Markers of muscle damage and strength performance in professional football (soccer) players during the competitive period

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Background: The present study investigated the impact of competitive soccer on the short-term changes in isometric strength of the adductor muscle group during the competitive season.

Methods: In this cohort study we evaluated the association between a serum marker of muscle damage [creatine phosphokinase (CPK)] and isometric strength of the adductor muscles of the hip in 30 professional football players (age: 26.7±2.9 years) during two seasons of the national top-level championship. Serum CPK level was determined the day before the match, 12–20, 36–48, 60–72 h after the match. The maximum voluntary isometric contraction force of the adductor muscles complex was determined immediately after having taken blood samples.

Results: There was evidence of a statistically significant positive association between age, body mass index (BMI), percentage of body fat, and muscle strength, and between weight and muscle strength. There was evidence of a statistically significant negative association between the level of CPK and the maximum isometric strength of the adductors of soccer players. Changes in CPK levels were associated with the muscle strength recovery trend (P<0.001). The strength/CPK ratio at different time points had a U-shaped curve.

Conclusions: Exercise induced muscle damage significantly affects the strength of the adductor muscle group of professional soccer players during the competitive period. The lower the CPK level, the greater the athletes' strength at a given time point. Also, the greater the decrease in CPK level, the greater the rate of strength restoration.

Keywords: Creatine kinase; muscle strength; adductors; isometric contraction; football

Submitted May 27, 2020. Accepted for publication Oct 25, 2020. doi: 10.21037/atm-20-2923 View this article at: http://dx.doi.org/10.21037/atm-20-2923

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Introduction

During a match, football players perform a multitude of movements, involving muscle contractions which eventually may cause muscle damage (1-3), with release of creatine phosphokinase (CPK) and other markers of such damage [for example, myoglobin and lactic dehydrogenase (LDH)] in the bloodstream (4-7). Exercise-induced damage to sarcomeres, cytoskeleton and sarcolemma is accompanied by transient muscle strength loss, and leads to delayed onset muscle soreness (DOMS) (8).

The reliability of serum CPK levels as a marker of muscle damage is debated (9,10). Some ethnic groups exhibit naturally increased levels of serum CPK following physical activity: this is not in any way pathological (6). Higher than normal levels of tissue CPK activity may increase the availability of cellular energy and improve myofibril contraction responses. Therefore, high levels of serum CPK, in the absence of muscle damage or other pathological conditions, may reflect the level of enzyme tissue activity. In such individuals, the serum CPK may be higher (6,11).

Serum CPK level is a marker of muscle microtrauma after exercise (12-14), and has been assessed during the competitive season in professional soccer players (15,16), using point-of-care testing (1,15,16). In some studies, CPK was used as an indicator of muscle fatigue, and there is an evident association between the level of CPK and very high-intensity running during the match (17,18).

In professional football (soccer), sports medicine professionals need objective and easy-to-perform tests to assess changes in strength in relation to the effects of exercise induced muscle damage. One of these tests is the five-second squeeze test, which is a valid indicator of sportsrelated hip and groin function (19).

In professional soccer, lower limb strength training produces measurable performance benefits. However, from a performance development view point, specifically considering fatigue and injury prevention, the advantages of increased lower body strength have received limited attention at the elite level of the game (20).

Pain associated with adductors muscles is the most common clinical finding on palpation and isometric muscle contraction in professional athletes, and can be a serious problem affecting the functional state and physical performance of the players. Weakness of the adductor muscles is associated with this clinical manifestation (21-23).

Every second football player suffers from pain in the

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adductor muscles during one football season, and every third football player begins the competition season with discomfort in the area close to the attachment of the adductor muscles to the pelvis, while a third of the players have residual discomfort dating back from the previous competitive season (24-26).

The present study investigated the impact of competitive soccer on the short-term changes in isometric strength of the adductor muscle group during the competitive season.

