

Cortisol and testosterone dynamics following exhaustive endurance exercise

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Abstract:

Purpose: Cortisol (C) and testosterone (T) are impacted significantly by prolonged endurance exercise with inverse responses. Increases in C are witnessed concurrently with decrements in T, possibly impacting recovery. This study was conducted to assess the dynamics of C and free T (fT) concentration and recovery time following an exhaustive endurance exercise session (EES).

Methods: 12 endurance-trained males ($X \pm SD$: VO_{2max} 66.3 ± 4.8 ml/kg/min, age 22.8 ± 3.1 years, body fat 11.0 ± 1.4 %, training 7.1 ± 3.2 years) completed a treadmill EES at ventilatory threshold (74.7 ± 4.6 % of VO_{2max} ; 96.9 ± 10.8 min). Basal blood C and fT were collected at baseline: -48 , -24 h, and immediately before (0 h) the EES as well as immediately ($+0$ h), $+24$ h, $+48$ h, and $+72$ h after the EES. Blood glucose (G) was measured to confirm no undue influence on C. Statistically data were analyzed with repeated measures ANOVA (LSD post hoc).

Results: C (nmol/L) increased significantly from -48 h (321 ± 59) to $+0$ h (701 ± 178) ($p < 0.001$), and displayed a baseline overshoot with $+24$ h (209 ± 67) being significantly lower than -48 and $+0$ h ($p < 0.03$). fT (pmol/L) decreased significantly from -48 h (161 ± 40) to $+0$ h (106 ± 38) ($p < 0.01$) and remained lower at $+24$ h (110 ± 33) and $+48$ h (129 ± 30) ($p < 0.001$). G remained stable throughout. A moderately negative correlation ($r = -0.636$, $p < 0.026$) was found between C and fT at $+0$ h.

Conclusions: EES recovery may require 48 h for C and 72 h for fT to return to baseline values. Furthermore, C and fT were only correlated immediately post-exercise. Future research should perform more frequent measurements throughout time course.

Keywords: Stress | Endocrine | Recovery | Overtraining

Article:

Abbreviations

<i>C</i>	Cortisol
<i>fT</i>	Free testosterone
<i>GnRH</i>	Gonadotropin releasing hormone

<i>Hb</i>	Hemoglobin
<i>Hct</i>	Hemocrit
<i>HPA axis</i>	Hypothalamic–pituitary–adrenal axis
<i>LH</i>	Luteinizing hormone
<i>REPANOVA</i>	Repeated analysis of variance
<i>RPE</i>	Rate of perceived exertion
<i>VO_{2max}</i>	Maximal volume of oxygen uptake
<i>VT</i>	Ventilatory threshold

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Introduction

Endurance running has been demonstrated to provide a large physiological stress to the body, resulting in large neuroendocrine system responses. Specifically, the hypothalamic–pituitary–adrenal (HPA) axis is stimulated to secrete the glucocorticoid hormone, cortisol. Cortisol has many physiological effects within the body that aid in substrate mobilization to improve running performance, but also affects the recovery process following an exercise bout (Brownlee et al. 2005; Daly et al. 2004). Testosterone, an anabolic–androgenic steroid that is secreted predominately from the Leydig cells of the testes, will tend to lower in concentration following exhaustive endurance exercise (Daly et al. 2005), reducing its anabolic affect within the body. These two hormones have long been studied as potential biomarkers for the catabolic or anabolic state of the body since they are highly involved with regulation of protein breakdown and synthesis (Hackney and Dobridge 2003). Due to the critical role these hormones play in protein turnover, it has been suggested they can be used to assess the recovery status of an athlete (Urhausen et al. 1995).

Although many studies have assessed the direction and magnitude of change in both of these hormones, most studies have assessed the concentrations either during an exercise bout or immediately following exercise completion. Some investigators have collected data on recovery processes, but most were only during the acute recovery phase (Hough et al. 2011; Thorpe and Sunderland 2012) or up to 24 h post-exercise (Daly et al. 2005; Duclos et al. 1997; Moore et al. 2005; Tremblay et al. 2005). Although this is valuable information, it is necessary to observe the concentrations of these hormones over a more extensive period of time to assess the time course required for the neuroendocrine system to return to a resting homeostatic state (Virus and Virus 2001). Such information is potentially vital to the coach and athlete in establishing timeline guidance for the number of rest days necessary after strenuous exercise, such as might occur in a competition.

