Applications of near-infrared spectroscopy in “anaerobic” diagnostics – SmO₂ kinetics reflect PCr dephosphorylation and correlate with maximal lactate accumulation and maximal pedalling rate

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ABSTRACT: We investigated the relationship of the time-dependent behaviour of muscle oxygen saturation SmO₂(t), phosphagen energy supply W₄₃(t) and blood lactate accumulation ΔBLC(t) during a 60-s all-out cycling sprint and tested SmO₂(t) for correlations with the end of the fatigue-free state tₚₛ maximal pedalling rate PRₘₐₓ and maximal blood lactate accumulation rate vLₐₘₐₓ. Nine male elite track cyclists performed four maximal sprints (3, 8, 12, 60 s) on a cycle ergometer. Crank force and cadence were monitored continuously to determine PRₘₐₓ, tₚₛ based on force-velocity profiles. SmO₂ of the vastus lateralis muscle and respiratory gases were measured until the 30th minute after exercise. W₄₃ was calculated based on the fast component of the post-exercise oxygen uptake for each sprint. Before and for 30 minutes after each sprint, capillary blood samples were taken to determine the associated ΔBLC. Temporal changes of SmO₂, W₄₃, and ΔBLC were analysed via non-linear regression analysis. vLₐₘₐₓ was calculated based on ΔBLC(t) as the highest lactate accumulation rate. All models showed excellent quality (R² > 0.95). The time constant of SmO₂(t) τₘₐₓ = 2.93 ± 0.65 s was correlated with the time constant of W₄₃(t) τₚₛ = 3.23 ± 0.67 s (r = 0.790, p < 0.012), vLₐₘₐₓ = 0.95 ± 0.18 mmol·l⁻¹·s⁻¹ (r = 0.768, p < 0.017) and PRₘₐₓ = 299.51 ± 14.70 rpm (r = -0.670, p < 0.049). τₚₛ was correlated with τₘₐₓ = 0.885, p < 0.001). Our results show a time-dependent reflection of SmO₂ kinetics and phosphagen energy contribution during a 60-s maximal cycling sprint. A high vLₐₘₐₓ results in a reduction, a high PRₘₐₓ in an increase of the desaturation rate. The half-life of SmO₂ desaturation indicates the end of the fatigue-free state.


INTRODUCTION

As an approach to the assessment of training and performance, the kinetics of skeletal muscle oxygenation are being examined more and more extensively in the laboratory and the field [1]. The application of the Beer-Lambert law to near-infrared spectroscopy (NIRS) allows for non-invasive and convenient measurement of changes in local muscle tissue oxygenation [2–4]. The procedure has been validated for use during dynamic exercise in adults [5], so that the NIRS signal obtained during exercise can be used to estimate the relationship between local O₂ supply and demand (SmO₂) or to describe changes in tissue blood volume at the site of O₂ exchange via changes in total haemoglobin and myoglobin mass (thb) [6].

Since the recovery of phosphocreatine (PCr) levels in muscle following activity or ischaemia depends primarily on production of ATP by mitochondria, which, in turn, depends on oxygen availability [7–10], oxygen supply and demand as reflected in NIRS and the levels of high-energy phosphates are closely correlated [3, 4]. For example, in muscles at rest, a relationship between dephosphorylation of PCr and desaturation has been demonstrated. Utilizing NIRS and magnetic resonance spectroscopy (MRS) in combination, Hamakura and colleagues [11] observed that arterial occlusion resulted in an exponential decrease in SmO₂. Upon reaching desaturation, the level of PCr decreased linearly, a phenomenon proposed to be required for maintenance of ATP levels.

The observed delay between SmO₂ desaturation and the reduction in PCr levels in muscle under ischaemic conditions at rest results from the interaction of different systems of energy supply. At rest, energy requirements are usually met almost exclusively by mitochondrial respiration. Subsequent transport of this energy from the mitochondria to the sarcomere involves a potential gradient created by rephosphorylation of PCr [10, 12–14]. Under ischaemic conditions at rest, the level of PCr in muscle remains stable until decreasing availability of oxygen lowers the rate of mitochondrial energy

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production; thereafter, the rate of PCr dephosphorylation reflects the energy consumption.

Boushel et al. [15] reported a two-fold higher rate of SmO₂ desaturation during exercise than at rest under ischaemic conditions and attributed this to a greater demand for energy in combination with glycolysis. However, the exact relationship between SmO₂, the level of PCr and glycolysis during maximal dynamic exercise, e.g., maximal bouts of cycling, remains to be elucidated. Under such conditions, we would predict a close temporal correlation between the kinetics of SmO₂ desaturation and PCr dephosphorylation, which could be influenced by the rate of glycolysis.

During exercise, the rate of ATP resynthesis is determined by the phosphorylation of ADP by enzymatic dephosphorylation of PCr in combination with glycolytic and mitochondrial ATP production [12, 16]. For instance, during twitch contractions, loss of ATP is buffered through usage of PCr by creatine kinase to phosphorylate ADP [13, 17]. Following such a contraction, the level of PCr recovers more rapidly than during the final period of stimulation, indicating the involvement of different mechanisms [17]. Chung and coworkers [17] reported that during exercise the phosphorylation status is closely coupled to the rate of glycolysis and oxidative phosphorylation. Assuming unchanged levels of creatine kinase activity, repophosphorylation of PCr depends on these two systems and deficient repophosphorylation leads to compensatory increases in mitochondrial and glycolytic ATP production [12, 17]. When the rate of ATP consumption changes suddenly, glycolysis, which occurs in the sarcoplasm, contributes more rapidly to the ADP/ATP gradient than mitochondrial respiration does [9]. During the initial phase of muscle contraction, an increase in oxidative phosphorylation is limited until the cardiopulmonary system has adapted to supply the greater amount of oxygen in demand [19].

