Introduction

Running economy (RE), the oxygen uptake or metabolic rate for running at a given submaximal speed, is one of the key determinants of distance running performance [4, 6, 14, 16, 18]. Understanding how RE changes with running speed is important for predicting performance of high-level runners and absolute energy expended. However, there is disagreement about the nature of the quantitative relationship between RE and submaximal speed. Most previous research has focused on a narrow range of running speeds (~2–4 m·s⁻¹) in average to good runners and found that oxygen uptake or metabolic rate (VO₂, mlO₂·kg⁻¹·min⁻¹ or E, kcal·kg⁻¹·min⁻¹) increases linearly with running speed [13, 14, 23, 25, 26]. One can also calculate the cost of transport (COT) or the amount of oxygen or energy needed to transport a kilogram of body mass a given distance forward (O₂COT, mlO₂·kg⁻¹·km⁻¹; ECOT, kcal·kg⁻¹·km⁻¹) by dividing oxygen uptake or metabolic rate by running speed [8, 28, 31, 32]. If the intercept is zero, a linear relationship between oxygen uptake or metabolic rate and running speed results in a COT that is independent of running speed [13, 23, 25, 26, 28]. However a positive intercept results in a decreasing COT with speed to some asymptotic value, whereas a negative intercept results in an increasing COT to some asymptotic value. Together, the idea of a linear relationship between oxygen uptake or metabolic rate and running speed, and an invariant COT define what we call the "traditional model" of running energetics.
The traditional model of running energetics has been supported by numerous investigations [8, 23, 30]. These studies measured the metabolic rate for running across a moderate range of submaximal running speeds (~2–4 m·s⁻¹) in average to good runners. In contrast, studies at faster speeds (>4.5 m·s⁻¹) in sub-elite to elite runners [7, 13, 14, 21, 34, 35] are not in agreement. Some studies report a linear relationship between metabolic rate and speed and thus a nearly invariant gross COT [13, 14, 23, 25, 26], whereas others have found that gross COT increased at faster running speeds [7, 25, 35, 36]. Additionally, no studies have compared average runners to sub-elite distance runners, and very few have looked at sub-elite runners at slow speeds. Given the lack of consensus and the importance of RE on distance running performance, a more comprehensive re-examination of the energetics of running is needed.

We sought to quantify the metabolic rate and COT of running in average and sub-elite distance runners over a wide range of submaximal running speeds from 1.78 m·s⁻¹ (4 MPH) to the speed corresponding to each individual’s lactate threshold. We hypothesized that the relationship between the oxygen uptake or metabolic rate and speed would best be described as linear for the average group and curvilinear for the sub-elite group.

Methods

This research protocol was reviewed and approved by the University of Colorado Institutional Review Board and was performed in accordance with the ethical standards outlined by Harris and Atkinson [12]. All subjects gave written informed consent before participating.

Subjects

Twenty healthy male runners (10 average and 10 sub-elite) participated. We classified subjects based on Morgan et al. [28] (Table 1). The average group ran at least three times per week and was capable of running 10 km in 40–60 min. The sub-elite group was capable of running 10 km in less than 30 min at sea level, or less than 31 min at the local altitude (~1600 m).

Experimental procedures

Subjects performed a series of trials on either a custom-made [20] or Treadmetrix (Park City, UT, USA) motorized force-measuring treadmill. Three subjects completed the protocol on the custom-made force treadmill before the axle of the non-drive roller failed. We then finished data collection on the Treadmetrix. Because both treadmills have rigid decks, there is no reason to expect differences in RÉ between the two treadmills. We used the same hand-held tachometer to verify running speed (Shimpo DT-107A, Electromatic Equipment Inc., Cedarhurst, NY, USA) on both treadmills. Subjects arrived to the laboratory 2 h post-prandial to help control for potential effects of diet on metabolic rate. During the experimental trials, we measured rates of oxygen uptake and carbon dioxide production to calculate metabolic rate using the caloric equivalents [11] determined from the respiratory exchange ratios (RER) with a widely accepted open circuit indirect calorimetry system (Parvomedics TrueOne 2400, Sandy, UT, USA) [5]. We calibrated the expired gas analysis system before each testing session using gas fractions calibrated with room air and a primary standard gas mixture within the physiological range (16.01 % O₂ and 4.01 % CO₂). We used a 3L syringe at five distinct flow rates within the expected range of the study protocol to calibrate flow. We considered calibration complete when recorded volumes were within 1 % of the calibration volumes, and gas fractions were within 0.3 % of calibration values (e.g., 20.93 ± 0.06 %).