Investigations that have observed hormonal concentrations for an extended duration following an exercise session have predominately used a field-based approach, in both team (Cunniffe et al. 2010; Elloumi et al. 2003; Ispirlidis et al. 2008; Kraemer et al. 2009) and individual (Lac and Berthon 2000) sporting competitions. Again, this provides important insight into the effects of competition recovery, but neglects potential variations that may exist following an exhaustive training session as can commonly occur amongst endurance athletes. Moreover, the lack of laboratory control of exercise intensity is a methodological limitation when assessing the true

time course of hormonal recovery. Therefore, the purpose of this study was to assess the dynamics of cortisol and testosterone concentration during recovery, following an exhaustive exercise bout in elite endurance runners.

Methods

Subjects

Twelve highly trained male endurance athletes volunteered to participate in this study. All subjects were free of any endocrine, musculoskeletal or cardiopulmonary system abnormalities or illnesses, and had been competing in and training for endurance-type activities for at least 5 days per week for the last 2 years. All subjects principally exercise trained by running; although, some also occasionally performed cycling or swimming. All subjects signed a written Informed Consent form, as approved by the Institutional Review Board at University of North Carolina at Chapel Hill.

Procedures

Subjects reported to the laboratory on seven separate occasions each time being required to adhere to the pre-assessment guidelines, which included being 2.5 h fasted, abstaining from strenuous activity, alcohol and sexual activity for 24 h, and abstaining from caffeine consumption for 12 h. At the initial session, subjects completed a written informed consent form, medical history questionnaire and training log, prior to anthropometric data (height, body mass, age) being recorded and undergoing a medical screening by a trained examiner, to ensure safety of participants in the study. Each subject then completed a modified Astrand maximal treadmill test to determine maximum oxygen uptake (VO_{2max}). Respiratory gases were collected via Truemax 2400 open-spirometry system (ParvoMedics Inc., UT) and VO_{2max} was determined by satisfying the following criteria: VO_2 did not increase with an increase by more than 0.15 l/min with an increase in workload, heart rate did not increase with an increase in workload, RPE > 18 (ACSM 2013). Ventilatory threshold (VT) was then calculated using the criteria of Wasserman (1984) and Yeh et al. (1983) to determine exercise intensity for the exhaustive exercise session (EES).

Following a 7-day wash-out period, participants returned to the lab for an initial blood draw, 48 h prior to the exhaustive exercise bout (-48 h). All subjects reported between 1300 and 1500 h, due to the relatively stable basal hormonal levels. First, height, body mass and body composition (skinfold measurements: chest, abdomen, thigh) were recorded. Subjects were then required to rest in a supine position for 30 min, prior to a blood sample being taken via the antecubital vein. This process was then repeated 24 h later (-24 h).

Subjects reported for the exhaustive exercise session having followed the pre-assessment guidelines. Subjects rested in a supine position for 30 min, prior to an indwelling, 20 gauge catheter being placed in the antecubital vein and a resting blood sample drawn (0 h). Subjects were fitted with a Polar heart rate (HR) monitor and were permitted time to warm-up and stretch. Subjects then completed a prolonged exercise run on the treadmill until volitional fatigue, running at 100 % of VT (within 3 %). Data were collected on VO_2 , HR, RPE, and %VT at 5, 30,

60, and 75 min into the exercise bout. Strong verbal encouragement was given to subjects towards the end of the exercise session to ensure a truly fatiguing exercise session. Subjects completed a ~5 min cool down, prior to a post-exercise blood sample being taken (+0 h). Subjects reported back to the laboratory at the same time of day 24 (+24 h), 48 (+48 h), and 72 (+72 h) hours after the exhaustive exercise bout, where blood samples were again drawn. Subjects were asked to maintain a similar diet and activity level compared to the pre-EES period during this 72 h of recovery, and the latter involved only activities of daily living and no exercise training.

Biochemical analysis

All blood samples were drawn in EDTA-treated tubes and placed on ice until processing. Whole blood aliquot samples were analyzed for hematocrit (Hct) and hemoglobin (Hb). Remaining sample aliquots were centrifuged at 4 °C for 15 min at 3000 rpm (Centra-8R IEC, MA). Separated plasma was frozen at -80 °C until hormonal analysis. Triplicate measurements of Hct and Hb were used to estimate plasma volume shifts during the fatiguing exercise bout (Dill and Costill 1974).