We analysed the association of CPK level to a series of anthropometric variables, and evaluated how this marker of muscle damage is associated with changes in the ability of the hip adductor muscle group to produce isometric strength. We wished to test the null hypothesis that there was no association between serum level of CPK and changes in the isometric strength of the hip adductor muscle group.

The aim of the study was to develop a relatively simple method to monitor the functional state of football players, and potentially to find a method to check and evaluate the adaptation to exercise and recovery in professional football.

We present the study in accordance with the STROBE reporting checklist (available at http://dx.doi.org/10.21037/ atm-20-2923).

Methods

All the procedures described in the present study were approved by our Local Ethics Committee (Sechenov Moscow State Medical University; Protocol No. 11-19, special meeting on 25.07.2019), and all participants signed a written informed consent. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

All athletes participating in the present study satisfied the following requirements:

- Participation in pre-season training camp;
- CPK level determination immediately prior to the beginning of the competitive season;
- CPK level determination at each of the 3 time points after three games, each with a minimum of 75-minute playing time;
- Identical physical loads in the training cycle (one training session every day 60–75 min);
- No adductor muscle injury 3 months prior to the first measurement.

Clinical and laboratory monitoring

In this cohort study, during two continuous seasons of

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the Premier League Championship (2017-2018 and 2018-2019), 30 professional male soccer players, all from the same professional team, were recruited (mean age: 26.7±2.9 years; height: 181.0±5.5 cm, body mass: 75.7±8.1 kg; percentage body fat: 10.1%±1.3%). No player was lost to follow-up. All the players were white Caucasians, and were members of their national soccer teams, played in the Russian Premier League, UEFA Europa League and Champions League. A finger-prick fasting blood sample allowed to analyse 32 µL of capillary blood to measure CPK concentration at five different time points: (I) before starting pre-season training after 2 or 3 weeks of rest; (II) 24 h before the match; (III) 12-20 h; (IV) 36-48 h; (V) 60–72 h after playing three matches during the competitive season (July-December, March-June in seasons of 2017/18, 2018/19). A total of 390 samples were taken. The Roche test-strips and the Roche Reflotron[®] Plus clinical chemistry system (Roche Diagnostics, F. Hoffmann-La Rosche, Basel, Switzerland) were used (15,27). The middle finger of the non-dominant hand of each player was wiped with an antiseptic solution and punctured with an automatic lancet. Blood was aspirated into the heparinized glass tube with an automatic pipette (preset volume 32 µL), and immediately pipetted onto the test strip. The measurement is based on the principle of reflectance photometry. The Reflotron[®] Quality Control (QC) Kit should be used with the Reflotron[®] test strips to ensure that the strips were working correctly and gave accurate and reliable results. The Roche Reflotron® Precinorm U QC kit was used for calibration procedures, which were performed: with each new lot of test strips; with each new shipment of test strip; whenever there was any concern that the test result may be wrong (test strip not stored correctly; when there was concern that a user may not have performed the test correctly); and every 30 days regardless of any other calibration procedures.

Strength testing

After the fingertip blood sample had been taken, the maximal voluntary isometric strength of the hip adductor muscle group was assessed. With the players lying supine, with their hands along the body, each player was asked to produce a 5 seconds maximal isometric contraction in three different positions, namely at 0°, 45° and 90° of hip flexion with 3 minutes rest between each maximal contraction (*Figure 1*). The Smart Groin Trainer dynamometer (NeuroExcellence, Braga, Portugal) was used for the measurement (in kg) (28). The dynamometer was placed

between the medial femoral condyles, enabling contraction of the pectineus, gracilis, short and long hip adductor muscles, and eliminating the influence of other muscles. Each player performed three attempts at each position, and the average per each position was used for statistical analysis. Before the test, each player warmed up performing passive resistance contraction and passive manual stretching of the adductors.

Anthropometric testing

Body composition analysis was performed using the "InBody720" body composition analyser, which employs bioelectric impedance (InBody CO, Seoul, Korea) (29,30). The analysis was performed the morning after a training session after an overnight fast.