The study took place over two days to minimize potential effects of fatigue from multiple trials. Day 1 trials began at 1.78 m·s⁻¹ (4 MPH) and day 2 trials began at 2.01 m·s⁻¹ (4.5 MPH). Subjects ran for 4 min at each speed. We increased treadmill speed by 0.45 m·s⁻¹ (1 MPH) in each subsequent stage during both days. During the final minute of each 4-min stage, we asked subjects to provide a rating of perceived exertion (RPE) on the Borg (6–20) scale [9], until they reached an RPE of 15 (speed at RPE of 15 = sRPE15) on both days. Previous data have demonstrated that an RPE value of 16 represents an intensity that corresponds closely to LT [22]. After the completion of each 4-min stage, we took a finger-prick blood sample to determine blood [La] and analyzed blood samples in duplicate with a YSI 2300 lactate analyzer (YSI, Yellow Springs, OH, USA). We monitored blood lactate concentrations [La] to assure a primary reliance on oxidative metabolism ([La] below lactate threshold, LT). We determined LT using the 1-mmol-above-baseline method described by Coyle et al. [4]. For subjects in our protocol this resulted in lactate values for all measured speeds under 2 to 3 mmol·l⁻¹.

Following the second session, subjects took a 10-min break and then completed a VO₂max test. Subjects ran at their sRPE15 on a level grade for two minutes. Then, we increased the grade by 1 % each minute until exhaustion. VO₂max was defined as the greatest 15-s mean value obtained. Our criteria for reaching VO₂max required a plateau in oxygen consumption (i.e., an increase in treadmill grade with no increase in oxygen consumption) and/or a respiratory exchange ratio (RER) over 1.15 [15]. All of our subjects reached VO₂max criteria.

Data analyses/statistics

We performed descriptive statistical analyses to determine means and standard deviations (SD) for average and sub-elite groups. We fit individual subjects’ linear and 2nd-order curvilinear regressions to the VO₂ and E values. R² values for each subject were used to assess the strength of fit for both regression methods. We used a paired samples t-test to compare the means of individual R² values for linear and curvilinear fits. A linear-mixed model was used to determine main effects of speed and group classification on mean O₂COT, and ECOT. We used this model to compare each of these variables across speed to values obtained at the fastest submaximal speed achieved in each group. All statistical analyses were done using RStudio software (version 0.99.892, Boston, MA, USA).

Results

All average subjects completed stages up to a speed of 3.58 m·s⁻¹ (9 stages) whereas all sub-elite subjects completed stages up to a speed of 5.14 m·s⁻¹ (16 stages) before reaching sRPE15. Importantly, a comparison of the individual regression equations revealed no differences between days (p > 0.05).
Oxygen uptake and metabolic rates were greater in the average group compared to the sub-elite group over comparable speeds (1.78–3.58 m·s⁻¹) (p < 0.01) (▶ Fig. 1–3). The R² values for each average subject’s linear and curvilinear fits for \( \dot{\text{V}}\text{O}_2 \) and \( \dot{\text{E}} \) vs. speed were not different (p > 0.05), but were different for the sub-elite subjects (p < 0.05) (▶ Tables 2 and ▶ 3).

