



# Short-term resource allocation during extensive athletic competition

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**Short-term resource allocation during extensive athletic competition**

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1 Figure

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Abbreviated title: Short-term resource allocation during competition

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**Abstract**

**Objectives:** Following predictions from life history theory, we sought to identify acute trade-offs between reproductive effort (as measured by psychological arousal) and somatic maintenance (via functional measures of innate immunity) during conditions of severe energetic imbalance.

**Methods:** Sixty-six male ultramarathon runners (ages 20 to 37 years) were sampled before and after a lengthy race. Saliva and sera were collected for testosterone and immunological analyses (hemolytic complement activity and bacterial killing ability). Lean body mass was assessed by bioelectrical impedance, and libido was measured using a slideshow of arousing and neutral images.

**Results:** Following predictions, there was a significant decrease in salivary testosterone levels (109.59 pg/ml versus 97.61 pg/ml,  $p < 0.001$ ) and arousal scores in response to provocative images (5.40 versus 4.89,  $p = 0.001$ ) between pre- and post-race time points. Additionally, participant bacterial killing ability ( $p = 0.035$ ) and hemolytic complement activity ( $p = 0.021$ ) increased between pre- and post-race.

**Conclusions:** Decreased libido and testosterone with concomitant heightened innate immune responses suggest a shift in energetic priorities away from reproduction and towards maintenance/defense during a period of energetic stress.

**Keywords:** Life history theory, testosterone, libido, innate immunity

Life history theory predicts the existence of trade-offs between competing physiological functions relating to reproduction, maintenance, defense, growth, and storage under conditions of limited environmental resources (Stearns, 1992; Reznik, Nunney & Tessier, 2000). Individuals that have developed effective mechanisms for both acquisition and optimal allocation of energy in their particular ecological niche are thereby advantaged (Lotka, 1922; Angiletta et al., 2003). According to evolutionary theory, every individual organism should exhibit behavior intended to enhance genetic contributions to subsequent generations. However, a life history strategy involving a greater allocation of resources towards reproduction necessitates a reduction in the resources available for other functions. Consequently, an individual may allocate energy to traits which enhance fecundity and fertility or invest in traits enhancing survivorship.

Fluctuations in the availability of environmental energy influence the relative investment in reproduction versus survivorship. This is exemplified by nutritional infecundability, whereby a negative shift in the relative balance between energy intake and expenditure adversely affects the fecundity in a range of mammalian species (Bronson, 1991; Wade et al., 1996; Mosley, 2012). Despite the pathological connotations, it is likely that conditions of sub-optimal nutrition will be encountered at some point during the lifetime of a mammal, and animals will have been selected to protect functions necessary for survival at the expense of less essential processes. Thus, available energy is partitioned according to a set of priorities that maximize the chance of survival and thereby optimize long term reproductive success (Wade, 1992).

The principle of allocation proposes the existence of negative correlations between investment in reproduction and somatic traits such as maintenance, growth,

or storage (Cody, 1966; Roff, 1992; Zera & Harshman, 2001). Despite the intuitive appeal of a stereotypical life history trade-off featuring a negative relationship between two traits, such negative covariation in traits are frequently absent when phenotypic comparisons are made between individuals within a population (Glazier, 2000). Furthermore, negative correlations are less likely to be reported in a field setting than in a tightly controlled laboratory study. This may be explained by inter-individual variation in resource acquisition exceeding variation in resource allocation (Van Noodwijk and de Jong, 1986).

Identifying quasi-experimental scenarios in which negative correlations are observable in the field would break new ground in the study of human life history trade-offs. While one cannot control individual energy intake in the field without compromising ecological validity, it may be possible to nullify the effect of this variation in resource acquisition by experimentally controlling energy balance. It is not the absolute values of energy intake and expenditure which force the individual to make resource allocation decisions, but the imbalance of the two (Bronson, 1991). Therefore, if the energy deficit is high enough, variation in food intake between individuals will be inconsequential; all individuals will be energetically stressed.