Consequently, during the first seconds of maximal exercise, the aerobic contribution to energy production is likely to be relatively small, and PCr is restored predominantly by glycolysis. As glycolysis is sensitive to pH and ATP hydrolysis during maximal exercise leads to a rapid reduction in muscle pH [20], the glycolytic flux may decline within a few seconds [12], whereas mitochondrial ATP synthesis increases with time [19]. In addition, acidic conditions also favour dissociation of O₂ [19].

When sprinting maximally, the PCr consumed is repophosphorylated by creatine kinase utilizing ATP produced by glycolysis and mitochondrial respiration. From the increase in the rate of glycolysis and associated production of lactate within seconds after the start of exercise [21, 22] it could be concluded that oxidation of lactate is likely to play a key role in energy equilibrium and, thereby, in SmO₂ kinetics. We hypothesize that during maximal sprint exercise, the kinetics of SmO₂ desaturation reflect those of PCr dephosphorylation and are correlated with the maximal rate of lactate accumulation vLa max as an indirect parameter of maximal glycolytic activity.

Maximal energy flow can only be generated until PCr stores are reduced to a maximum of half [23]. The time span for maximal
energy flow and maximal power output seems to be well reflected by the half-life of PCR dephosphorylation, which is a function of its time constant. In our recent work, we presented a new approach to determine the end of the time interval of maximal energy flow (fatigue-free time span) the timespan up to the first systematic deviation from the fatigue-free force-velocity (F/v) profile (\(t_b\)) [24]. Assuming a temporal overlap of the SmO\(_2\) desaturation and PCR dephosphorylation kinetics, a correlation between \(t_f\) and the time constant of SmO\(_2\) can be hypothesised.

Further assuming a correlation between the maximal rate of energy, the rate of PCR dephosphorylation and muscle fibre type composition [25], with maximal pedalling rate being dependent on the proportion of fast-twitch muscle fibres in the muscles involved [26], we also predict an increase in the rate of PCR dephosphorylation and of SmO\(_2\) desaturation with increasing pedalling rate. According to the model presented, direct measurement of SmO\(_2\) should provide a powerful approach to assessment of changes in the levels of high-energy phosphates and lactate with time.

**MATERIALS AND METHODS**

**Participants**

Nine male (21.1 ± 3.4 years, 184.9 ± 5.1 cm, 89.6 ± 6.4 kg, BFP: 13.4 ± 1.7%, \(\text{VO}_{2\text{max}}\): 53.9 ± 4.6 ml kg\(^{-1}\) min\(^{-1}\), \(P_{\text{max}}\): 1744 ± 111 W) (mean ± standard deviation) elite track cycling sprinters took part in this study. Since our study design required high-level neuromuscular and metabolic performance, only athletes who had demonstrated closely linear F/v profiles \((R^2 > 0.95)\) in previous tests and who had shown constant performance across all races per day in track cycling sprint events at international championships were included.

Participants were requested to refrain from consuming alcohol and from intense training and asked to maintain their normal drinking and eating habits 24 h prior to the experimental session. All participants provided written informed consent for participation in this study. The study was approved by the institute’s ethical committee and performed in accordance with the Declaration of Helsinki.

**Exercise protocol**

Each participant visited the laboratory once, completing four maximal sprints in a seated position on a cycle ergometer. As the testing protocol was similar to the daily training and competition among these track cyclists, who had also previously carried out similar performance tests in the laboratory, no additional test familiarisation was considered necessary. All participants used their own cycling shoes and pedals during these efforts. The ergometer settings resembled actual competition demands.

The participants warmed up before each sprint with 6 minutes of low-intensity cycling (1–1.5 W kg\(^{-1}\) body weight), followed by a 3-s maximal sprint. Participants rested passively for 10 minutes between warm-up and testing. Four maximal sprints (3, 8, 12, 60 s) were performed in an isokinetic mode at a cadence of 120 rpm on an SRM cycle ergometer (Schoberer Rad Messtechnik GmbH, Jülich, Germany) with a 9-kg flywheel, in 7th gear. With these settings, the pedal force decreased linearly during the initial acceleration phase and the target cadence of 120 rpm was reached after 3–4 s.

The participants were asked to accelerate as rapidly as possible from a simulated rolling start at approximately 20 rpm to 120 rpm at a self-selected time-point and to continue sprinting with maximal effort until the end of each sprint, with the investigators providing emphatic verbal encouragement throughout the entire test. 30 min after the 12-s sprint, each participant performed 6 s of maximal-cadence, low-resistance cycling (the motoric test) on an ergometer [24]. The settings enabled the athletes to reach a cadence of ≥ 160 rpm within the first 3 s, which generated additional data in the high-frequency cadence range. Combination of these data with the fatigue-free rate of pedalling during the acceleration phase of each sprint allowed more valid determination of the F/v profiles [27].

