
An Intensive Winter Fixture Schedule Induces a Transient Fall in Salivary IgA in English Premier League Soccer Players

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An intensive winter fixture schedule induces a transient fall in salivary IgA in English Premier League soccer players

Submission Type: Short Communication

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Abstract

We examined the effects of an intensive fixture schedule on salivary IgA (SIgA) concentration in professional soccer players from the English Premier League. Salivary samples were obtained from 21 males who participated in 7 games over a 30-day period during December 2013 and January 2014 (games 1-5 occurred in a 15-day period). Salivary-IgA decreased (P < 0.05) at 2 days post-game 3 (45 ± 9 μg.mL⁻¹), 4 (52 ± 7 μg.mL⁻¹) and 5 (41 ± 10 μg.mL⁻¹) compared with game 1 (139 ± 25 μg.mL⁻¹). When the normal fixture schedule resumed (i.e. one game per week), SIgA returned towards baseline such that game 6 and 7 values were not different (P > 0.05) from game 1 (91 ± 18 and 99 ± 21 μg.mL⁻¹, respectively). Data demonstrate for the first time that a congested winter fixture schedule induces detectable perturbations to mucosal immunity in professional soccer players.

**Keywords:** football, illness, infection,
**Introduction**

The fixture schedule of competitive soccer games in the English Premier League (EPL) typically involves one game per week, usually on a Saturday-to-Saturday basis. In this way, the fitness and sports medicine staff is considered to have sufficient time to implement recovery strategies, manipulate training loads, and provide optimal energy intake to ensure players prepare appropriately for the subsequent game. In contrast, the winter period in the EPL is characterised by an intensive fixture schedule where players can play multiple games per week with limited recovery time. Such fixture schedules are especially prevalent over the Christmas and New Year period where it is not uncommon for players to participate in multiple games per week with only forty-eight hours recovery between consecutive games. This increase in workload may be particularly problematic given that susceptibility to illness and infection is exacerbated during the winter months (Orhant et al. 2010; Fricker et al. 2005).

In an attempt to monitor athlete wellbeing and assess susceptibility to infection, it is commonplace in endurance sports to monitor basal salivary immunoglobulin-A (SIgA; the predominant immunoglobulin expressed in mucosal fluids) during times of intensive training and competition (Gleeson et al. 1995). As a non-invasive assessment, routine monitoring of SIgA can be practically relevant given that it not only correlates with the onset of illness (Neville et al. 2008) but is also sensitive to physical stress (Gleeson and Pyne, 2000) and appears to vary in accordance with exercise intensity and duration (Mackinnon & Hooper, 1994; Mackinnon & Jenkins, 1993; Nieman et al. 2002). In relation to soccer-specific exercise, the stress of an acute training session induces a transient fall in SIgA that returns to basal levels within 18 hours (Fredericks et al. 2012). To the authors’ knowledge, however, no
researchers have yet examined the variation in basal SIgA levels in elite soccer players during a period of intensive competition that is characterised by multiple games per week.

The aim of the present study was to therefore monitor resting SIgA levels in professional soccer players during an intensive winter fixture schedule. We studied professional players from an EPL team over a thirty-two day period throughout December 2013 and January 2014 where seven games were completed (five of which occurred in a fifteen day period). We hypothesised that SIgA would display a significant decrease during the intensive fixture schedule but would return to basal levels when the normal schedule of one game per week was resumed.

Methods

Participants. Twenty-one soccer players from a professional EPL soccer team (age: 26 ± 4 years; body mass: 80.9 ± 7.4 kg; height: 1.83 ± 0.07 m) provided informed written consent to participate. The study was approved by the Ethical Committee of Liverpool John Moores University.

Salivary IgA Analysis. Samples were obtained two days after each game in the morning period (between 0900 and 1000 h and prior to breakfast). Given that soccer match play induces a reduction in SIgA levels that return to basal levels within 18 hours (Fredericks et al. 2012), we reasoned that collection of samples 2 days post-game would allow us to ascertain the effects of the acute suppression in SIgA levels from that associated with more chronic levels of stress. Participants were provided with an Oral Fluid Collector (OFC; IPRO Interactive, Oxfordshire, UK) consisting of a synthetic polymer based material on a polypropylene tube. The OFC contains a
volume adequacy indicator, providing a clear colour change (changing from white to dark blue) when 0.5 mL (± 20%) is collected. Participants were instructed to swallow any saliva present within the oral cavity before placing the collection device on top of the tongue. Salivary IgA concentration was analysed immediately upon collection by author 3 according to a real time lateral flow device in accordance with the manufacturer’s guidelines (IPRO Interactive, UK). This method has previously been validated against ELISA ($R^2 = 0.78$) in 208 samples collected from a cohort of EPL soccer players with a range of SIgA concentrations from 28 – 628 μg.mL$^{-1}$ (Dunbar et al. 2011).