This study uses male participants from running ultramarathons to investigate the relative investment between reproduction and survivorship. Ultramarathons, defined as races lasting over 4 hours (Knez et al., 2006) (but often several days in length). Recent work suggests that high levels of energy expenditure cause metabolic adaptation to reduce total energy output (Pontzer et al. 2016), and the high energetic costs of locomotion necessitated by ultramarathons are expected to necessitate physiological trade-offs. This is due to energy deficit induced by a combination of the high levels of energy expenditure in physiological systems linked

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2  
3 to locomotion, and the lack of opportunity to ingest any significant meals (Knechtle  
4  
5 and Bircher, 2005; Knechtle et al., 2005).  
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7  
8 Testosterone functions as a physiological regulator, influencing the allocation  
9  
10 of energetic resources between life history traits (Muehlenbein and Bribiescas, 2005;  
11  
12 Hau, 2007). From a physiological perspective, testosterone plays a significant role in  
13  
14 stimulating muscle growth (Kadi, 2008), which is beneficial in both inter-sexual  
15  
16 (Lavrakas, 1975; Dixon, 2003; Fan, 2005; Frederick and Haselton, 2007) and intra-  
17  
18 sexual (Bribiescas, 2001; Dijkstra and Buunk, 2002) selection. Testosterone is also  
19  
20 implicated in behavioral aspects of human male reproductive effort. Sex drive, or  
21  
22 libido, is a vitally important motivational force in human behavior (Darwin, 1871;  
23  
24 Bancroft, 1988; Ariely and Loewenstein, 2006). In addition to its role in generating  
25  
26 sexual motivation (Beach, 1976; Sherwin, 1988; Baumeister et al., 2001),  
27  
28 testosterone is important in modulating qualities believed to be beneficial in the male  
29  
30 mating effort, such as confidence and assertiveness in social situations (Elias, 1981;  
31  
32 Bagatell et al., 1993; Ellison, 2003; Morley, 2003). Consequently, we measured  
33  
34 testosterone levels, lean body mass, and libido of participants.  
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39 Measures of investment in immune function, a proxy for the body's investment  
40  
41 in survivorship, were taken through the use of two functional assays of innate  
42  
43 immune responses (bacteria killing assay and hemolytic complement assay). The  
44  
45 purpose of the bacteria killing assay is to measure the functional ability of integrative  
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47 immunological components, including opsonizing proteins and antibodies, to lyse a  
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49 known quantity of *Escherichia coli* bacteria relative to a positive control (Muehlenbein  
50  
51 et al. 2011). The hemolytic complement assay serves to measure the ability of the  
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53 antibody-dependent pathway of the complement system to lyse pathogens. These  
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55 assays assess innate immune responses only. While a comprehensive assessment  
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of immunocompetence would require examination of the adaptive immune response as well, these functional assays are arguably more relevant than simple measures of inflammation (e.g., C-reactive protein or sIgA), and meaningful results have previously been reported in humans and nonhuman primates (Prall and Muehlenbein, 2014; Georgiev et al., 2015; Prall and Muehlenbein, 2015; Prall et al., 2015).

This paper aims to build upon current knowledge of human male energetic investment in reproduction and survivorship through analysis of male ultramarathon runners. While past studies predominantly consider trade-offs acting on a long-term evolutionary scale, the present study employs the conceptual framework provided by life history theory as a reference to consider acute trade-offs. It was hypothesized that markers of investment in reproductive function (Demas et al., 2011) will decrease in magnitude, allowing for an increase in markers indicative of investment in short-term survival. This is due to the high costs of male mating effort (Ellison, 2003) and the suggestion that survival may be prioritized over other processes (Bronson, 1991).