For the average group, \( \text{O}_2 \text{COT} \) and ECOT decreased by 10.2% and 7.9% respectively from speeds of 1.78 to 2.68 m·s⁻¹, but did not change over the moderate running speed range of 2.68–
3.58 m·s⁻¹. In the sub-elite group, O₂COT and ECOT decreased by 14.1 % and 14.4 %, respectively, from speeds of 1.78 to 2.68 m·s⁻¹ and did not change over the moderate running speed range of 2.68 to 3.58 m·s⁻¹ (Fig. 3). However, the sub-elite group’s O₂COT and ECOT increased at faster running speeds. For example, O₂COT and ECOT at 5.14 m·s⁻¹, the fastest submaximal speed achieved by all sub-elite subjects, were 7.9–9.6 % and 9.6–12.8 % greater than O₂COT and ECOT across the moderate speed range from 2.46 to 4.02 m·s⁻¹ (p<0.05).

Discussion
We accept our hypothesis that the relationships between oxygen uptake and metabolic rate (VO₂ and ˙E) and speed are linear for the average runners, and curvilinear for sub-elite runners.

Average group
In accordance with the traditional model of running energetics, our average runners sustained only a moderate range of submaximal running speeds (mean speed range: 1.78–4.08 m·s⁻¹). This population is not capable of maintaining steady-state oxygen consumption at faster running speeds because these speeds would correspond to intensities greater than LT [1, 2]. Over the range of submaximal speeds achieved, the average group demonstrated equally strong linear and curvilinear fits characterizing oxygen uptake or metabolic rate and speed.

We found that the average group’s COT was elevated at slow running speeds (1.78–2.46 m·s⁻¹) compared to moderate running speeds. It should be noted that oxygen uptake and metabolic rate for running at these slow speeds is rarely reported. Over the more typically measured moderate speed range of 2.68–3.58 m·s⁻¹, we found that the gross COT was independent of running speed, similar to previous studies [14, 23, 25, 26].

Sub-elite Group
As expected, sub-elite runners sustained faster submaximal running speeds than average runners, which allowed us to measure the oxygen uptake and metabolic rate over a wider range of speeds (mean speed range: 1.78–5.23 m·s⁻¹). The majority of previous investigations of sub-elite to elite runners report linear relationships between metabolic rate and running speed over narrow speed ranges [7, 13, 14, 21]. Over the wider range of speeds sustained by sub-elite subjects (1.78–5.14 m·s⁻¹), the oxygen uptake and metabolic rate vs. speed relationships were best described by curvilinear fits. Our data are in agreement with the findings of Steudel-Numbers and Wall-Scheffler [35], who reported a curvilinear relationship between these variables over a similar wide range of speeds (~2.01–4.91 m·s⁻¹).

We found that the sub-elite group’s COT was greater at both slow and fast running speeds than at moderate running speeds. Although this finding contradicts the traditional model of running energetics, it is not unprecedented. For example, when we converted Daniels and Daniels [7] VO₂ data for elite marathon runners (4.83–6.17 m·s⁻¹) to O₂COT, we calculated a ~9 % increase in O₂COT at 6.17 compared to 4.83 m·s⁻¹ (180–197 mlO₂·kg⁻¹·km⁻¹). Our data are also consistent with Tam et al. [35] who reported significant increases in O₂COT at 5.0 m·s⁻¹ compared to 3.33 m·s⁻¹ in elite distance runners. Tam et al. suggested that this finding could be explained by the increasing contribution of aerodynamic resistance to the metabolic rate during their over-ground running protocol. However, this does not explain our findings or those of Daniels and Daniels [7], because treadmill running involves negligible aerodynamic resistance. Pugh [29] compared track and treadmill running to estimate the energy cost of overcoming air resistance. Two of his four subjects inexplicably had a non-linear relationship between speed and VO₂ on the treadmill. Our data also support the findings of Mayhew [24] as well as Steudel-Numbers and Wall-Scheffler [35], who reported increases in O₂COT and ECOT at slow and fast running speeds compared to moderate speeds in runners during treadmill running over similar speed ranges (~2.33–4.67 m·s⁻¹ and ~2.01–4.9 m·s⁻¹). Thus, the traditional model of running energetics seems appropriate for a narrow range of moderate running speeds (2.68–3.58 m·s⁻¹), but not slower (<2.68 m·s⁻¹) or faster speeds (>3.58 m·s⁻¹). The cost of generating force hypothesis appears to explain the near constant COT for moderate speeds [30]. At very slow running speeds with slow stride frequencies, the spring-mass model for running may not be valid [10]. At faster running speeds, it appears that running adopts more flexed leg postures, which reduce the mechanical advantage that requires recruitment of more muscle volume [3].

