4.5 h runners than the 3–3.5 h group ($P < 0.001$) and the 2.5–3 h athletes ($P < 0.001$), while additional differences were observed between the 3–3.5 h runners ($P = 0.005$) and the 3.5–4 h group and against the >4.5 h group ($P < 0.001$). Training load was shown to be significantly higher for the 2.5–3 h group compared to the 3.5–4 h group ($P = 0.002$) and the >4.5 h runners ($P = 0.001$). Meanwhile, those in the 3.5–4 h ($P = 0.004$) and >4.5 h groups ($P < 0.001$) exhibited significantly lower training loads in comparison to the 3–3.5 h runners.

When considering training speeds (average), these were compared to the speed associated with both LT1 and LT2 derived from the treadmill test. For LT1, the 4–4.5 h group showed a significant difference of $-1.0 \pm 2.0 \text{ km}\cdot\text{h}^{-1}$ ($P = 0.02$). For LT2, significant differences were encountered for the 2.5–3 h group ($P < 0.001$), 3–3.5 h runners ($P = 0.001$), 3.5–4 h group ($P < 0.001$), 4–4.5 h group ($P < 0.001$) and for the >4.5 h runners ($P = 0.008$) with differences of $-4.2 \pm 2.4 \text{ km}\cdot\text{h}^{-1}$, $-3.7 \pm 4.4 \text{ km}\cdot\text{h}^{-1}$, $-2.9 \pm 4.1 \text{ km}\cdot\text{h}^{-1}$, $-3.6 \pm 2.2 \text{ km}\cdot\text{h}^{-1}$ and $-3.8 \pm 3.5 \text{ km}\cdot\text{h}^{-1}$, respectively. Training speeds were also compared to the average speed during the marathon indicating significant differences for the 2.5–3 h group ($P < 0.001$) of $-3.4 \pm 2.1 \text{ km}\cdot\text{h}^{-1}$, 3–3.5 h runners of -2.3 ± 8.3 ($P = 0.02$) and the 4–4.5 h group ($P = 0.01$) of $-3.0 \pm 3.3 \text{ km}\cdot\text{h}^{-1}$. From the blood lactate data, a series of training zones were discerned of which the steady zone, denoted as the difference between LT1 and LT2, was of interest. Speed ranges were calculated for this zone for each group and were shown to be $3.4 \pm 0.7 \text{ km}\cdot\text{h}^{-1}$ (2.5–3 h), $2.7 \pm 0.8 \text{ km}\cdot\text{h}^{-1}$ (3–3.5 h), $2.6 \pm 0.9 \text{ km}\cdot\text{h}^{-1}$ (3.5–4 h), 2.6 ± 0.9 (4–4.5 h) and $2.2 \pm 0.8 \text{ km}\cdot\text{h}^{-1}$ (>4.5 h), with significant differences observed between 2.5–3 h and 3–3.5 h ($P = 0.01$), 2.5–3 h and 3.5–4 h ($P = 0.002$), 2.5–3 h and 4–4.5 h ($P = 0.04$) and 2.5–3 h and >4.5 h ($P = 0.000$) groups. Additional differences were also observed between the 3–3.5 h and >4.5 h groups ($P = 0.03$).

Association between training characteristics and physiological variables

Race speed was shown to be related to both LT1 ($r = 0.791$) and LT2 ($r = 0.721$) as well as the O_2 cost ($\text{mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$) at LT1 ($r = 0.701$), while the $\dot{\text{V}}\text{O}_{2\text{max}}$ ($\text{mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$) at LT2 was also significantly related to race speed ($r = 0.719$). However, $\dot{\text{V}}\text{O}_{2\text{max}}$ showed a weaker but still significant association with race speed ($r = 0.535$). When considering the training characteristics, weekly training distance was shown to be significantly correlated with race speed ($r = 0.494$) as well as being correlated with both $\dot{\text{V}}\text{O}_{2\text{max}}$ ($r = 0.453$) and LT2 ($r = 0.510$). Distance completed per session was shown to be significantly related to race speed ($r = 0.563$), LT1 ($r = 0.479$) and LT2 ($r = 0.531$) but not with $\dot{\text{V}}\text{O}_{2\text{max}}$. Meanwhile, training load ($r = 0.525$) but not training volume was correlated with race speed, while load was shown to be associated with $\dot{\text{V}}\text{O}_{2\text{max}}$ ($r = 0.468$) and LT2 ($r = 0.486$), but not with LT1.