Hormonal concentrations of cortisol (C) and free testosterone (fT) were assessed in all samples, while glucose (G) was assessed in only -48, -24, 24, 48, 72 h, to exclude blood glucose changes as a confounding factor. All assessments were completed using highly specific, single-antibody sole phase radioimmunoassay kits that were specific for each hormone (DPC Inc., Los Angeles, CA, USA). All hormonal measurements were made with duplicate determination and were performed by highly trained technicians. Between and within coefficients of variation for all assays were less than 10 % for all biochemical analyses.

Statistics

Repeated measures analysis of variance (REPANOVA) over time was used to assess differences in cortisol, testosterone and glucose concentrations (SPSS version 21.0; IBM, Armonk, New York, USA). Fisher LSD post hoc procedures were used to identify significant differences between each level. Pearson-Product moment correlations were used to assess the relationship between cortisol and testosterone immediately following the EES (+0 h). Significance for all statistical tests was set a priori at $\alpha \leq 0.05$.

Results

Subjects all presented VO_{2max} values that confirmed they were highly trained aerobic endurance athletes (Table 1). The exercise session was successful in exhausting the athletes, and resulted in athletes running at 74.7 ± 4.6 % of VO_{2max} for 96.9 ± 10.8 min (details presented in Table 2). Means and standard deviations for fT, C, and G are presented in Table 3. All values reported were within clinically accepted normative values, and support the notion that our subjects were healthy and without endocrine disorders at the time of the study (Tietz 1995).

Table 1. VO_{2max} test ($X \pm SD$)

Age (years)	BMI (kg/m^2)	VO_{2max} (ml/kg/min)	VT (% of VO_{2max})	RPE (6–20 scale)	HR_{max} (bpm)	Running Speed (km/h)
22.8 ± 3.1	22.1 ± 0.8	66.3 ± 4.8	74.65 ± 4.6	19.3 ± 0.8	198.8 ± 4.7	17.8 ± 1.4

BMI body mass index, *VO_{2max}* maximal oxygen uptake, *VT* ventilatory threshold, *RPE* rate of perceived exertion, *HR_{max}* maximum HR at the end of the test, *running speed* maximal running speed

Table 2. Exhaustive exercise session ($X \pm SD$)

Time (minutes)	VO_2 (ml/kg/min)	HR (bpm)	RPE (6–20 scale)	VO_2 (% of VT)
5	47.6 ± 6.3	154.9 ± 3.8	11.6 ± 0.7	95.9 ± 2.9
30	49.8 ± 6.5	160.8 ± 4.2	13.1 ± 1.1	100.3 ± 2.7
60	51.2 ± 6.8	168.0 ± 5.2	15.1 ± 0.7	103.0 ± 3.0
75	52.7 ± 7.1	172.4 ± 5.5	16.6 ± 0.7	106.2 ± 3.2

VO₂ oxygen uptake, *HR* heart rate, *RPE* rate of perceived exertion, *VT* ventilatory threshold

Table 3. Testosterone, cortisol and glucose concentrations ($X \pm SD$)

	-48 h	-24 h	0 h	+0 h	+24 h	+48 h	+72 h
Testosterone (pmol/L)	161.3 (40.2)	158.6 (46.4)	167.1 (48)	106.1 (38.6)*	110.3 (33)*	129.8 (30.8)*	166.2 (30.0)
Cortisol (nmol/L)	321 (59.5)	326.75 (57.5)	331.8 (42.9)	700.8 (177.7)*	209.3 (67.1)*	309.3 (72.3)	340.5 (40.5)
Glucose (mg/dL)	79.3 (9.4)	83.4 (7.2)	78.4 (9.4)		77.3 (10.2)	82.1 (12.7)	78.4 (10.5)

*Significantly different ($p < 0.05$) compared to pre-value

Free testosterone

The REPANOVA and subsequent post hoc analysis revealed a significant decrease in fT following EES compared to pre-EES time points (-48 to 0 h; $p = 0.053$ to $p = 0.001$) that persisted for up to +48 h into recovery, before returning to baseline (pre-EES) levels at +72 h.