Measurement of physical activity during matches

All the measurements of physical activity during matches were carried out using the InStat[®] kinematic system (Moscow, Russia) (31).

Statistical analysis

Data were entered into Excel tables using a tablet computer. Statistical analysis was performed using IBM SPSS Statistics software, version 23.0. The Wilcoxon signed-rank test was used to compare independent samples. Spearman's rank correlation coefficient has been used for correlation analysis. P<0.05 was considered statistically significant. Individual coefficients of changes in serum levels were calculated: CPK_PRE was divided by CPK_BAS, CPK_ POST1 by CPK_PRE, CPK_POST2 by CPK_POST1, CPK_POST by CPK_POST2. This allowed to normalise for the individual variability in serum CPK levels, and calculate their increase rate.

Results

The results of the measurements at all time points (24 h before the match, and 12–20, 36–48, and 60–72 h after the match) in three matches during the first 1/3, second 1/3 and third 1/3 of the competitive season were compared using the Wilcoxon signed-rank test. No significant difference was detected (P=0.39), and therefore measurements from the first match were used for further analysis for all athletes.

The mean and standard deviation (SD) pre-match



Figure 1 The adductor squeeze test at 0° (A), 45° (B) and 90° (C) of hip flexion.

strength values were: 0°, 53.2±7.4 kg; 45°, 38.9±7.5 kg; 90°, 45.7±7.3 kg.

Measurements of physical activity during each match were carried out in each player. No statistically significant differences in the level of physical activity were found. The average value of the team total distance at the speed of more than 19.8 km/h during all matches was 8,111±315 m, with the average number of high-intensity activities at the speed of more than 19.8 km/h during all matches being 615.5±49.0.

There was no evidence of a statistically significant association between the age of the players, body mass index (BMI), body fat percentage and maximal isometric strength of hip adductor muscle group in the three different hip flexion positions. There was evidence of a statistically significant positive association between the height and the body mass of the players and the maximal isometric strength of the hip adductor muscle group in each of the flexion position in all measurements (*Table 1*).

There was no evidence of a statistically significant association between serum CPK levels and BMI, percentage of body fat, and body mass. There was evidence of a statistically significant positive association between baseline CPK concentration, 24 and 48 h post-game CPK levels and players' age (*Figure 2, Table 2*). Interestingly, the serum CPK level at 72 h was associated with height.

The association between serum CPK level and maximal isometric strength of the hip adductor muscle group in three different hip flexion positions was studied. Serum CPK levels varied significantly in different players, and increased with the age of the athlete, as shown above, and also varied inter-individually. There was evidence of a statistically significant negative association between serum CPK levels and maximal isometric strength of the hip adductor muscle group at 0° (P<0.001; r=-0.418), 45° (P=0.06; r=-0.202) and 90° flexion positions (P=0.001; r=-0.363) (*Figure 3*).

Moreover, the changes in serum CPK level were associated with the rate of restoration of isometric strength: the lower the serum CPK level after the match, the higher the strength restoration rate (*Figure 4*).

There was a U-shaped CPK concentration/isometric strength ratio trend. A significant decrease was observed for the POST1 measurements 24 h after the match (P<0.001). 72 h after the match, the ratio has returned to basal values (P>0.05) (*Figure 5*).

Discussion

The present study evidenced a statistically significant negative association between serum CPK level and the

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Table 1 The relationship of age, height, weight, BMI, percentage of body fat with maximum isometric strength of the hip adductor muscle group (PRE: day before the match, POST1: 12–20 h after the match, POST2: 36–48 h, POST3: 60–72 h)