**Materials & Methods**

Self-reported heterosexual male athletes (total of 66) were recruited at the 2013 North Downs Way 100 (102.6 miles) race, held 10-11<sup>th</sup> August (<http://www.centurionrunning.com/north-downs-way-100-2015/>). Athletes received an email explaining the study prior to race day, and were invited to participate. The study was approved by the University of Cambridge Human Biology Research Ethics Committee. The race is a relatively high profile event, with course records of



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2  
3 15:44:39 and 20:10:39 (hr:min:s) for men and women, respectively. Participants had  
4  
5 a mean age of 29.4 years (range 20-37 years), and all were of European descent.  
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7  
8 Samples and measurements were taken both before and after the race from  
9  
10 athletes who completed the full race distance in order to determine the effects of the  
11  
12 energetic deficit caused by race participation. Measurements included height,  
13  
14 weight, and lean body mass (via bioelectrical impedance analysis). Five ml saliva  
15  
16 and 5ml blood serum samples were collected for testosterone measurement and  
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18 bacteria killing and hemolytic complement assays. While all of the pre-race  
19  
20 measurements were taken between 1800 and 2200 hours, the practicalities of  
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22 varying finish times (ranging from 22:30 hours on the day of race start to 12:00 hours  
23  
24 the following day) meant that post-race measurements were not standardized by  
25  
26 time. All post-race measurements were collected within 15 minutes of completion of  
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28 the race course following established methods (Davies and Thompson, 1986;  
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30 Malarkey et al., 1993; Berg et al., 2008; Lucas et al., 2008; Stuempfle et al., 2010).  
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### 34 ***Reproductive effort***

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36 Testosterone was measured from saliva, rather than serum. Salivary analysis  
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38 facilitates repeated sampling, enhances subject compliance, and provides reliable  
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40 data (Dabbs, 1990; Shankar and Dandekar, 2012). Salivary measurements may  
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42 underestimate testosterone levels; however, this is more problematic in females, and  
43  
44 this study sought to track changes in testosterone levels within subjects rather than  
45  
46 absolute differences between individuals (Shirtcliff et al., 2002).  
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49  
50 A 5ml saliva sample was collected in multiple aliquots using the Salimetrics  
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52 Saliva Collection Aid (#5016.02). Subjects refrained from eating, drinking, chewing  
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54 gum, or brushing teeth in the 30 minutes preceding saliva collection.  
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56 Cotton/polyester swabs were avoided and participants were screened for mouth  
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injuries such as open sores to prevent blood contamination. Samples were immediately stored at -18°C, and were frozen at -80°C within 48 hours of collection. All samples were analyzed within two weeks of collection, using the DRG Salivary Testosterone ELISA kit (SLV-3013). Pre- and post-race hormone levels were tested for normality, which was confirmed. Intra-assay CV = 6.4%, inter-assay CV = 10.26%. The high and low controls were within established values.

**Lean body mass**

Lean body mass was measured using bioelectrical impedance analysis (BIA) (BodyStat Quadscan4000). It is appreciated that *in vivo* measurements cannot measure body composition directly, but rather make predictions from other physiological metrics. BIA was chosen because of its speed, simplicity, high precision and suitability for assessing short-term changes in individuals (Johnson et al., 1985; Roubenoff, 1996; Wells and Fewtrell, 2006). In order to avoid the inherent problems of predicting total body water (TBW), regression equations for converting between impedance and TBW were avoided. Instead, the simple index of 1/impedance, which reliably predicts lean mass index (lean mass/height<sup>2</sup>), was used (Wells et al., 2007).

**Libido**

For the purposes of this study, the terms libido and sexual desire are used to refer to desire to partake of sexual activity (Levin, 1994). This attitude is viewed as being ever-present on a continuous scale, and is responsible for inducing a desire for sexual activity upon stimulation (Levin, 1994). A measurement of libido was taken both before and after the ultra-marathon race, using near-nude female images as stimuli. Visual stimuli have been shown to be powerfully sexually provocative for

men (Quiney et al., 1996; Graziottin, 2000; Hietanen and Nummenmaa, 2011). All participants self identified as being heterosexual.