Cortisol

The REPANOVA and post hoc analysis revealed increases and decreases in C following EES compared to pre-EES. At +0 h, C was increased ($p < 0.001$) and decreased at +24 h ($p < 0.002$) before returning to baseline (pre-EES) levels.

Glucose

No differences were observed in glucose concentrations between any measurement times.

Correlations

The +0 h C and +0 h fT showed a significant moderate negative correlation ($r = -0.636$, $p = 0.026$), presented in Fig. 3.

fT:C ratio

The REPANOVA revealed significant differences with post hoc analysis showing fT:C was significantly decreased at +0 h (0.167 ± 0.084) relative to all other time points ($p < 0.005$). The fT:C returned to pre-EES values at +24 h, with no significant differences between +48 h or +72 h compared to any pre-EES values. +24 h and +48 h (0.579 ± 0.239 vs. 0.441 ± 0.153) did, however, approach significance ($p = 0.086$) relative to each other (power = 0.69, effect size 0.67).

Discussion

This study attempted to provide a more extensive understanding of the effects of exhaustive exercise on testosterone and cortisol concentration dynamics during recovery. The pre-exercise C and fT values were within normal ranges (Hellman et al. 1970; Zitzmann and Nieschlag 2001) and followed regular responses to exercise as previously observed in the literature (Kuoppasalmi et al. 1980).

The extended time frame with which these hormonal markers were assessed clearly demonstrate that 48–72 h of recovery may be required to return C and fT levels to resting levels following exhaustive exercise (Figs. 1, 2). This time to recovery is more extensive than has previously been observed, even in response to competitive rugby (38 h) and soccer (24 h) matches (Cunniffe et al. 2010; Ispirlidis et al. 2008), although some studies report an elevation in fT 6 days following a rugby competition (Elloumi et al. 2003). This elevation in testosterone is a curious finding relative to the 48 h of testosterone depression observed both in the present study and in triathletes (Urhausen and Kindermann 1987), and may be related to the high-intensity, intermittent nature of the sport of rugby.

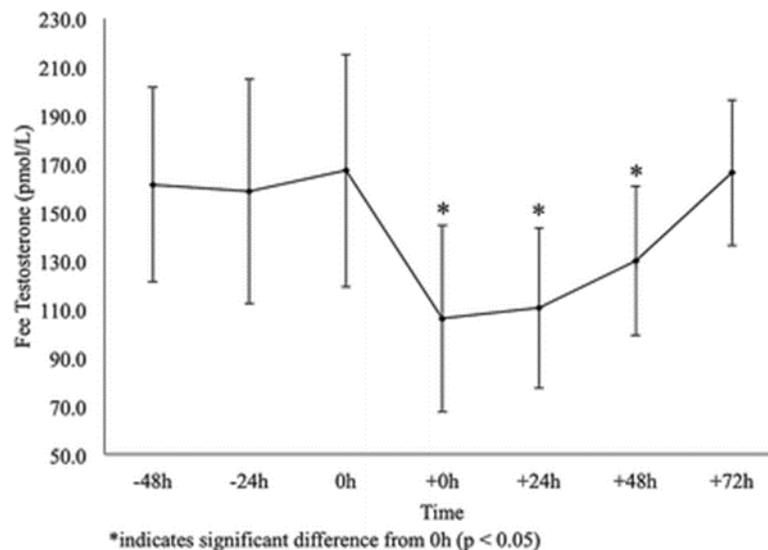


Fig. 1 Testosterone dynamics over the course of the study. The depicted values are mean \pm SD