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Muscle strength (kg)	Age (years)	Body mass (kg)	Height (cm)	Fat %	BMI (kg/m ²)
PRE_0	P=0.35	R=0.38, P=0.04*	P=0.19	P=0.18	P=0.38
PRE_45	P=0.68	R=0.44, P=0.01*	R=0.42, P=0.02*	P=0.13	P=0.45
PRE_90	P=0.21	R=0.52, P=0.004*	R=0.33, P=0.07	P=0.15	P=0.31
POST1_0	P=0.53	R=0.32, P=0.08	P=0.25	R=-0.37, P=0.05*	P=0.43
POST1_45	P=0.68	R=0.41, P=0.02*	R=0.4, P=0.03*	R=-0.33, P=0.07	P=0.44
POST1_90	P=0.40	R=0.48, P=0.008*	R=0.35, P=0.06	R=-0.33, P=0.07	P=0.36
POST2_0	P=0.55	R=0.37, P=0.04*	P=0.17	P=0.13	P=0.33
POST2_45	P=0.67	R=0.43, P=0.02*	R=0.43, P=0.02*	R=-0.34, P=0.07	P=0.43
POST2_90	P=0.35	R=0.48, P=0.007*	R=0.36, P=0.05*	R=-0,33, P=0.08	P=0.32
POST3_0	P=0.40	R=0.35, P=0.05*	P=0.23	P=0.14	P=0.31
POST3_45	P=0.60	R=0.46, P=0.01	R=0.42, P=0.02	P=0.16	P=0.43
POST3_90	P=0.26	R=0.48, P=0.007	R=0.34, P=0.06	R=-0.31, P=0.09	P=0.37

Statistically significant results are indicated with *. BMI, body mass index.

Table 2 Relationship of age, height, weight, BMI, percentage of body fat with serum CPK level (BAS-control measurement before pre-seasontraining, PRE: on the day before the match, POST1: 12–20 h after the match, POST2: 36–48 h, POST3: 60–72 h)

CPK (U/L)	Age (years)	Height (cm)	Body weight (kg)	BMI (kg/m ²)	Percentage of body fat
CPK_BAS	R=0.50, P=0.006*	P=0.12	P=0.90	P=0.70	P=0.21
CPK_PRE	P=0.52	P=0.89	P=0.50	P=0.29	P=0.50
CPK_POST1	R=0.40, P=0.03*	P=0.25	P=0.24	P=0.65	P=0.99
CPK_POST2	R=0.42, P=0.02*	P=0.64	P=0.17	P=0.24	P=0.89
CPK_POST3	P=0.19	R=0.40, P=0.03*	P=0.42	P=0.25	P=0.26

Statistically significant results are indicated with *. BMI, body mass index; CPK, creatine phosphokinase.

maximal isometric strength of the hip adductor muscle group during the recovery period after a professional football match. In addition, the maximal isometric strength of the hip adductor muscle group in the professional football players under study showed evidence of a statistically significant association with body height and mass at all hip flexion positions. These findings carry valuable practical implications, as, considering these time dependent patterns of isometric strength changes, it will allow strength and conditioning coaches to better plan training during the competitive season.

A major strength of the present study is that this investigation was conducted on a cohort of elite professional male soccer players: this is a difficult task to accomplish during the competitive season of professional soccer. The results obtained are therefore a real reflection of the daily work in professional clubs.

We acknowledge that serum CPK levels alone may not accurately reflect the structural damage to muscle cells. Serum CPK levels can be affected by hydration status before eccentric exercise (9), hyperphosphatemia and hypermagnesemia (32), and may vary widely within subject groups of comparable male volunteers. Serum CPK levels may be also high from contraction-induced damage to the various muscles or following direct muscle injury.

Unfortunately, at present not all analytical methods and techniques are suitable for a quick, reliable and painless early diagnosis of muscle fatigue leading to muscle

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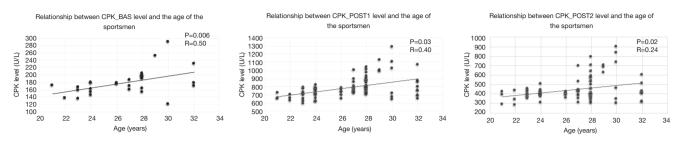


Figure 2 Relationship between the concentration of CPK and age of the soccer players. CPK, creatine phosphokinase.