Discussion

The purpose of this study was to provide insight into the physical and training characteristics of recreational marathon runners, with a completion time of 229.1 ± 48.7 min. Previous works have alluded to the importance of physical characteristics,^{6,7} training volume,¹⁰ training intensity,¹³ anthropometry,¹⁴ age¹⁵ and sex⁴ to distance running success, but to date very little data are available for the recreational runner.

$\dot{\text{V}}\text{O}_{2\text{max}}$

The mean score for maximal oxygen uptake across all participants was $53.9 \pm 7.4 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$ with a range of $39.4\text{--}79.7 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$, in broad agreement with previous works where runners with a reported finish time of >180 min exhibited a value of $58.7 \pm 1.9 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$.⁶ However, of significance within this study is the data showing a profound difference between groups based on finishing times, indicating that as race speed increases and by association finish time decreases there is a concomitant increase in $\dot{\text{V}}\text{O}_{2\text{max}}$. Although the significance of $\dot{\text{V}}\text{O}_{2\text{max}}$ to marathon performance has been debated,^{2,16} it is apparent that a well-developed oxygen delivery system is fundamental to marathon performance where it has been estimated that elite runners operate at ~65–85% $\dot{\text{V}}\text{O}_{2\text{max}}$ utilizing as much as 92% of their maximal stroke volume.¹⁷ Indeed, a significant correlation was shown between $\dot{\text{V}}\text{O}_{2\text{max}}$ and race speed in this group of recreational runners, again in broad agreement with that reported previously of $r = 0.68$ for runners of a similar ability (139–298 min).^{15,18} Thus, the data suggest that as with elite runners, $\dot{\text{V}}\text{O}_{2\text{max}}$ is

important to recreational marathon performance but is not pivotal once a threshold of $53.0 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$ is exceeded as reflected in Table 1.

Previous works^{7,19,20} have suggested that $\dot{V}O_{2\text{max}}$ is sensitive to training intensity, assuming an adequate training volume is undertaken, with intensities of 80–100% $\dot{V}O_{2\text{max}}$ being most closely associated with its development. Such training has been associated with increased mitochondrial density and enhanced lactate removal which have been shown as a function of the imposed lactate concentration.^{7,21} Interestingly, in the current study, training load, which is a function of intensity, frequency and duration rather than training volume, a composite of duration and frequency was shown to be positively correlated with $\dot{V}O_{2\text{max}}$, thus implying that the discriminating variable as previously reported is the intensity of work. The positive correlation between $\dot{V}O_{2\text{max}}$ and distance completed per training session and distance per week reinforces the notion of a minimum training volume for the development of $\dot{V}O_{2\text{max}}$, suggesting that a minimum training distance per session is ~15 km with a minimum training distance of 56.2 ± 14.8 km per week. This finding is supported by earlier works,⁶ where training distance per week for the slower runners (>180 min) was 57 km which was associated with a $\dot{V}O_{2\text{max}}$ of $58.7 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$. The $\dot{V}O_{2\text{max}}$ for the fastest runners (2.5–3 h) of $63.3 \pm 7.7 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$ was in agreement with previous works $65.6 \pm 1.2 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$ and $60.6 \pm 9.7 \text{ mL}\cdot\text{kg}^{-1}\cdot\text{min}^{-1}$,^{16,33} with similar marathon race times to those reported in this study. Although it has previously been argued that the $\dot{V}O_{2\text{max}}$ is not pivotal to marathon success, this and previous studies lend support to the notion that marathon race speed is partly dependent on $\dot{V}O_{2\text{max}}$. This is highlighted by the observed significant relationship, which was reported in previous studies of elite and nonelite marathoners.^{7,16}