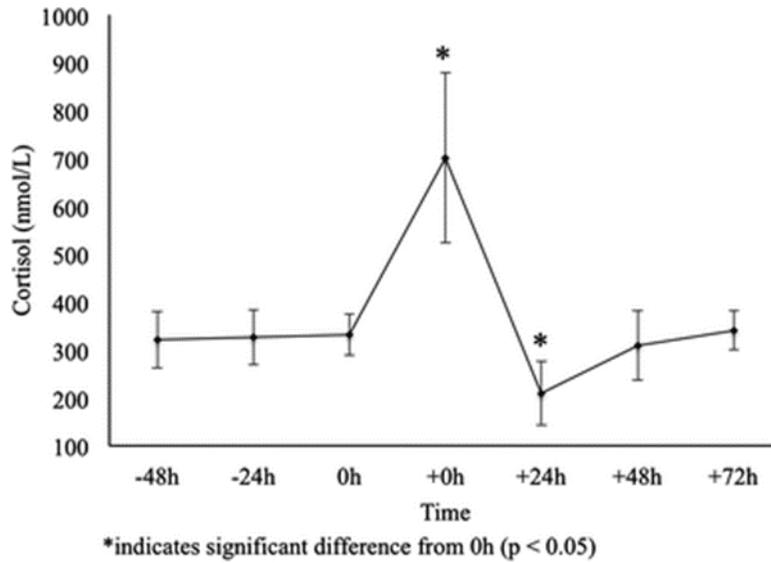


Fig. 2 Cortisol dynamics over the course of the study. The depicted values are mean \pm SD

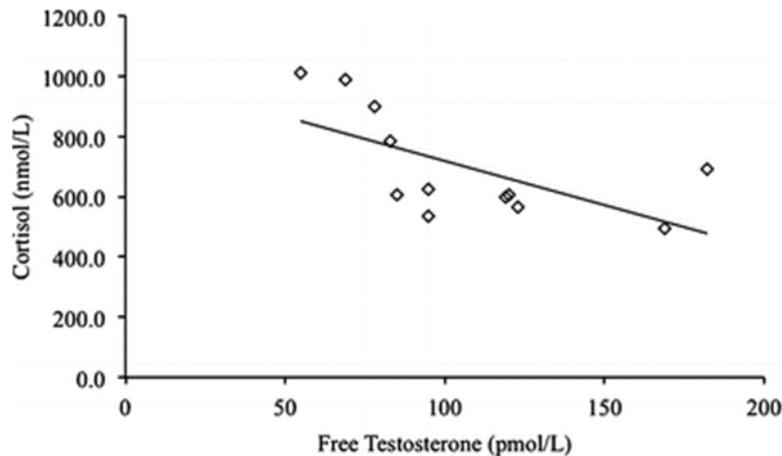


Fig. 3 The relationship between testosterone and cortisol immediately post-exhaustive exercise

Following exercise, the negative relationship between cortisol and testosterone (Fig. 3) suggests that the elevated cortisol levels may be acting as a testosterone suppressor. Indeed, this mechanism has been suggested since early investigations demonstrated that orally ingested cortisol resulted in testosterone suppression without changes in follicle-stimulating hormone or luteinizing hormone (Doerr and Pirke 1976). In agreement with the point, an investigation by Cumming et al. (1983) found that elevated levels of both endogenous and exogenous glucocorticoids resulted in the suppression of testosterone. Evidence suggests this may be due to direct glucocorticoid suppression of steroidogenesis at the Leydig cells (Daly et al. 2005), inhibition of luteinizing hormone (LH) and gonadotropic-releasing hormone (GnRH) (Hackney and Dobridge 2003) or inhibition of testicular luteinizing hormone receptors (Bambino and Hsueh 1981). Although elevations in glucocorticoids have been demonstrated to show an inverse relationship with fT (Bambino and Hsueh 1981; Kuhn et al. 1986), Daly et al. (2005) found either no relationship or a positive relationship between cortisol and fT in the 24 h following exhaustive exercise, even though a negative relationship existed for C and total T 60 min into

recovery. This was suggested to possibly be a result of concurrent release of C and fT from the adrenal cortex, or as a result of plasma volume shifts (Daly et al. 2005).

The present results suggest that cortisol returns to baseline more rapidly than testosterone (48 vs. 72 h). In fact, following the large elevations in cortisol concentration immediately post-exercise, the subsequent 24 h are characterized by a decrease in cortisol that overshoots baseline values and must then gradually increase to return to baseline levels. In agreement with our findings, Kraemer et al. (2009) found cortisol levels in American football players to be depressed for 44–48 h following a collegiate match, a similar pattern as that observed in other exercise stressors such as wrestling (Passelergue and Lac 1999), and rugby (Cunniffe et al. 2010; Elloumi et al. 2003). This cortisol depression has been suggested to be a result of an enhanced anabolic “rebound” stimulus during recovery (Cunniffe et al. 2010). However, it may also be a result of nutritional status of the athlete prior to and following the exercise stimulus. Future studies should be aware of carbohydrate intakes during these periods, as this will be a major influence on endocrine responses, even when sampling occurs at consistent time points (Lane et al. 2010). The rapid decrease in cortisol concentration is in contrast to testosterone concentrations which present a much more gradual increase back to baseline values, following the post-exercise suppression. The decoupling of the cortisol and testosterone relationship following +0 h, however, is possibly due to the fast response of the HPA axis and the short half-life of cortisol (Few 1974), resulting in increased cortisol clearances; whereas the hypothalamic–pituitary–gonadal axis requires extended time to adequately respond and stimulate gonadal testosterone production following glucocorticoid suppression of GnRH and LH.