Tests were separated by approximately 2 hours, during which the athletes were asked to eat a light meal, corresponding to the individual’s calculated energy expenditure during the preceding test. The next test began when comparable levels of blood lactate and blood glucose were observed.

**Equipment and Measurements**

Data collection started from 3 minutes before the beginning of the standardized warm-up and ended 30 minutes after exercise, while participants remained seated, with as little movement as possible.

A commercially available continuous-wave NIRS device was used (Moxy Monitor; Fortiori Designs LLC, MN, USA) to continuously measure SmO\(_2\) and tHb (total mass of haemoglobin and myoglobin) of the vastus lateralis muscle (VL), as this muscle has been used most commonly in recent studies using NIRS in cyclists [1]. The sensors were fixed in place using medical adhesive tape (Hypafix; BSN Medical, DE) and covered with a compatible commercially available light shield to eliminate possible ambient light intrusion. The device uses four wavelengths (680, 720, 760 and 800 nm) to assess absorbency via the modified Beer-Lambert law resulting in a relative concentration of SmO\(_2\) as a percentage in the following equation: oxygenated Hb + Mb / oxygenated Hb + Mb + deoxygenated Hb + Mb = SmO\(_2\) [28]. The device detectors are spaced at 12.5 mm and 25 mm from the emitter. The sampling rate was set to 0.5 Hz, which samples the four wavelengths over 10 cycles for an averaged output every second and gathered using the Idiag Moxy software (Idiag AG, CH). With the emitter-detector spacing of 25 mm, a penetration depth of 15.0 mm can be expected (ibid.). The adipose tissue thickness (ATT) of the subjects at the measurement site was individually measured and none breached the upper threshold of 15.0 mm; therefore ATT interference is considered negligible.

Net crank torque (M) and angular velocity (\(\omega\)) were monitored continuously at a sample rate of 500 Hz with an SRM power meter (Schoberer Rad Messtechnik GmbH, Jülich, Germany). The average data from units of 10 Hz were used for calculations.
Before and directly after each sprint, as well as 1, 3, 5, 7, 10, 15, 20, 25 and 30 minutes after the 8- and 12 s-sprints, 20 µl of capillary blood was collected from the hyperaemic ear lobe for haemolysis and enzymatic-amperometric determination of lactate and glucose ( Biosen, EKF Diagnostics, Magdeburg, Germany). Since accumulation of blood lactate during the 3-s test was expected to be limited and the participants had already performed a considerable number of such samplings, capillary blood was only collected during the first, third, fifth and tenth minutes following this sprint to ensure sufficient blood flow, prevent sample contamination and enhance comfort.

Respiratory gases were measured continuously with a breath-by-breath portable gas analyser (Metamax 3B, Cortex, Leipzig/Germany). Prior to each test the gas analyser was calibrated using a known concentration of gases and a 3-litre syringe following the manufacturer’s recommendations.

Data processing

Raw data from the ergometer were obtained at a sample rate of 10 Hz. From these values, the mean tangential force F at both pedals, averaged over one revolution (taking into account individual crank length), as well as the corresponding pedalling rate, was derived.

To determine fatigue-free maximal pedalling rate, the fatigue-free force-velocity (F/v) profile was created based on the pedalling rate and corresponding mean crank force during the first 3–4 cycles of pedalling (≥ 3 s in duration, data point in the cadence range from 30 to 120 rpm) in each sprint, during which the crank force decreased linearly. As recommended by Dunst et al. [27, 29] 1 or 2 revolutions with very high pedalling rate (data point in the cadence range ≥ 160 rpm) attained during the first 3 s of the motoric test and its corresponding mean crank force were incorporated into these calculations. The data points with the best F/v relationship were selected. Since pedalling rate is directly proportional to the tangential speed of motion at the pedal, the F/v profile was based on pedalling rate (as specific movement velocity) and corresponding mean crank force F [29]. Parameters of the linear model function

\[ F(v) = a \cdot v + b \]  

were determined by linear regression analysis and characterise the fatigue-free relationship between mean pedal force (N) and specific movement velocity v (rpm) (ibid.). Maximal pedalling rate was calculated as \( PR_{\text{max}} = -b \cdot a^{-1} \).

With the onset of fatigue, reflected in reduced energy production by the main propulsion muscles, the mean pedal force decreases at each cadence so that the F/v profile at time t \( F(v,t) \) is lower than the fatigue-free maximum \( F(v) \). We defined the timespan up to the first systematic deviation from \( F(v) \) by the time-point \( t_F \), after which \( F(v,t) \) fell below and thereafter never returned to the fatigue-free maximal \( F(v) \) [24]:

\[ t_F = \min \{ t \in [s; T] | F(v,t) < F(v) \ \forall \ t \in [s; T]; t > t_F \}, \]

where \( s = 0 \) represents the initiation and \( T \) the termination of each performance test.

Parameters of the exponential model function of \( \text{SmO}_2(t) \) were determined by non-linear regression analysis based on the \( \text{SmO}_2 \) data of the participants’ 60-s-sprint [30]:

\[ \text{SmO}_2(t) = \text{SmO}_2^{\text{Base}} - \text{SmO}_2^{\text{A}} (1 - e^{-\frac{t - \text{TD}}{\tau}}) \]

with \( \text{SmO}_2^{\text{Base}} \) as the resting level before the start of a sprint, \( \text{SmO}_2^{\text{A}} \) as the amplitude of the \( \text{SmO}_2 \) drop and \( \tau_{\text{SmO}_2} \) as the time constant of desaturation.