**Training and match load.** Training load was monitored on all training days using global positioning system (GPS) technology (STATSports Technologies Ltd©, Viper Software 2.0.6.1, Ireland). Match performance data were collected using a multiple-camera computerised tracking system (Prozone Sports Ltd®, Leeds, UK). Four players reported soft tissue injuries during the course of data collection and no player reported illness or infection. An overview of training load and match load data is shown in Table 1.

**Statistical Analysis.** Changes in SIgA were analysed using a one-way repeated measures general linear model (GLM) (SPSS, version 17, Windows). Additionally, changes in SIgA between players who played >50% of the total minutes played in the first 5 games (i.e. the congested schedule) versus those who played <50% of total minutes were also assessed using two-way repeated measures GLM. Differences between specific time-points were located using Bonferonni post-hoc analysis. Finally, changes in match load data during the 7 games (from those outfield players who started each game) were assessed using a between groups GLM. Data are presented as means ±SD with P values <0.05 indicating statistical significance.
Results
Salivary IgA significantly changed (P < 0.001, \( \eta^2 = 0.232 \)) during the 32-day data collection period (see Figure 1A). Specifically, SIgA was significantly decreased on the sample points after game 3 (P = 0.001), 4 (P = 0.003) and 5 (P = 0.002) when compared with game 1. However, when the normal schedule of competition and training was resumed (i.e. one game per week), SIgA returned towards basal levels. As such, SIgA levels at 2 days post-game 6 and 7 was not different from game 1 values (P = 0.09 and 0.13, respectively). When players were classified into those who played >50 % minutes of the total minutes played in games in 1-5 versus those who played <50 % of total minutes, changes in SIgA were still apparent (P = 0.001), with no differences evident between groups (P = 0.79) (see Figure 1B). Total distance (P = 0.43), high-intensity distance (P = 0.16) and sprint distance (P = 0.09), as assessed in the starting outfield players, also did not differ between the 7 games (see Table 1).

Discussion
Confirming our hypothesis, we provide novel data by demonstrating that SIgA displays a significant decline in professional EPL soccer players during an intensive winter fixture schedule. Our data are practically relevant as they demonstrate that SIgA provides a non-invasive assessment that is sensitive to the changes in either the physical and/or psychological stress associated with a short-term period of intensive competition.

The stress of intense exercise typically induces a transient fall in SIgA levels that tend to return towards baseline levels within hours to days providing there is sufficient recovery and no infection is present (Walsh et al. 2011). Our data are in agreement with this assertion as we also show that SIgA at two days post-game 2 was not
significantly different to that observed at two days post-game 1. This result was not unexpected given that both games were played within a five day period and hence recovery time was likely sufficient. However, SlgA at two days post-games 3, 4 and 5 were significantly different from those values observed post-game 1 and 2. Such data therefore suggest that the scheduling of games in this way (i.e. 4 games in 11 days) is not appropriate for optimal recovery given that detectable perturbations to mucosal immunity were clearly evident. It is noteworthy, however, that indicators of physical match load did not change over the data collection period though interpretation of this data is complicated by the fact that squad rotation was practiced and also that 7 games is likely insufficient to make associations between physical performance and related physiological variables such as SlgA, in accordance with the high match-to-match variability apparent in elite level soccer (Gregson et al. 2010).

Interestingly, the decline in SlgA was evident in players that played >50% (n = 11) and <50 % (n = 10) of the total minutes played in games 1 to 5 (i.e. the congested fixture schedule). It is difficult to comment on the role of training load between games in modulating SlgA responses given that both cohorts of players performed different training loads in order to promote recovery in ‘starters’ but yet also promote match readiness for ‘non-starters’ (see Table 1). Nevertheless, these data suggest that decrements in SlgA may also be due to additional factors such as the seasonal time of year (i.e. winter) but perhaps also the travel stress and psychological stress associated with this intense period of competition. Indeed, all 21 players were part of the squad that travelled to each game.