Before the race, subjects practiced using the Self-Assessment Manikin (SAM, these practice scores were not analyzed - Crabbe et al., 2007), a pictogram exhibiting levels of affect on a nine-point scale (Lang, 1980; Hodes et al., 1985). The SAM has gained popularity in such work as it is a low-cost and relatively easy method for quickly determining affective response in many contexts (Bradley and Lang, 1994). Subjects were randomly divided into two groups. The first group was shown a 45-photo slide show (slide show A) before the race and a different 45-photo slide show (slide show B) after the race, and this order was reversed for the second group. Each slide show comprised 15 provocative near-nude female images, and 30 International Affective Picture System (IAPS) images (15 positive valence, 15 negative valence), and were viewed under standardized, private, and relaxed conditions. The order of images within each slide show was pseudo-random such that two photos from the same category (provocative near-nudes, positive and negative) could be viewed consecutively (Smith, 2012). Immediately after each picture was viewed, the participant used the SAM to rate their valence and arousal experienced whilst viewing the picture. Athletes were not asked to rate the same image more than once, to avoid possible development of habituated responses (Rosen, 1973; Freund et al., 1974; Rubin and Henson, 1976; Heiman, 1977; Julien, 1984). See Appendix for further details of this protocol used.

### ***Innate immune function***

*In vitro* bacteria killing assays were used with serum to measure innate immunity. After a test-run to optimize dilutions, serum was diluted 1:12 in L-glutamine supplemented CO<sub>2</sub> Independent Media (Gibco #18045). A single

lyophilized *E. coli* pellet (MicroBiologics Epower Microorganisms #0483E7) was reconstituted in sterile phosphate buffered saline and then diluted into a working solution, which produced approximately 200-300 colonies per 20µl of aliquot. Aliquots of bacteria working solution were added to diluted serum in a microcentrifuge tube, vortexed, and incubated for 30 minutes. After incubation, the samples were spread on trypticase soy agar plates (BD BBL #211043) in triplicate and incubated overnight at 37°C. The number of colonies on each plate the next day were counted, and the percent bacteria killed for each sample relative to a positive control (media and bacteria only) was calculated.

Serum was also used to measure the classical pathway of complement protein activity via a hemolytic complement assay (Sinclair and Lochmiller, 2000; Demas et al., 2011). Following test runs to optimize dilutions, serum was diluted 1:90 and 1:180 in dextrose gelatin veronal buffer (Lonza BioWhittaker #10-539) and pipetted in duplicate onto a round-bottom 96-well plate. Sheep red blood cells (MP Biomedicals #55876) were washed in sterile phosphate buffered saline, and further diluted to 0.6% in veronal buffer. Anti-sheep red blood cell antibodies (Sigma #S1389-1VL) were diluted to 1:40 in veronal buffer. Both the diluted antibodies and sheep red blood cells were then added to each sample well, and the plate was vortexed. Following incubation for 1.5 hours at 37°C, the plate was centrifuged, and the supernatant transferred to a new 96-well round-bottom plate. The absorbance of this supernatant was read at 405nm. Results of this assay are expressed as CH50 units, or the inverse of the dilution predicted to cause 50% hemolysis (Mayer, 1948).

Statistics

Paired samples t-tests were performed to compare pre- and post-race metrics. Associations between baseline (pre-race) metrics were explored using Pearson correlation analysis. All analyses were performed using SPSS v21, and significance set at  $<0.05$ . *N* values vary due to some athletes not wishing to participate in certain aspects of the study (e.g., blood testing).

## Results

### *Height and weight*

There was a significant decrease in both height ( $n = 66$ , pre-race height  $M = 176.9\text{cm}$ , post-race  $M = 176.4\text{cm}$ , mean change =  $-0.508\text{cm}$ ; 95% confidence interval  $-.846 - -.169$ ) and weight ( $n = 66$ , pre-race  $M = 76.9\text{kg}$ , post-race  $M = 75.1\text{kg}$ , mean change =  $-1.83\text{kg}$ ; 95% confidence interval  $-2.30 - -1.37$ ) during the race.

### *Testosterone*

There was a significant decrease in salivary testosterone levels from before ( $n = 52$ ,  $M = 109.59\text{ pg/ml}$ ) to after the race ( $n = 52$ ,  $M = 97.61\text{ pg/ml}$ , mean change =  $-12.0\text{ pg/ml}$ ; 95% confidence interval  $-14.9 - -9.03$ ).