Running economy

Running economy, defined as the O_2 cost²² showed no significant difference across running speeds from 10 kph to 13 kph, equating to treadmill speeds where participants from all group bandings registered a score. As would be anticipated irrespective of race speed, there was a linear response between the O_2 cost of running and treadmill speed, with the slope of the regression line for all groups being in broad agreement with that reported previously for athletes of a similar race speed.¹⁵ However, when expressed as fractional utilization ($\% \dot{V}O_{2\text{max}}$), notable differences were observed, with those in the 2.5–3 h group utilizing $63.5 \pm 6.1\%$ compared to those in the >4.5 h group of $86.2 \pm 7.2\%$ across running speeds of $10\text{--}13 \text{ km}\cdot\text{h}^{-1}$, with those in the 3–3.5 h group utilizing 68.8

$\pm 6.1\%$ and the 3.5–4 h group utilizing $76.1 \pm 6.5\%$. These responses are closely aligned to those witnessed previously³² across similar running speeds; however, it should be noted that when comparing to elite athletes,^{2,7} these responses are significantly higher for the same running speeds. The implication of RE to marathon running speed is well established in elite populations suggesting that successful performances are associated with lower fractional utilization of $\dot{V}O_{2\text{max}}$ ($\% \dot{V}O_{2\text{max}}$). Various postulations have been attributed to the development of RE, but a consistent component is training volume,^{2,23,24} indeed positive correlation ($r = 0.62$) between RE and the number of training years, taken as a proxy for volume.²⁵ However, of note was the suggestion that athletes' "most" economical training speeds are those which they regularly train at.²⁶ In the current study, average training speed was for all groups at and around LT1 corresponding to the aerobic base training zone, with the racing speeds for all groups either being at LT1 or between LT1 and LT2. Despite the assignment of the association between training volume and RE in the literature, no such association was evident within our cohort. However, given the complex nature of RE, the lack of association with a single training component should not be surprising. Indeed, changes in RE have been attributed to nefarious variables, including lower limb flexibility, running mechanics, thermoregulation, running surface, muscle stiffness and limb morphology.^{9,27} Given the reported differences in these variables between trained and untrained runners groups homogeneous for such components as $\dot{V}O_{2\text{max}}$ or aerobic capacity, no such association was evident. Despite the lack of evidence to support an association between training characteristics and RE in this group of runners, it should be noted that RE was still of uppermost significance to marathon running success.

Blood lactate responses

The LT2 characterized by the systematic increase in the blood lactate concentration in response to exercise intensity and reflecting the inability of fatty acid metabolism to sustain oxidative phosphorylation has been previously cited as a potent predictor of endurance performance.^{28,29} A significant relationship has been shown between marathon race speed and LT2,^{2,6,10,15,21,30} in the range of 0.78–0.98, depending on the training/racing status of the athletes. These findings are in broad agreement with those reported in this study where a significant correlation of 0.72 was observed, despite proportionally slower race speeds compared to those reported previously. The significant differences observed for both the speed and associated lactate concentration at LT2 between

each of the groups were coupled with this finding, conforming to previously established principles of a rightward and downward shift in the lactate–speed profile in response to endurance training.^{2,8,31,32} LT2 has been shown to be highly sensitive to both training intensity and volume as reflected by significant responses to both continuous and interval-based training in moderately trained individuals.³³ Within this recreational group of marathon runners, training volumes as reflected by both distance per session and distance per week as opposed to training intensity were shown to be associated with the speed at LT2. Indeed, this finding is reinforced by the positive association between LT2 and training volume, despite the apparent lack of association with training intensity. As with the $\dot{V}O_{2\max}$ responses, some caution should be applied to these findings given that we cannot discern from the data how much of the training was completed above and below LT2 and how intense the interval-based sessions were. However, what is fascinating is the average training speed in comparison to those associated with both LT2 and LT1; not surprisingly all groups were training at speeds which were significantly slower than that associated with LT2. Training below this point has been associated with the development of the aerobic base and associated physiological components including increased stroke volumes, capillarization, mitochondrial density and increased muscle glycogen content;³³ however, as Midgley et al³⁴ highlighted, there has to be a minimum intensity for this type of training below which the adaptive responses are not induced. The aerobic base range is denoted by the difference between LT1 and LT2 and of note within this group of runners was that a significant difference was evident between the 2.5–3 h group and all other groups highlighting a greater aerobic base range for these runners. Of interest was that all groups completed their training with an average speed \geq LT1. Previous works^{7,10,35,36} suggest that marathon runners typically apply the majority of their training (59.5–87.0% week⁻¹) to what were termed long slow distance runs, below LT1. Furthermore, it has also been observed that, depending on the period of the training year, high-performance endurance athletes would typically devote 70–90% of their training to intensities to blood lactate concentrations of \leq 2 mM, corresponding to LT1.^{7,11}