For each sprint test, excess post-exercise oxygen consumption (\( \dot{\text{VO}}_{\text{EPOC}} \)) was determined by non-linear regression analysis using the following bi-exponential 4-parameter model to establish the phosphagen energy supply [31]:

\[ \dot{\text{VO}}_{\text{EPOC}}(t) = A_{\text{FC}} \cdot e^{-\frac{\text{TD}}{\tau_a}} + B_{\text{SC}} \cdot e^{-\frac{\text{TD}}{\tau_b}} + \dot{\text{VO}}_{\text{Base}} \]

where \( A_{\text{FC}} \) and \( B_{\text{SC}} \) represent the amplitudes of the fast component (associated with post-exercise resynthesis of creatine phosphate) and the slow component (associated with post-exercise lactate oxidation), \( \tau_a \) and \( \tau_b \) the corresponding time constants and \( \dot{\text{VO}}_{\text{Base}} \) the asymptotic resting oxygen uptake at time \( t \rightarrow \infty \).

For equations (3) and (4), a time delay (TD) was used to compensate for any delays in the measurement data.

Replenishment of high-energy phosphates was estimated based on the product of the amplitude \( A_{\text{FC}} \) and time constant \( \tau_s \) of the fast component (\( \dot{\text{VO}}_{\text{PPO}} = A_{\text{FC}} \cdot \tau_s \)). Phosphagen energy supply was calculated from the latter by [16, 32]:

\[ W_{\text{PCr}} = \dot{\text{VO}}_{\text{PCr}} \cdot CE \]

employing a caloric equivalent (CE) of 20.9 J ml⁻¹ [16].

Based on \( W_{\text{PCr}} \) of the single sprint tests, time-dependent energy contribution of the phosphagen energy system during the 60-s test was analysed with the following mono-exponential 2-parameter model by non-linear regression analysis:

\[ W_{\text{PCr}}(t) = W_{\text{PCrTOT}} \cdot (1 - e^{-\frac{t}{\tau_{\text{PCr}}}}) \]

where \( W_{\text{PCrTOT}} \) represents the maximum phosphatic energy supply at time \( t \rightarrow \infty \) and \( \tau_{\text{PCr}} \) represents the corresponding time constant.

Since the time-point when maximal blood lactate concentration occurs is unpredictable, the dynamics of blood lactate response were calculated by non-linear regression analysis using a 3-parameter model to determine maximal blood lactate concentration [33, 34]:
ΔBLC(t) = B(·(-1/k3)e^(-k3·t)+1/k4·e^(-k4·t))+C (8),

where B may represent the amplitude of lactate formation rate in the muscle, k3 and k4 represent the time constants of appearance and disappearance of lactate in the blood and C depicts the limit value of blood lactate accumulation during maximal exercise.

tauc reflects the time delay of the onset of lactate accumulation in the blood [16], so that ΔBLC (tauc)=0. Maximal rate of lactate accumulation (VLamax, mmol l^-1·s^-1) was calculated by:

\[ \frac{\Delta BLC(t)}{\tau_{SmO2}} \]

The kinetics of oxygen uptake during the 60 s-sprint were described by Barstow & Molé [37]:

\[ \dot{V}O_2(t) = \dot{V}O_{2a} \cdot \left(1 - e^{-t/\tau_{SmO2}} \right) + \dot{V}O_2(0) (10), \]

where \( \dot{V}O_2(t) \) represents the oxygen uptake at time t. \( \dot{V}O_2(0) \) is the initial oxygen uptake or starting value immediately before the start of exercise. \( \dot{V}O_{2a} \) defines the amplitude or denotes the change between the initial and current oxygen demand. To investigate the dependence of oxygen saturation on oxygen availability, the time-dependent quotient of changes in oxygen saturations (ΔSmO2(t) = SmO2(t)/SmO2(t + 1)100 [%]) and changes in oxygen uptake (ΔVO2(t) = VO2(t)/VO2(t + 1))100 [%]) was calculated.

**Statistical analyses**

All data were checked for normality using the Shapiro-Wilk test and are presented as mean ± SD. The Pearson product-moment correlation test was used for analysing interrelationships between variables. Pearson’s correlation coefficient r (small: r ≥ 0.1; medium: r ≥ 0.3; large: r ≥ 0.5) was employed as a measure of effect size. Statistical significance was set at an alpha level of < 0.05. Linear and nonlinear regression analysis was used to study the relationship between data. The least squares method was used in the regression analysis to determine the model parameters and time delays of the various model functions. The absolute difference was used to determine the bias of measurements. The quality of the regression analyses was examined by calculating the coefficient of determination. Mathematical analysis and statistical tests were processed using IBM SPSS Statistics version 24 for Windows (IBM Corp., Amonk, NY, USA), Office Excel 2016 (Microsoft Corporation, Redmond, WA, USA) and MATLAB 9.10.0 R2021a (The MathWorks, Inc., Natick, MA, USA).