### *Impedance*

There was a significant decrease in bioelectrical impedance from before ( $n = 66$ ,  $M = 496.80$ ) to after the race ( $n = 66$ ,  $M = 454.90$ , mean change =  $-41.9$ ; 95% confidence interval  $-53.3 - -30.5$ ).

There was a significant increase in  $1/\text{impedance}$  ( $1/z$ ) from before ( $n = 66$ ,  $M = 0.00204$ ) to after the race ( $n = 66$ ,  $M = 0.00223$ , mean change =  $0.000185$ ; 95% confidence interval  $.000134 - .000235$ ).

### *Response to visual stimuli*

While there was a significant decrease in arousal scores from before (n = 52, M = 5.29) to after the race (n = 52, M = 5.04, mean change = -.248; 95% confidence interval -.389 – -.108), there was no change in either valence (n = 52, before M = 5.37, after M = 5.29, SD = 0.69, mean change = -.0722; 95% confidence interval -.168 - .0236) or dominance (n=52, before M = 5.19, after 5.16, SD = 0.67, mean change = -.0325; 95% confidence interval -.121 - .0556).

Further investigation considered how the three photo subtypes (provocative, positive valence, and negative valence) contributed to the overall decrease in arousal in response to the entire 45-photo slide show. This revealed a significant decrease in arousal scores in response to provocative near-nude images from before (n = 52, M = 5.40) to after the race (n = 52, M = 4.89, mean change = -.501; 95% confidence interval - .782 – -.220). In contrast, there was no significant difference in the arousal scores in response to positive valence images (n = 52, before M = 5.19, after M = 5.10, mean change = -.0833; 95% confidence interval - .230 – .0637) or in response to negative valence images (n = 52, before M = 5.27, after M = 5.11, mean change = -.160; 95% confidence interval -.365 – .0445). The difference in the total arousal score before and after the race can therefore be explained by a decrease in arousal in response to the near-nude provocative images.

**Bacteria killing assay**

There was a significant increase in bacterial killing ability from before (n = 34, M = 30.0 percent killing) to after the race (M = 41.0 percent killing, mean change = 11.0; 95% confidence interval .847 – 21.2)

**Hemolytic complement assay**

There was a significant increase in hemolytic complement from before ( $n = 34$ , 124.60 CH50) to after the race ( $n = 34$ , 139.40 CH50, mean change = 14.9; 95% confidence interval 2.41 – 27.3)

These results are summarized in Table 1. Figure 1 shows percent change in measures of reproductive effort and immune function.

INSERT TABLE 1 AND FIGURE 1 ABOUT HERE

### ***Correlation analyses***

Consideration of both baseline (pre-race) and post-race metrics revealed a positive correlation between testosterone and total arousal (pre-race  $n = 52$ ,  $r = 0.33$ ,  $p = 0.017$ ; post-race  $n = 52$ ,  $r = 0.35$ ,  $p = 0.011$ ). This result is unsurprising as testosterone has been implicated in sexual arousal.

The only other significant pre-race correlations came between the subgroups of imagery contributing to the total arousal score, although there was a significant positive correlation between post-race impedance and the provocative imagery subset of arousal scores ( $n = 52$ ,  $r = 0.31$ ,  $p = 0.025$ ). We do not propose an explanation for this.

### **Discussion**

It was hypothesized that the energy deficit associated with ultramarathon participation would impose an energetic allocation trade-off, causing an increased investment in short-term survival (as measured by innate immune function) relative to reproductive effort. This hypothesis is supported by the observation of a significant increase in bacteria killing assay and hemolytic complement assay metrics, and a decrease in testosterone and arousal scores.