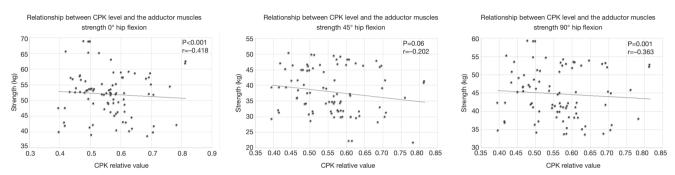


Figure 3 Relationship of the serum CPK level with the maximum isometric strength of the hip adductor muscle group (kg) at 0°, 45°, and 90° of hip flexion. CPK, creatine phosphokinase.

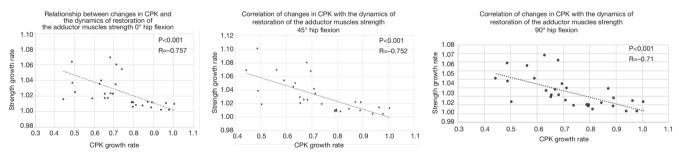


Figure 4 Relationship between changes in CPK and the kinetics of restoration of isometric strength of the adductor muscle strength at 0°, 45°, and 90°. CPK, creatine phosphokinase.

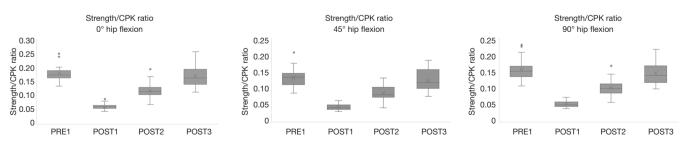


Figure 5 Strength/CPK ratio at different time points 0°, 45°, and 90°. CPK, creatine phosphokinase.

microdamage at the morphological level. In real life, such test diagnosis should be easy to perform without stopping from

the main activity. Muscle biopsies have revealed similar ultrastructure damage to the Z band of muscle fibres in the absence of differences in muscle soreness (33). In various studies, muscle biopsies have been performed to try and quantify muscle damage following exercise (6,34). However, by definition they only give a snapshot of what is happening in the small portion of the muscle investigated. Therefore, muscle biopsies may not give the full picture of the damage to the muscle group(s) exercised, and the biopsy procedure may itself cause damage to muscle fibres (35). Muscle biopsy is nevertheless a direct method to examine muscle tissue, but it is invasive, and requires specialized equipment and expertise.

Other additional indirect indices of muscle damage such as magnetic resonance imaging studies and ultrasound assessment are useful, but can only identify existing structural damage (36).

There is a number of biochemical serum markers of inflammation and stress, such as lipid hydroperoxides, malondialdehyde, myoglobin, lactate, alpha-tocopherol, retinol, uric acid, which can assist in quantifying and substantiating muscle disturbance parameters (37). However, most of these indicators are too indirect and their changes are multifactorial.

The present study focussed on performing routine, affordable assessment of serum CPK level and strength of the hip adductor muscle group: these muscles are involved in soccer-specific repetitive movements, such as side running, kicking, crossing, and curling the ball (38). The injury rate of this anatomical region is 1.015–1.133 injuries per 1,000 h playtime, and comprises 11–16% of all soccer-related traumatic injuries (23).

Computer-assisted isokinetic dynamometry is widely used to assess hip muscle strength in athletes (39). This equipment is expensive, requires specialised training and experience, and, given its size and weight, cannot readily be used for pitch side purposes. Hand-held dynamometers are easy to use and inexpensive, but their reliability can be a problem (40). The disadvantages of isokinetic and handheld dynamometers led us to employ the device used in the present investigation, which has the advantage of portability, is not dependent on the strength of the examiner, and is easily placed in an anatomically reproducible location over the extremities to test (41).

We chose the isometric type of contraction during

testing, because isometric strength testing exerts a lower load on the skeletal system in comparison to eccentric contractions, minimising pain and the risk of trauma (42). Thus, we believe that isometric testing is most beneficial for diagnostic and prophylactic use.