Training characteristics

The most striking difference among the groups was the training distance completed per week with runners in the 3.5–4 h, 4–4.5 h and >4.5 h groups all covering significantly less distance than those in the faster 2.5–3 h and 3–3.5 h groups. However, in comparison to elite and nationally ranked runners with finish

times of ~140 min, these distances are even for those in the fastest group substantially lower; ~145 km.^{10,11} Of note was the difference for training frequency, expressed as sessions·week⁻¹ between elite/nationally ranked runners (8.1 ± 2.8)¹⁰ and the recreational runners in this work (4.6 ± 1.2). It would appear to be the difference in training frequency that distinguishes these athletes as exhibited by the similarity in km·session⁻¹, 14.5 ± 23.6 compared to 17.9 ± 9.1 for the recreational and high-performance runners, respectively. In a comparable group of runners, with a finish time of 231.9 ± 31.7 min, a similar km·session⁻¹ to those in the current study was observed (12.1 ± 15.4) suggesting that for the recreational runner the limiting factor for race speed progression is the amount of time per week that can be devoted to training. The longest run per week was shown to be similar for all groups apart from those in the 2.5–3 h (37.3 ± 5.8 km) group, with these runners completing a significantly longer training run, ~6 km further than those in the other groups; however, in comparison to elite runners (39.0 ± 1.6 km), little difference is evident.

Typically, a recreational runner completes $\sim 3.7 \pm 1.6$ runs per week compared to elite and nationally ranked runners averaging $\sim 14.1 \pm 5.2$ runs per week.¹⁰ It was observed in this group of recreational runners that the average number of training sessions completed was 4.6 ± 1.2 , with a range of 2–7 sessions per week. Of note was the lowest frequency of sessions (4.1 ± 1.3) for the 3.5–4 h group when compared to the other groups. It appears that this relatively low training frequency was compensated for by a significantly higher training distance per week compared to the slower runners and a distance per session that was no different to those in the faster groups. Previous work for elite athletes has suggested that there is a consistency regarding training frequency with ranges of 10–14 sessions per week being reported for endurance runners across an array of distances and disciplines,³⁸ suggesting that the discriminating variables for performance are duration and intensity.³⁴ There was no difference for either LT1 or LT2 when expressed as % $\dot{V}O_{2\max}$ between the five groups of runners; however, absolute training speed was shown to be significantly faster for those in the faster groups 2.5–3 h and 3–3.5 h compared to those in the slower groups a difference of 3.4 km·h⁻¹, equitable to the differences witnessed for LT1, 2.3 km·h⁻¹ and LT2, 4.6 km·h⁻¹. Training intensity has been highlighted as the predominant factor in the adaptive process, assuming a minimum training intensity is achieved,³⁴ with the minimum being a function of the physiological status of the individual, coupled with the duration threshold to establish the adaptive threshold which, for any given physiological component, has to be surpassed to drive the physiological response.^{24,37}

Conclusion

The data presented for a recreational-based group of marathon runners offer some insight into the nature of training and associated physiological status of these runners. Across the spectrum of race completion times, noticeable differences were observed for $\dot{V}O_{2\max}$, RE (% $\dot{V}O_{2\max}$), LT1 and LT2 ($\text{km}\cdot\text{h}^{-1}$ and mM). The data suggest that training frequency coupled with absolute training speed are the determinants of race speed progression and that for similar groups of marathon runners the emphasis in training should be to maximize running speed within the time available, while achieving a minimum training intensity and duration threshold.

Acknowledgment

There were no funding sources associated with this study.

Disclosure

The results of this study are presented in a clear, honest manner, with no falsification, fabrication or inappropriate data manipulation. There are no personal relationships to disclose with this study. The authors report no other conflicts of interest in this work.

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