Evidence from sports medicine and related fields generally support the findings of increased immune function following intense exercise. These findings include temporary increases in the pro-inflammatory cytokines interleukin-6 (IL-6) and tumor necrosis factor- $\alpha$  (TNF- $\alpha$ ), granulocyte colony stimulating factor (GCSF), monocyte chemoattractant protein 1 (MCP-1), and other immunological factors (Nieman, 2012). Natural killer (NK) cell function and numbers increase following bouts of exercise as well (Shephard and Shek, 1994; Woods et al., 1999; McFarlin et al., 2008), and lysozyme, an important anti-bacterial component of innate mucosal immunity, shows a significant increase in male runners after two hours of running (Costa et al., 2012). In endurance athletes of both sexes, both leukocyte and lymphocyte counts are increased after an hour of intense exercise (McFarlin et al., 2008). While there are some reports of declining immune measures after exercise (e.g., sIgA), these results could be due to a number of different parameters, including sample collection, hydration, and even environmental factors (reviewed in Bishop and Gleeson, 2009).

On the whole, however, strenuous exercise is associated with changes in several inflammatory mediators (e.g., IL-6) which are released from injured and/or contracting muscle cells during intense and prolonged exercise (Petersen and Pedersen, 2005). This leads to local neutrophil and monocyte invasion and phagocytosis of debris from damaged myocytes (Evans and Cannon, 1991; Weight et al., 1991; Northoff et al., 1994). Changes in core body temperature have also been shown to affect leukocyte and neutrophil counts during bouts of exercise, although this effect is also mediated by environmental temperature (Mestre-Alfaro et al., 2012; Sureda et al., 2015). Given the systemic effects of many immune molecules, it may be that localized increases induce or contribute to a general



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3 increase in immune function, as measured here. However, because these same  
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5 molecules and cells may also be involved in minimizing or repairing the damage  
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7 caused by intense physical exertion (Tidball, 2005), we maintain that this represents  
8  
9 an additional investment in somatic maintenance, rather than a confounding factor.  
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11 Furthermore, measuring multiple aspects of immune function (assays which do not  
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13 measure inflammation, i.e., complement activity and bactericidal capabilities) may  
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15 mitigate the effects of locally produced immune mediators on our results, as different  
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17 immune responses are often driven by different mechanisms.  
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21 Although human males do not face the same energetic challenges as females  
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23 (i.e., gametogenesis, gestation, and lactation), and energetic investment in  
24  
25 spermatogenesis is minimal (Bagatell and Bremner, 1990; Elias, 1992), mating effort  
26  
27 is still metabolically demanding. Energy costs include competition and mate  
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29 attraction, as well as protection of and provisioning for mates and offspring  
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31 (Muehlenbein and Bribiescas, 2005). While male mating effort may therefore be  
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33 considered as largely behavioral, it does have important physiological correlates  
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35 stemming from the need to signal underlying quality (Zahavi, 1975; Ellison, 2003).  
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39 The Immunocompetence Handicap Hypothesis (Folstad and Karter, 1992),  
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41 which builds upon Zahavi's (1975) handicap hypothesis for the evolution of  
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43 secondary sexual characteristics, suggests that testosterone mediates signals of  
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45 reproductive status and quality in humans and other animals. In humans, muscle  
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47 mass provides one such signal (Griggs et al., 1989; Kadi, 2008). Muscularity  
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49 provides benefits in sexual selection from both inter- and intra-sexual perspectives  
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51 (Lavrakas, 1975; Bribiescas, 2001; Dijkstra and Buunk, 2002; Frederick and  
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53 Haselton, 2007; Gallup et al., 2007). However, skeletal muscle mass accounts for  
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55 approximately 20% of human male basal metabolic rate (Elias, 1992). While this  
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figure is lower than the percentage of body mass accounted for by muscle (42% for a male, 36% for a female on average) (Marieb and Hoehn, 2010), suggesting that muscle may be a relatively cheap tissue at rest, it is the large size of some muscle groupings and high rates of energy consumption during activation that ensure muscles comprise a significant portion of the human energy budget (McArdle, Katch and Katch, 2001). Skeletal muscle is therefore an expensive tissue to maintain and thereby constrains the amount of energy available for competing physiological functions such as immune responses (Muehlenbein and Bribiescas, 2005). Thus, a highly muscular phenotype, which is dependent on the anabolic effects of testosterone (Bhasin et al., 1996; Tsai and Sapolsky, 1996), presents an energetic handicap.