There is a significant correlation between isometric strength and isokinetic strength of a given muscle group (43). The results of the present study confirm that post-exercise CPK changes are closely associated to maximal isometric strength (44), and maximum isometric strength immediately after eccentric exercise is negatively associated with peak CPK activity (45). The present study identified a U-shaped CPK concentration/adductors isometric strength ratio curve. A significant decrease was observed for the POST1 measurements 24 h after the match:by 72 h after the match, the ratio had returned to basal values. Other studies evidenced a positive association between CPK peak activity and maximal isometric strength of lower extremity extensor muscles (46). The study of this biological behaviour can be used to assess the recovery of CPK level and adductors strength relative to each other during the competitive period, and to fine tune personalized training schemes during each microcycle.

Exercise modality can affect the appearance of CPK in blood serum. Following eccentric training, CPK serum levels can peak between 72 h (47) and 96 h (48) to 120 h (49). Training status may affect this time response. Eccentric resistance training elicits a significantly different increase in CPK serum levels in resistance trained and untrained men: serum levels of CPK peak at 24h in the resistance trained group, whilst they continue to raise and peak at 72 h in the untrained group (50).

We are aware that we report a snapshot within a competitive season. However, we point out that the results of our observations over three games at different stages of the competitive seasons were not significantly different, and therefore they likely reflect the actual response to the intense competition requirement of modern professional football. We are not sure that extending such observation to more games would allow to paint a different picture, and that it would impact on the practical implications delineated in the present study.

The question of the effect of the level of load on the increase in CPK could be the subject of future investigations in this field. We acknowledge that we did not study the pattern of motor activity and did not test physical qualities in the days of changes in physical performance and muscle strength: this could be incorporated in future

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experimentations. However, it should be kept in mind that we dealt with elite professional football players, for whom undertaking scientific research could be difficult and considered invasive of their performance capabilities.

We only studied the relationship between the serum level of CPK and muscle strength. Therefore, we could not ascertain how the level of CPK and a decrease in the strength of the adductors are related to the dynamics of various other parameters of physical performance, and how the level of daily physical activity can affect the indicators studied. This could be interpreted as a weakness of our work, and could be the subject of future endeavours.

The players taking part in the present study are all white Caucasian. This reflects the composition of the teams studied, and should prompt similar investigations in other ethnic groups, defining whether the findings of the present study are applicable to the whole population of elite professional football players.

Conclusions

Indirect muscle microtrauma after a professional soccer match significantly affects muscle strength parameters in professional soccer players during the competitive period. There is a strong negative association between serum CPK levels and adductor muscle strength—the faster the decrease in serum CPK levels, the faster the recovery of the isometric strength of the hip adductor muscle group. Serial serum CPK level measurement using finger prick sampling, combined with isometric dynamometry, is an easy and convenient method to check and evaluate the adaptation to exercise and recovery in professional football.

Acknowledgments

Funding: None.

Footnote

Reporting Checklist: The authors have completed the STROBE reporting checklist. Available at http://dx.doi. org/10.21037/atm-20-2923

Data Sharing Statement: Available at http://dx.doi. org/10.21037/atm-20-2923

Peer Review File: Available at http://dx.doi.org/10.21037/ atm-20-2923

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at: http://dx.doi. org/10.21037/atm-20-2923). The authors declare no conflict of interest.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. All the procedures described in the present study were approved by our Local Ethics Committee (Sechenov Moscow State Medical University; Protocol No. 11-19, special meeting on 25.07.2019), and all participants signed a written informed consent. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

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Cite this article as: Khaitin V, Bezuglov E, Lazarev A, Matveev S, Ivanova O, Maffulli N, Achkasov E. Markers of muscle damage and strength performance in professional football (soccer) players during the competitive period. Ann Transl Med 2021;9(2):113. doi: 10.21037/atm-20-2923

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