**RESULTS**

Figure 1 illustrates the time-course of power output, oxygen uptake and blood lactate concentration in an exemplary 60-s test.

After the initial acceleration phase, the athletes reached the specified cadence of 120 rpm within the first 4 s in all sprint tests. To control the quality of the performance within the tests, individual fatigue-free F/v profiles were compared. No statistically significant differences were found.

Considering only the best F/v profile of each athlete (the profile with the highest calculated maximal power output), mean maximal force was 1343 ± 107 N and mean maximal cadence was 300 ± 15 rpm. The slope of the F/v profile was a = -4.51 ± 0.53 N rpm^-1. The coefficient of determination R^2 for the F/v profile was calculated to be ≥ 0.99 for all athletes. The time-point immediately before the first systematic deviation from the fatigue free F/v profile was located at t_f = 2.06 ± 0.32 s.

Mean power output, ΔBLC and the model parameters of EPOC(t) of the different sprint tests are shown in Table 1.

The average R^2 of all EPOC-models was 0.90 ± 0.03. The mean EPOC(t) of a 60-s test with the different components is illustrated in Figure 2.

Mean model parameters of oxygen desaturation (equation 3) of the VL were SmO2Base = 75.13 ± 8.60%, SmO2a = 65.94 ± 9.36% and τ_{SmO2} = 2.93 ± 0.65 s. An R^2 > 0.99 was calculated. The time-point at which modelled ΔSmO2(t) started to increase systematically above 100% and therefore SmO2 values differ from the model function was τ_{SmO2} = 32.60 ± 4.96 s. An example of SmO2 data and the determined model SmO2(t) of an athlete’s 60-s sprint test is shown in Figure 3.

TbH values ranged between 12.95 ± 0.33 mg dl^-1 and 12.74 ± 0.31 mg dl^-1 with 0.22 mg dl^-1 as the largest difference observed. The maximal value of TbH was reached at 24.8 ± 22.8 s, the minimal value at 15.3 ± 16.0 s, not indicating any time-dependent statistical differences.

Mean parameters of oxygen uptake kinetics were VO2a = 3.70 ± 0.44 l min^-1, τ_{VO2} = 14.73 ± 3.00 s and VO2(0) = 0.68 ± 0.20 l min^-1 with a mean R^2 of 0.85 ± 0.10.

The time-dependent quotient of changes in oxygen saturation and changes in oxygen uptake (ΔSmO2(t) ΔVO2(t)^-1) was >> 1 in the relevant period of exercise (t_{test} < t ≤ 60 s).

Based on WP, of the different sprint tests, the parameters of equation 6 to describe the time-dependent energy supply of the phosphagen energy system WP(t) were WP/Ctot = 53.02 ± 9.30 kJ and τ_{WP} = 3.23 ± 0.67 s with an R^2 of > 0.98.
Based on $\Delta BLC$ of the different sprint tests, the parameters of equation 8 describing the time-dependent blood lactate accumulation $\Delta BLC(t)$ were $B = 1.59 \pm 0.40$ mmol·l$^{-1}$·s$^{-1}$, $k_3 = 0.07 \pm 0.02$ s, $k_4 = 0.62 \pm 0.28$ s and $C = 18.78 \pm 2.99$ mmol·l$^{-1}$ with an $R^2$ of $> 0.99$. $t_{\text{totl}}$ was determined at 2.09 ± 0.41 s. Mean maximal rate of lactate accumulation was $v_{L_{\text{max}}}$ = 0.95 ± 0.18 mmol·l$^{-1}$·s$^{-1}$.

The mean kinetics of SmO$_2$, $W_{\text{PCr}}$, $\Delta BLC(t)$ and $La(t)$ in the 60-s sprint tests are shown in Figure 4. SmO$_{2A}$ showed a significant strong correlation with SmO$_{2B}$ (r = 0.911; p < 0.002). $\tau_{\text{SmO}_2}$ was positively correlated with $\tau_{\text{PCr}}$ (r = 0.790, p < 0.012), $v_{L_{\text{max}}}$ (r = 0.768, p < 0.017) and $t_{\text{totl}}$ (r = 0.822, p < 0.001). $t_{\text{Ff}}$ was positively correlated with $\tau_{\text{SmO}_2}$ (r = 0.885, p < 0.001) and $\tau_{\text{PCr}}$ (r = 0.781, p < 0.001).

**FIG. 1.** Example of the time-course of power output $P$ (W), oxygen consumption $VO_2$ (l·min$^{-1}$), and blood lactate concentration $BLC$ (mmol·l$^{-1}$) during the 60-s cycling sprint test protocol.

**TABLE 1.** Mean power output ($P_{\text{mean}}$), mean blood lactate accumulation ($\Delta BLC$), mean parameters of excess post exercise oxygen consumption EPOC(t) and mean phosphagen energy supply ($W_{\text{PCr}}$) of the 3-, 8-, 12- and 60-s sprint test (n = 9).