The measurement of SlgA in sporting contexts is typically used as a marker of over-training (Mackinnon & Hooper, 1994; Gleeson et al. 1995) and to predict susceptibility to infections (Neville et al. 2008). Indeed, the latter authors observed
that a fall in SIgA to levels <40% of the individual’s baseline value (when considered healthy) indicates a 50% chance of contracting infection within 3 weeks. Although our data are limited in that our baseline sample was restricted to post-game 1 (therefore are not entirely representative of true basal levels obtained in pre-match days), we observed that a significant cohort of our sample presented with values less than 40% of basal on those samples collected post-game 3 (n = 14), 4 (n = 8) and 5 (n = 4). The collection of such data provided additional objective information to support the implementation of specific strategies (especially that of reduced training loads) to potentially minimise any risk of infection (Nieman & Bishop, 2006) and maintain player availability for team selection. Indeed, post-game 2 we initially observed reductions in SIgA (albeit not yet statistically significant), therefore all players underwent a reduction in training load compared to habitual loads typically experienced during a one game per week schedule, even for those ‘non-starter’ players (See Table 1). Although we observed no illness in our players during the data collection period, this was likely due to the resumption of the normal one game per week schedule following game 5.

In summary, we show for the first time that a congested winter fixture schedule is sufficient to induce detectable perturbations to mucosal immunity in professional EPL soccer players. Monitoring of SIgA during intense competition schedules may obviate the requirement to implement intervention strategies to help maintain athlete wellbeing and availability for team selection. Considering the field-based nature of the present study, obvious limitations include the inability to control environmental temperature and players’ energy intake etc. Nevertheless, the use of highly trained professional soccer players in a ‘real-world’ environment also strengthens the ecological validity of our data.
References


Table 1 – Overview of training load and match load data during the 32-day data collection period. Players were classified as those who started each game (STARTERS) and those who did not start (NON-STARTERS). H, denotes home game and A denotes away games. HSR denotes high speed running, > 19.8 km/h/ Sprinting denotes speeds >25km/h. *RECOVERY consisted of indoor activities, which usually included ice baths, massage and foam rolling.

Figure 1 – A) Salivary IgA concentration during the 32-day data collection period, as observed in all 21 players. Black bars represent game days and samples were collected on the morning of two days post-game. * denotes significant difference from day 3 i.e. game 1 values, P < 0.05. B) Salivary IgA concentration during the 32-day data collection period in those players who played >50% of the total minutes played in games 1 to 5 versus those players who played <50% of the total minutes.
Figure 1

(A)

(B)

- Players >50%
- Players <50%
<table>
<thead>
<tr>
<th>Day</th>
<th>Activity</th>
<th>STARTERS</th>
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<th>NON-STARTERS</th>
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<tbody>
<tr>
<td></td>
<td></td>
<td><strong>Duration (min)</strong></td>
<td><strong>Total Dist (m)</strong></td>
<td><strong>HSR (m)</strong></td>
<td><strong>Sprint Dist (m)</strong></td>
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<td>1</td>
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<td>90 ± 13</td>
<td>10476 ± 1657</td>
<td>677 ± 226</td>
<td>282 ± 166</td>
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<td>3</td>
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<td>2137 ± 248</td>
<td>11 ± 10</td>
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<td>6</td>
<td>Match (A)</td>
<td>92 ± 9</td>
<td>10841 ± 1280</td>
<td>845 ± 277</td>
<td>364 ± 145</td>
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<tr>
<td>8</td>
<td>Training</td>
<td>42 ± 6</td>
<td>1444 ± 395</td>
<td>7 ± 7</td>
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<td>9823 ± 1058</td>
<td>632 ± 184</td>
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<tr>
<td>11</td>
<td>Training</td>
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<td>508 ± 443</td>
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<td>Match (H)</td>
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<td>597 ± 222</td>
<td>197 ± 91</td>
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<td>14</td>
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<td>95 ± 9</td>
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<td>19 ± 21</td>
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<td>2690 ± 987</td>
<td>31 ± 23</td>
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<td>49 ± 26</td>
<td>3494 ± 1855</td>
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