Implications for performance predictions
Traditionally, it has been thought that the oxygen uptake and metabolic rate for competitive runners can be calculated from the COT measured at any submaximal running speed or predicted from linear extrapolation of the oxygen uptake or metabolic rate vs. speed relationship [13]. Velocity at VO₂max (vVO₂max) is commonly determined from a linear extrapolation of the oxygen uptake or metabolic rate vs. speed relationship up to VO₂max [6, 8, 27]. The traditional model of running energetics implies that linear extrapolation of submaximal running speeds is an appropriate method for estimating the metabolic rate at all speeds [8, 16, 27]. However, our finding of a curvilinear relationship between oxygen uptake or metabolic rate and speed demonstrates that this assumption is not always valid. In our sub-elite group, vVO₂max calculated from linear
| Subject | Slope  | Intercept | R²  | Quadratic Coefficient | Linear Coefficient | Constant | R²  |
|---------|--------|-----------|-----|-----------------------|-------------------|----------|-----|
| Average |        |           |     |                       |                   |          |     |
| 1       | 10.255 | 4.610     | 0.990 | 0.574                 | 6.410             | 10.653   | 0.992 |
| 2       | 10.735 | 0.667     | 0.986 | 0.860                 | 5.552             | 7.965    | 0.989 |
| 3       | 12.291 | -2.835    | 0.996 | 0.601                 | 8.666             | 2.271    | 0.997 |
| 4       | 14.626 | -5.291    | 0.978 | 1.845                 | 3.504             | 10.372   | 0.985 |
| 5       | 12.266 | -0.5012   | 0.985 | 2.109                 | 0.497             | 15.049   | 0.995 |
| 6       | 11.347 | 6.779     | 0.997 | 0.085                 | 10.836            | 7.499    | 0.997 |
| 7       | 11.164 | 5.985     | 0.985 | 1.200                 | 4.68              | 14.831   | 0.988 |
| 8       | 10.510 | 7.221     | 0.994 | 0.328                 | 8.637             | 9.778    | 0.995 |
| 9       | 10.825 | 5.781     | 0.991 | -0.472               | 13.354            | 2.551    | 0.991 |
| 10      | 10.676 | 2.457     | 0.985 | 1.672                 | 0.971             | 15.706   | 0.994 |
| Mean ± SD | 11.470 ± 1.306 | 2.487 ± 4.340 | 0.989 ± 0.006 | 0.879 ± 0.824 | 6.289 ± 4.153 | 9.667 ± 4.790 | 0.992 ± 0.004 |
| Sub-elite |      |           |     |                       |                   |          |     |
| 11      | 13.271 | -1.868    | 0.991 | 0.981                 | 6.477             | 8.850    | 0.996 |
| 12      | 10.834 | 0.403     | 0.973 | 1.827                 | -1.819            | 20.364   | 0.997 |
| 13      | 10.667 | 0.595     | 0.975 | 1.585                 | -0.306            | 17.904   | 0.994 |
| 14      | 13.306 | -4.431    | 0.971 | 2.092                 | -2.118            | 21.177   | 0.997 |
| 15      | 13.424 | -3.389    | 0.990 | 0.884                 | 6.905             | 7.435    | 0.995 |
| 16      | 11.602 | 0.763     | 0.976 | 1.818                 | -0.988            | 20.623   | 0.996 |
| 17      | 12.521 | -2.075    | 0.990 | 0.996                 | 5.402             | 9.452    | 0.996 |
| 18      | 12.128 | 1.093     | 0.987 | 0.693                 | 7.174             | 9.114    | 0.990 |
| 19      | 12.227 | -0.598    | 0.963 | 2.716                 | -5.971            | 27.349   | 0.999 |
| 20      | 13.040 | -5.497    | 0.980 | 1.944                 | -0.4213           | 15.738   | 0.998 |
| Mean ± SD | 12.302 ± 1.008 | -1.500 ± 2.340 | 0.980 ± 0.009 * | 1.554 ± 0.647 | 1.434 ± 0.648 | 15.801 ± 6.783 | 0.996 ± 0.002 * |