<table>
<thead>
<tr>
<th>Parameters</th>
<th>3 s</th>
<th>8 s</th>
<th>12 s</th>
<th>60 s</th>
</tr>
</thead>
<tbody>
<tr>
<td>$P_{\text{mean}}$ (W)</td>
<td>1151 ± 116</td>
<td>1395 ± 109</td>
<td>1371 ± 102</td>
<td>715 ± 42</td>
</tr>
<tr>
<td>$\Delta BLC$ (mmol·l$^{-1}$)</td>
<td>0.66 ± 0.25</td>
<td>4.73 ± 0.65</td>
<td>8.16 ± 0.90</td>
<td>17.70 ± 2.62</td>
</tr>
<tr>
<td>$A_{FC}$ (ml·min$^{-1}$)</td>
<td>1985 ± 436</td>
<td>2740 ± 381</td>
<td>2878 ± 501</td>
<td>3032 ± 447</td>
</tr>
<tr>
<td>$\tau_{a}$ (min)</td>
<td>0.82 ± 0.14</td>
<td>0.84 ± 0.12</td>
<td>0.84 ± 13</td>
<td>0.85 ± 13</td>
</tr>
<tr>
<td>$B_{SC}$ (ml·min$^{-1}$)</td>
<td>195 ± 167</td>
<td>418 ± 133</td>
<td>577 ± 119</td>
<td>981 ± 298</td>
</tr>
<tr>
<td>$\tau_{b}$ (min)</td>
<td>5.72 ± 2.52</td>
<td>6.46 ± 3.38</td>
<td>8.03 ± 3.10</td>
<td>9.36 ± 5.11</td>
</tr>
<tr>
<td>$VO_{2\text{Base}}$ (ml·min$^{-1}$)</td>
<td>391 ± 58</td>
<td>391 ± 43</td>
<td>411 ± 58</td>
<td>532 ± 106</td>
</tr>
<tr>
<td>$W_{\text{PCr}}$ (kJ)</td>
<td>31.83 ± 6.98</td>
<td>46.50 ± 3.79</td>
<td>51.24 ± 9.52</td>
<td>53.73 ± 9.96</td>
</tr>
</tbody>
</table>

**Abbrev.:** $P_{\text{mean}}$: Mean power output; $\Delta BLC$: Blood lactate accumulation; $A_{FC}$: Amplitude of the fast component of EPOC(t); $\tau_{a}$: Time constant of the fast component of EPOC(t); $B_{SC}$: Amplitude of the slow component of EPOC(t); $\tau_{b}$: Time constant of the slow component of EPOC(t); $VO_{2\text{Base}}$: Oxygen consumption at rest as the limit value of EPOC(t); $W_{\text{PCr}}$: Anaerobic alactic energy supply calculated based on the fast component of EPOC(t).
Insights into the physiology of sprint diagnostics with NIRS

DISCUSSION

In accordance with Hamaoka et al. [2, 11], Jones et al. [38], Hamaoka & McCully [39] and Vasquez-Bonilla et al. [40], our results demonstrate a significant correlation between desaturation behaviour of SmO$_2$ and the time-dependent change in PCR. In addition, significant correlations of SmO$_2$(t) with maximal pedalling rate and maximal lactate accumulation rate were found.

All models show excellent quality, qualifying them as suitable for describing the time-dependent behaviour of SmO$_2$, phosphagen energy supply (W$_{PCr}$(t)), blood lactate accumulation (ΔBLC(t)) and oxygen uptake (V$_{O2}$(t)) in an isokinetic 60-s all-out test on a cycle ergometer.

W$_{PCr}$(t) was described by a mono-exponential function that shows a steep increase in phosphagen energy contribution in the first seconds of maximal exercise. After approximately 10 s, the rate of phosphagen energy contribution is significantly reduced and at approximately 12 s W$_{PCr}$(t) reaches a quasi-steady state close to the maximum, which is mathematically reflected by the limit value of the function. According to the results of di Prampero and colleagues [41] the calculated amount of total W$_{PCr}$ of ~ 53 kJ, assuming an active muscle mass of circa 30% in cycling exercises [42], corresponds to about 26 mmol phosphagen split per kg active wet muscle. This value is in the same order of magnitude as the total phosphagen content of the resting muscle [16]. The time to maximum is in line with previous observations. Barclay [13] reported that during maximal exercise, the energy received from the phosphagen system peaks after 10–15 s. Therefore, a glycolytic and mitochondrial contribution sufficient to maintain a quasi-steady state of

p < 0.008). Furthermore, a negative correlation of τ$_{SmO2}$ and PR$_{max}$ (r = -0.670, p < 0.049) was found.

The associations between these parameters are shown in Figure 5.

Using the mean time constant of SmO$_2$(t) in combination with W$_{PCrTOT}$ to estimate W$_{PCr}$(t) results in a maximum error of 1.91 kJ. The results of the linear regression analysis comparing the two functions at each second during the 60-second sprint are shown in Figure 6.
re- and dephosphorylation of PCr in the working muscles is assumed at the time-point of near zero energy contribution of the phosphagen system. This state is also reflected in the desaturation behaviour of SmO\textsubscript{2} up to ~30 s.