Additionally, high levels of testosterone have a number of other detrimental health effects, including increased incidence of prostate cancer, oxygen radical production, reduced tissue and organ maintenance, and injury associated with aggressive confrontational behavior (Muehlenbein, 2006; Lassek and Gaulin, 2009). As such, testosterone is an effective mediator of quality signalling due to the significant metabolic costs of a muscular physique (Zahavi, 1975; Graffen, 1990; Andersson, 1994), immunomodulation (Muehlenbein and Bribiescas, 2005; Muehlenbein et al., 2006; Muehlenbein and Watts, 2010), and the negative health effects of high levels of the hormone.

The high energetic demands associated with muscle mass mean that during periods of energetic deficits there may be a suppression of testosterone levels. This leads to reduced somatic reproductive effort through decreasing muscle mass (Bribiescas, 2001) and frequency of behaviors associated with mating. This study has demonstrated a decrease in electrical impedance. Although this could be in part

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2  
3 due to reductions in hydration (Talluri *et al.*, 1999), it is suggested that a breakdown  
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5 of muscle tissue also contributed to these measurements (Friedl *et al.*, 1994).  
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8 Our findings of both increased innate immunity and decreased testosterone  
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10 levels in runners lends further support to the considerable literature on relationships  
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12 between testosterone and immune function (Bouman, 2005).  
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15 This study demonstrated an acute-level trade-off between reproduction and  
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17 survivorship in ultramarathon runners. Because this is a highly trained and physically  
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19 active group, these precise results may not be generalizable to the wider population.  
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21 Such conditioning may buffer our participants from any negative health  
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23 consequences of such strenuous exercise and nutritional imbalance. On the other  
24  
25 hand, it could be argued that endurance running has cross-cultural ecological  
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27 relevance due to its importance in activities such as persistence hunting, as  
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29 observed in the Kalarhari in Africa and the Tarahumara tribe of Northern Mexico  
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31 (Liebenberg, 2006; Pennington, 1963). Indeed, endurance running may have played  
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33 a role in our evolution (Carrier, 1984; Lieberman *et al.* 2006, Longman, 2015). We  
34  
35 therefore maintain that results in different populations should be broadly comparable,  
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37 given the universality of general life history theory predictions. Although we have  
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39 used two immune measures, we have measured only one arm of the immune  
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41 system, which is not indicative of total immunocompetence. Future avenues of  
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43 research should include adaptive immune measures and use similar approaches to  
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45 consider trade-offs between other life history traits (i.e., growth). In addition to  
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47 controlling for both core and environmental temperature, repeated post-race  
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49 measures at different intervals could also help distinguish the precise contribution of  
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51 muscle-produced immunological mediators on increased immune function.  
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**Author contributions**

DL, JTS and JCKW designed the study as a whole, with further input from IDS, and DL collected the data. MPM contributed financially to laboratory analyses, and oversaw the laboratory analyses performed by ECS and SPP. DL performed statistical analyses, and all authors edited the manuscript.

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## Appendix

Changes in general arousal that may be elicited by such prolonged exercise were controlled for, using the International Affective Picture System (IAPS) (Lang et al., 1988; Lang et al., 2005; Lang et al., 2008). The IAPS is a large collection of color photographs designed for distribution and research as standardized affective materials. The system includes normative ratings of each photograph with respect to valence (pleasure), arousal, and dominance.

A pilot study of 30 subjects (age- and sex-matched to race athletes) was conducted to collect SAM-rated valence and arousal data on 30 provocative near-nude female photographs. These photographs were obtained from a dating website, where visitors had given scores out of 10 to each image. The photographs chosen for the study had a mean score of 8.0 from a sample size of over 2000 website visitors. The pilot study enabled a determination of mean valence (6.45), arousal (5.11) and dominance (5.64). These thirty images were evenly split to make two groups, with equal mean valence, arousal, and dominance (see Table 2 below).