Individual linear and curvilinear fit slopes, intercepts, curvilinear coefficients, quadratic coefficients, and $R^2$ values for VO$_2$ vs. speed relationships up to the speed at lactate threshold for average and sub-elite groups. * = Significant difference between linear and 2nd-order polynomial models (p < 0.05)
### Table 3
Linear and curvilinear models for $\dot{E}$ vs. speed.

| Subject | Slope | Intercept | $R^2$ | Quadratic Coefficient | Linear Coefficient | Constant | $R^2$ |
|---------|-------|-----------|-------|------------------------|--------------------|----------|-------|
| **Average** |
| 1 | 0.053 | 0.015 | 0.993 | 0.003 | 0.032 | 0.049 | 0.995 |
| 2 | 0.056 | -0.006 | 0.984 | 0.006 | 0.022 | 0.041 | 0.989 |
| 3 | 0.063 | -0.021 | 0.996 | 0.005 | 0.035 | 0.018 | 0.999 |
| 4 | 0.075 | -0.039 | 0.977 | 0.009 | 0.019 | 0.041 | 0.984 |
| 5 | 0.065 | -0.013 | 0.981 | 0.012 | -0.002 | 0.075 | 0.992 |
| 6 | 0.058 | 0.026 | 0.998 | 0.001 | 0.049 | 0.038 | 0.999 |
| 7 | 0.056 | 0.022 | 0.982 | 0.007 | 0.020 | 0.069 | 0.986 |
| 8 | 0.055 | 0.026 | 0.994 | 0.002 | 0.042 | 0.044 | 0.994 |
| 9 | 0.057 | 0.020 | 0.990 | -0.001 | 0.061 | 0.014 | 0.990 |
| 10 | 0.054 | 0.004 | 0.979 | 0.009 | 0.002 | 0.075 | 0.988 |
| Mean ± SD | 0.059 ± 0.006 | 0.003 ± 0.022 | 0.987 ± 0.008 | 0.005 ± 0.004 | 0.028 ± 0.020 | 0.046 ± 0.021 | 0.992 ± 0.005 |

| **Sub-elite** |
| 11 | 0.066 | -0.013 | 0.984 | 0.008 | 0.014 | 0.069 | 0.994 |
| 12 | 0.051 | 0.001 | 0.978 | 0.007 | -0.001 | 0.082 | 0.996 |
| 13 | 0.055 | -0.006 | 0.971 | 0.009 | -0.009 | 0.096 | 0.994 |
| 14 | 0.065 | -0.028 | 0.958 | 0.011 | -0.020 | 0.112 | 0.990 |
| 15 | 0.070 | -0.026 | 0.986 | 0.006 | 0.026 | 0.046 | 0.994 |
| 16 | 0.058 | -0.002 | 0.972 | 0.010 | -0.010 | 0.104 | 0.995 |
| 17 | 0.063 | -0.016 | 0.989 | 0.006 | 0.021 | 0.051 | 0.997 |
| 18 | 0.062 | -0.005 | 0.985 | 0.005 | 0.024 | 0.055 | 0.991 |
| 19 | 0.061 | -0.009 | 0.960 | 0.014 | -0.033 | 0.135 | 0.998 |
| 20 | 0.066 | -0.035 | 0.976 | 0.011 | -0.009 | 0.083 | 0.998 |
| Mean ± SD | 0.061 ± 0.006 | -0.014 ± 0.012 | 0.977 ± 0.011 * | 0.009 ± 0.003 | 0.0004 ± 0.020 | 0.083 ± 0.029 | 0.995 ± 0.003 * |