Similar to \(W_{\text{PCr}}(t)\), \(SmO_2(t)\) can be described mono-exponentially and reflects the PCr dephosphorylation behaviour well. \(SmO_2(t)\) also shows a steep desaturation in the first few seconds of the test. After a subsequent decrease in the desaturation rate after approximately 10 s, \(SmO_2(t)\) also approaches a plateau after approximately 12 s. Unlike \(W_{\text{PCr}}(t)\), \(SmO_2(t)\) eventually increases slightly after the initial steep desaturation from the beginning of the test (see Figure 3). Maximal voluntary effort was ensured retrospectively by applying our previously published method [29]. Even though individuals continued their efforts at maximal voluntary intensity, relative \(SmO_2\) extraction decreases.

\[ y = 0.996x, \quad R^2 = 0.987, \quad \text{Bias} = 0.2550 \text{~kJ} \]
The correlation of the time constants $\tau_{\text{SmO}_2}$ and $\tau_{\text{PCr}}$ and the high agreement of the time courses indicate a close relationship between PCr dephosphorylation and SmO$_2$ desaturation in maximal cycling sprints. This statement is supported in contemporary exercise physiological literature [4, 42]. If the O$_2$ stored in the tissue primarily serves PCr resynthesis between twitch contractions during exercise, the SmO$_2$ desaturation rate should depend on the PCr dephosphorylation rate [7, 9]. This is in line with di Prampero [41] and Francescato and colleagues [44], who described a transfer of O$_2$ from Hb on Mb on the working muscle with the onset of exercise. The differences of the time constants calculated in this study could be explained by measurement inaccuracies (especially in determination of the fast component of EPOC, which represents the greatest uncertainty in this study), the difference between local and systemic measurement or by the additional inhibiting influence of high glycolytic activity on PCr dephosphorylation state, whereas the shift of the cell pH to acidic due to high ATP hydrolysis additionally favours O$_2$ dissociation [19].

Our results indicate that the time-point immediately before the first systematic deviation from the fatigue-free $Fv$ profile could reflect the half-life SmO$_2$ and $W_{\text{PCr}}$ and could indicate the onset of accumulation of lactate in the blood. Assuming an activation of glycolytic and oxidative ATP resynthesis after a depletion of 10–12 mmol PCr [16], which, applied to our data, corresponds to approximately 40–45%, the temporal proximity of these parameters seems plausible.

$\Delta \text{BLC}(t)$ was described by a bi-exponential function using $t_{\text{max}}$ to describe the delayed onset of blood lactate accumulation [41]. After a mean $t_{\text{max}}$ of 2.09 s, $\Delta \text{BLC}(t)$ rises steeply in an approximately linear manner until the tenth to twelfth second of exercise. After 10–12 s the increase in blood lactate accumulation is reduced, reflecting a sudden decrease of glycolytic activity caused by pH reduction in the active muscle cells [12, 20]. Extrapolated $\Delta \text{BLC}$ for 30 s of maximum effort is in accordance to the values reported by Beneke et al. [45] and Leithäuser et al. [46], which provides confirmation of model validity.

The onset of sudden significant reduction of lactate accumulation seems to coincide with the time-point when PCr dephosphorylation and SmO$_2$ desaturation approach their respective plateaus (see Figure 7). A positive correlation between the maximal lactate accumulation rate and the time constant of O$_2$ desaturation of the VL was observed. The higher $vL_{\text{max}}$, the higher $\tau_{\text{SmO}_2}$ and the longer it takes for the SmO$_2$ desaturation to reach a steady state. Considering the correlation of SmO$_2$ desaturation and PCr dephosphorylation, a higher $vL_{\text{max}}$ representing a higher rate of PCr replenishment via glycolysis between twitch contractions can be assumed, reducing the need for oxidative dephosphorylation of PCr due to a smaller initial alactic debt in the dominant working muscle.

Hautier et al. [26] and Hansen et al. [47] found a positive correlation in the proportion of fast glycolytic IIX muscle fibres in the VL with maximal pedalling rate. A significant correlation between the maximal pedalling rate and the time constant of SmO$_2$ desaturation was found. As fast contracting glycolytic muscle fibres have the highest ATPase activity, highest maximal energy flow rate and the fastest dephosphorylation of PCr [48, 49], the correlation between $\tau_{\text{SmO}_2}$ and $\tau_{\text{PCr}}$ as well as $PR_{\text{max}}$ and the muscle fibre spectrum seems plausible.

Although the potential for a high rate of glycolysis increases with the proportion of fast fibres [25], no correlation between $PR_{\text{max}}$ and $vL_{\text{max}}$ was found within this sample of elite power athletes. This result suggests that the specific enzyme activity required for high glycolytic energy flow is not exclusively morphologically determined, but represents a specific training adaptation.

The previously described sudden resaturation of SmO$_2$ in the VL after approximately 30 s of maximal exercise could be a multifactorial phenomenon reflecting the onset of a decrease in the active muscle fibre spectrum in the muscle, the delayed response of the cardiovascular system to rapid changes in peripheral energy demand and the haemodynamics of the capillary beds to maintain oxygen supply to the muscle.

Thomas and Victor [19] reported that a kind of contraction-induced sympathetic ischaemic state prevails in the muscles, which does not allow the influx of oxygenated blood for the very first seconds of maximal exercise. If this was the case, resaturation of SmO$_2$ in the working muscle would not be possible initially. With accumulation of metabolites in the interstitial fluid (K$^+$, lactate) [40], vasomotor relaxation occurs, mediated by the endogenous vasodilator nitric oxide (NO), attenuating sympathetic vasoconstriction in active muscles and contributing to greater hyperaemia during exercise [19]. Although such an effect cannot be excluded, our results do not indicate any significant reduction of blood flow and thus no restriction of oxygen availability during exercise due to temporary ischaemic states. Thus, a non-significant and time-related non-systematic change in $t\text{Hb}$ by a maximum of < 2% could be detected.