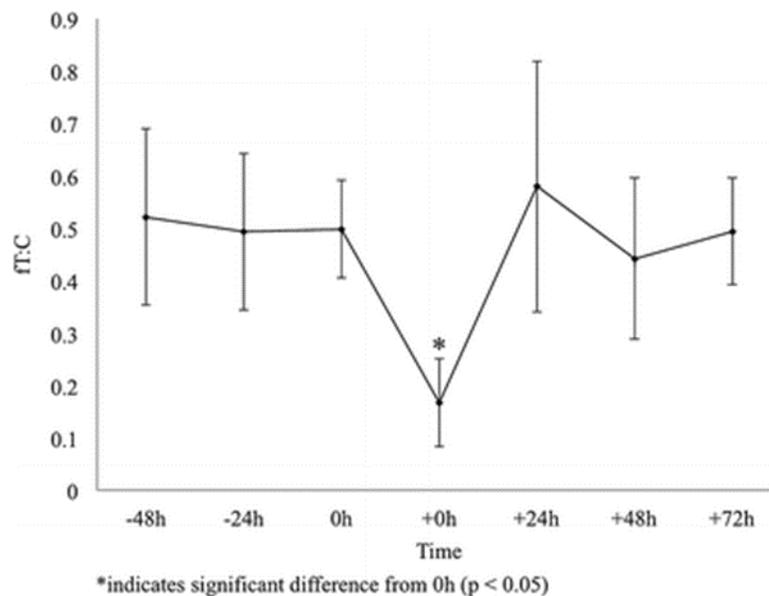


Fig. 4 Free testosterone to cortisol ratio over the course of the study

This difference in individual hormone recovery dynamics is a likely reason for the intriguing values observed in the fT:C during recovery (Fig. 4). Although not significant, the fT:C suggests an oscillating pattern in the 48 h post-exercise, prior to returning to pre-exercise values. Coupled with the variation in cortisol and testosterone recovery dynamics, these results support the

hypothesis that a negative relationship only exists between these hormones when cortisol is significantly elevated (>160 % of baseline values) (Brownlee et al. 2005). Our results show that cortisol concentrations are significantly reduced below baseline resting values within 24 h of recovery from an exhaustive exercise bout and although further research is needed to confirm this finding, these results challenge the notion that fT:C is a useful metric for monitoring the anabolic/catabolic status of an athlete during recovery as has been previously suggested (Banfi and Dolci 2006), especially during the acute recovery phase.

The present study was limited by the frequency of sample collection; future studies should employ more frequent blood sampling so the exact nature of the fT:C dynamics could be more appropriately categorized. In addition, the present study included only 12 male subject endurance athletes and limited biomarker analysis. The sample size clearly affected the statistical power of our fT:C ratio analysis (postmortem power calculation = 0.69; effect size = 0.66). To reach a statistical significant finding, a calculated sample size of 16 would be required, thereby resulting in data for fT:C only trending towards significance (i.e., +24 h and +48 h). Future studies should also expand on these results by analyzing additional neuroendocrine biomarkers (e.g., prolactin, growth hormone, catecholamines), specifically in women and those of varying fitness levels undertaking varied training modalities.

Conclusion

To summarize and conclude, following an exhaustive exercise bout, 48 h of recovery may be required for cortisol to return to baseline values and testosterone may require up to 72 h of recovery. Furthermore, cortisol and testosterone appear to be only correlated immediately post-exercise, although the ratio of these two hormones may follow an oscillating response, thus requiring future studies to perform more frequent measurements throughout the same time course. These findings support the notion that following strenuous exercise, similar to a prolonged competitive event, several days of rest may be necessary to re-establish a complete hormonal equilibrium.

Notes

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Compliance with ethical standards

Conflict of interest. The authors declare no conflicts of interest.

Ethical approval. All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethics standards.

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