Individual linear and curvilinear fit slopes, intercepts, quadratic coefficients, linear coefficients, and $R^2$ values for $\dot{E}$ vs. speed relationships up to the speed at lactate threshold for average and sub-elite groups. * = Significant difference between linear and 2nd-order polynomial models ($p<0.05$).
extrapolation of data collected at a series of moderate speeds ranging from 2.68–3.58 m·s⁻¹, averaged 0.80 m·s⁻¹ faster than for faster speeds ranging from 4.47 to 5.14 m·s⁻¹ (6.45 vs. 5.67 m·s⁻¹). $v\dot{VO}_{2\text{max}}$, calculated using a linear extrapolation over a wider range of speeds from 2.68 to 5.14 m·s⁻¹ (a range excluding slow running speeds) results in a nearly 0.33 m·s⁻¹ faster $v\dot{VO}_{2\text{max}}$ than if calculated using a curvilinear extrapolation (5.45 vs. 5.77 m·s⁻¹). Thus, $v\dot{VO}_{2\text{max}}$ values are often overestimated when calculated from linear extrapolation of moderate speed values. More valid estimates of sub-elite $v\dot{VO}_{2\text{max}}$ can be obtained from linear extrapolation at a series of fast running speeds (>4.47 m·s⁻¹) or a curvilinear extrapolation at a wider range of speeds.

Recent world record performances in the marathon have motivated predictions of the ultimate marathon performance from a variety of physiological parameters. Joyner et al. [19] argued that breaking the 2-h marathon will require exceptional running economy. Extrapolating the metabolic rate from our sub-elite runners at 2-h marathon pace demonstrates that a curvilinear extrapolation results in a 15% greater predicted metabolic rate compared to a linear extrapolation. Because this extrapolation is from treadmill running, this predicted difference would likely be greater for road racing due to the nonlinear relationship between air resistance and speed. Consequently, a sub-2-h marathon may be more difficult than previously appreciated.

**Limitations**

Competitive distance runners compete at speeds well above LT during a 10 km race. To our knowledge, it is not yet clear whether, or how, the total metabolic rate (both oxidative and non-oxidative) of running changes at these intensities [2, 17]. We intentionally did not examine the metabolic rate for running at speeds corresponding to intensities greater than LT in order to ensure only oxidative metabolism. As a result, we were not able to quantify the metabolic rate for running at a competitive race pace. Further, we are not able to ensure that the relationships observed here will hold true up to race pace. Finally, we performed this investigation at a moderate altitude of ~1600 meters albeit with a subject population that was acclimatized to this environment.

**Future directions**

We have not yet elucidated the biomechanical and physiological basis for the relationships reported here. Previous investigations have demonstrated correlations between biomechanical parameters and the metabolic rate for running. Some correlates include ground contact time, vertical ground reaction force, elastic energy return, and vertical center of mass oscillation [33]. It is possible that changes in running mechanics may, in part, explain the curvilinear metabolic rate vs. running speed relationships that we observed.

**Conclusions**

We conclude that the traditional model of running energetics is appropriate only over a narrow range of moderate running speeds (~2.68–3.58 m·s⁻¹). Because average runners are capable of sustaining only a moderate range of submaximal running speeds, the traditional model of running energetics accurately describes the metabolic rate for running in these populations. However, this model significantly underestimates the ECOT at slower (speeds < 2.68 m·s⁻¹) and faster speeds (speeds > 3.58 m·s⁻¹) by up to 12.8% (5.14m·s¹). In order to best model the relationship between metabolic rate and running speed of sub-elite competitive runners, we suggest that future investigations measure subjects’ metabolic rate at a series of four to five fast running speeds that approach a runners’ competitive race pace and use curvilinear fitting to best extrapolate the race pace.

**Acknowledgements**

This research was partially supported by an unrestricted gift from Saucony Inc. The views expressed are those of the authors and do not reflect those of Saucony Inc.

**Conflict of Interest**

The authors do not have any conflicts of interest to declare.

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