Although $V_{O_2}(t)$ increased exponentially from the initial value of ~15% to ~85% of the athletes' $V_{O_2\text{max}}$ during the 60-s sprint, oxygen uptake kinetics cannot explain the sudden resaturation of SmO$_2$ after ~ 30 s of exercise, which, at a nearly constant $t\text{Hb}$, has to represent a resaturation of Hb and Mb in the VL.

The $V_{O_2}(t)$ time constant ($t_{V_{O_2}}$ ~ 15 s) leads to a steep increase in oxygen uptake during the first 15 s of maximal sprinting. After 30 s of exercise, about 90% of the $V_{O_2\text{max}}$ was reached. In the second half of the sprint, oxygen uptake only increased by < 10% with increments of < 0.75% s$^{-1}$.

The relationship of resaturation rate and changes in oxygen uptake indicates a decoupling of resaturation and oxygen uptake in the second half of exercise ($\Delta\text{SmO}_2(t) > 100\%$, $\Delta V_{O_2}(t) < 100\%$ $\forall$ 30 < $t$ ≤ 60 s). From our results, we suggest that the resaturation during exercise is neither an effect of blood flow restriction nor a delayed response of the cardiovascular system to rapid changes in peripheral energy demand.

Considering the exponential decrease in maximal power output [29] and the reported exponential increase in blood lactate
concentration, it can be speculated that the progressive increase in resaturation of Hb and Mb from mid-exercise onwards may reflect the onset of rephosphorylation of PCr in fatigued high-glycolytic IIx fibres, which no longer contribute to power output [29, 50].

The assumption that resaturation during maximal exercise is an indication of a decrease in the number of working motor units in the dominant working muscles is also supported by the findings of Vasquez-Bonilla et al. [40]. In a repeated sprint protocol, a decrease in desaturation amplitude was observed in elite female football players with a simultaneous decrease in sprint performance in each bout. This is consistent with our observations of NIRS signalling during repeated cycling sprints.

Finally, Figure 7 illustrates the possible interpretation of SmO₂ kinetics based on the results of our study.

LIMITATIONS

The study is based on a considerable amount of mathematical modelling of physiological responses during maximal exercise. Despite the high quality of the individual models and plausibility of the overall modelling, the simplifying character of a model in favour of the identification of interrelationships and systematic behaviour has to be considered.

The models are based on data points derived from several maximal sprint tests. The validity of the models requires a high...
reliability of performance, metabolic stress and their measurements. Despite our efforts to standardise the tests and ensure similar test conditions, it must be pointed out that it is an indirect reconstruction of the metabolic response during the 60-s effort, so that misinterpretation cannot be excluded completely.

Due to the limited number of participants, elite training status and the possible sources of error presented, further investigations are required to check our results, especially in different cohorts. When recruiting participants, care must be taken to ensure a sufficient level of capability and proficiency, as the method used here requires the athletes to sustain maximal neuromuscular performance. If this performance is below the time-dependent maximum, metabolism in the fibres of the active muscles may vary, altering total metabolic response and its systematics.

PRACTICAL APPLICATIONS

For practical application, the $\text{SmO}_2$ kinetics of the muscle with the highest desaturation within the main propulsive/working muscles can be used as a representation for the PCr dephosphorylation state of the system.

By combining the value of total alactic energy derived from oxygen uptake of the fast component of EPOC with the time constant of $\text{SmO}_2(t)$ derived from NIRS measurements, PCr dephosphorylation kinetics can be estimated on the basis of a single sprint with an error $< 2$ kJ.

A direct measurement of the lactate accumulation rate based on NIRS measurements does not seem to be possible, but the time interval from the onset to the maximum activation of the lactate accumulation rate can be represented by NIRS measurements.

CONCLUSIONS

Our results indicate that $\text{SmO}_2$ kinetics of the dominant working muscle reflect the time-dependent phosphagen energy contribution and the dephosphorylation of high-energy phosphates in a 60-s maximal sprint in elite track cyclists. The rate of $\text{SmO}_2$ desaturation is dependent on the maximal pedalling rate and the maximal lactate accumulation rate reflecting the maximal glycolytic energy flux. While higher $\text{PR}_{\text{max}}$ accelerates desaturation, higher $\text{vLa}_{\text{max}}$ reduces it. The half-life of $\text{SmO}_2$ represents the end of the fatigue-free state and could indicate the beginning of accumulation of lactate in the blood. Reaching an $\text{SmO}_2$ plateau after maximal desaturation seems to be temporally associated with a reduction in lactate accumulation rate.

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Conflicts of interest/Competing interests

On behalf of all authors, the corresponding author states that they have no conflicts of interest.

Ethics approval

All procedures were approved by the ethics committee and conducted in accordance with the Declaration of Helsinki.

Authors’ contributions

AKD designed and conducted the study and analysed the data. AKD, CH, CM and AF were responsible for drafting the paper and revising it critically with respect to important intellectual content. AKD, CH, CM and AF approved the final version of this manuscript for publication.

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REFERENCES


Insights into the physiology of sprint diagnostics with NIRS