INSERT TABLE 2 ABOUT HERE

Thirty non-sexual positive IAPS images were then selected to match the pilot-derived arousal, valence, and dominance scores as closely as possible. These thirty images were split to make two groups of fifteen (Group A numbers: 1500, 1560, 2075, 2155, 2160, 8090, 5330, 5890, 8220, 7250, 7280, 7291, 7440, 7499, 8021; Group B numbers: 1540, 1590, 2152, 2158, 2224, 8208, 5628, 5990, 8467, 7279, 7289, 7402, 7450, 7660, 8060).

A further thirty non-sexual negative IAPS images were selected with matched arousal and dominance, but with a negative valence (3.39). Again, these thirty images were split to make two groups of fifteen (Group A numbers: 1220, 1274, 2055.1, 2120, 2683, 2692, 2710, 3181, 3212, 3300, 6021, 6200, 6561, 6836, 9101; Group B numbers: 1111, 1275, 2039, 2457, 2691, 2694, 2717, 3160, 3213, 3301, 6022, 6250.1, 6570, 6834, 9102). This design allowed the possibility of detecting any effects the exercise may have on the arousal valuations of both positive and negative pictures. Table 3 shows the average valence, arousal and dominance scores of each image subset.

Independent samples t-tests have confirmed that there were no significant differences between slide show A and slide show B in any of the three metrics (valence, arousal, or dominance). Furthermore, Levene's Test confirmed there were no significant differences in the variances of any variable.

INSERT TABLE 3 ABOUT HERE

For Peer Review

	Pre-race	Post-race	Comparison of pre- to post-race	
	Mean (SD)	Mean (SD)	t	p
Height (cm)	176.90 (6.69)	176.40 (6.53)	3.00	0.004
(n = 66)				
Weight (kg)	76.90 (8.58)	75.10 (8.26)	7.91	< 0.001
(n = 66)				
Testosterone	109.60 (17.60)	97.60 (17.50)	8.18	< 0.001
(pg/ml)				
(n = 52)				
Impedance	496.80 (58.90)	454.90 (50.9)	7.32	< 0.001
(n = 66)				
Valence Total	5.36 (0.69)	5.29 (0.69)	1.51	0.136
(n = 52)				
Arousal Total	5.29 (0.86)	5.04 (0.99)	3.55	0.001
(n = 52)				
Dominance Total	5.19 (0.65)	5.16 (0.67)	0.74	0.463
(n = 52)				
Haemolytic	124.6 (30.20)	139.40 (33.30)	2.43	0.021
complement assay				
(CH50) (n = 34)				
Bacteria killing	29.98 (36.70)	41.00 (39.60)	2.20	0.035

assay (% killing)

( $n = 34$ )

Table 1: Descriptive statistics for pre-race and post-race metrics, with independent t-test comparisons.

For Peer Review

Table 2: Average valence, arousal and dominance scores of each image subset.

	Prov A	Prov B	t (p)
Valence	6.49 (0.516)	6.41 (0.739)	0.315 (0.185)
Arousal	5.19 (0.712)	5.02 (0.641)	0.692 (0.420)
Dominance	5.47 (1.070)	5.81 (0.840)	-0.947 (0.192)



Table 3: Mean (standard deviation) valence, arousal and dominance for each image subset, independent t-test (p-value) reported no significant difference between the image subsets. The valence, arousal and dominance scores for the 2 subsets of provocative images (abbreviated below) were obtained through a pilot study.

	Prov A	Prov B	t (p)	IAPS+ A	IAPS+ B	t (p)	IAPS- A	IAPS- B	t (p)
Valence	6.49 (0.516)	6.41 (0.739)	0.315 (0.185)	6.54 (0.223)	6.56 (0.238)	-0.332 (0.742)	3.45 (0.431)	3.33 (0.607)	0.590 (0.560)
Arousal	5.19 (0.712)	5.02 (0.641)	0.692 (0.420)	5.15 (0.502)	5.07 (0.415)	0.503 (0.619)	5.10 (0.600)	5.20 (0.956)	-0.352 (0.727)
Dominance	5.47 (1.07)	5.81 (0.840)	-0.947 (.192)	5.65 (0.927)	6.01 (0.481)	-1.298 (0.205)	4.85 (0.552)	5.01 (0.753)	-0.675 (0.506)

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Percent Change

