Analysis of the aerobic-anaerobic transition in elite cyclists during incremental exercise with the use of electromyography

Alejandro Lucia, Oscar Sánchez, Alfredo Carvajal, José L Chicharro

Abstract

Objectives—To investigate the validity and reliability of surface electromyography (EMG) as a new non-invasive determinant of the metabolic response to incremental exercise in elite cyclists. The relation between EMG activity and other more conventional methods for analysing the aerobic-anaerobic transition such as blood lactate measurements (lactate threshold (LT) and onset of blood lactate accumulation (OBLA)) and ventilatory parameters (ventilatory thresholds 1 and 2 (VT1, VT2)) was studied.

Methods—Twenty eight elite road cyclists (age 24 (4) years; \( V_{\text{O}2\text{MAX}} \) 69.9 (6.4) ml/kg/min; values mean (SD)) were selected as subjects. Each of them performed a ramp protocol (starting at 0 W, with increases of 5 W every 12 seconds) on a cycle ergometer (validity study). In addition, 15 of them performed the same test twice (reliability study). During the tests, data on gas exchange and blood lactate levels were collected to determine VT1, VT2, LT, and OBLA. The root mean squares of EMG signals (rms-EMG) were recorded from both the vastus lateralis and the rectus femoris at each intensity using surface electrodes.

Results—A two threshold response was detected in the rms-EMG recordings from both muscles in 90% of subjects, with two breakpoints, EMGT1 and EMGT2, at around 60–70% and 80–90% of \( V_{\text{O}2\text{MAX}} \) respectively. The results of the reliability study showed no significant differences (p>0.05) between mean values of EMGT1 and EMGT2, obtained in both tests. Furthermore, no significant differences (p>0.05) existed between mean values of EMGT1 in the vastus lateralis and rectus femoris, and VT, and LT (62.8 (14.5) and 69.0 (6.2) and 64.6 (6.4) and 68.7 (8.2)% of \( V_{\text{O}2\text{MAX}} \) respectively), or between mean values of EMGT2 in the vastus lateralis and rectus femoris, and VT, and OBLA (86.9 (9.0) and 88.0 (6.2) and 84.6 (6.5) and 87.7 (6.4)% of \( V_{\text{O}2\text{MAX}} \) respectively).

Conclusion—rms-EMG may be a useful complementary non-invasive method for analysing the aerobic-anaerobic transition (ventilatory and lactate thresholds) in elite cyclists.

Methods

SUBJECTS

Twenty eight elite male cyclists (age 24 (4) years; height 177.1 (5.2) cm; body mass 67.2 (6.0) kg; all values mean (SD)) participated in
this study. Sixteen subjects were professional road cyclists with a minimum competition experience of three years, and some of them had won several professional races. The other 12 were elite road cyclists (competition experience at least two years in the amateur category). Written informed consent was given before participation in the experiments, in accordance with the institutional human subjects guidelines (Complutense University of Madrid).

STUDY PROTOCOL
Before each exercise testing session, subjects were familiarised with the equipment and procedures used in this investigation. In addition, they were previously instructed to refrain from intense training during the day before testing.

Fifteen subjects were randomly selected for the reliability study. Each of them performed two exercise tests on a bicycle ergometer (Ergometrica 900; Ergo-line, Barcelona, Spain) on different occasions and separated from each other by a period of no more than five days. Each of the two tests consisted of a ramp protocol until exhaustion, starting at 0 W. The power output was increased by 5 W every 12 seconds and pedalling cadence was kept constant at 70–80 rpm. The selection of a ramp-like protocol instead of a graded steady state test was chosen for two reasons: (a) in previous research conducted in our laboratory this type of protocol was used to analyse physiological responses—that is, ventilatory thresholds and lactate kinetics—in elite/ professional cyclists during incremental exercise; (b) previous research showing a non-linear increase in the EMG response to incremental exercise used exercise protocols comparable with that used in our investigation. For the validity study, each of 28 subjects performed a single bicycle ergometer test following the above protocol.

Exercise tests were terminated (a) voluntarily by the subjects, (b) when pedalling cadence could not be kept at least at 70 rpm, or (c) when established criteria of test termination were met. Each test was performed under similar conditions (21–24°C and 45–55% relative humidity).

ANALYSIS OF EXPIRED GAS AND DETERMINATION OF VENTILATORY THRESHOLD
During the tests, gas exchange data were collected continuously using an automated breath by breath system (CPX; Medical Graphics, St Paul, Minnesota, USA). The measuring instruments were calibrated before each test and the necessary environmental adjustments made. VT1 was determined using the criteria of an increase in the ventilatory equivalent for oxygen (VE/VO2) with no increase in the ventilatory equivalent for carbon dioxide (VE/VECO2) and the departure from linearity of VE, whereas VT2 was determined by using the criteria of an increase in both VE/VO2 and VE/VECO2. Two independent observers detected VT1 and VT2, following the criteria previously described. If they did not agree, the opinion of a third investigator was included.

ANALYSIS OF BLOOD LACTATE
Blood samples (25 μl) for the measurement of blood lactate (YSI 23L; Yellow Springs Instruments, Yellow Springs, Ohio, USA) were taken from fingertips at rest, every two minutes during the test, and immediately after termination of exercise.

LT was determined by examining the “lactate concentration-power output (W)” relationship during the tests, using the methodology previously described by Weltman and co-workers. Thus the greatest power output that was not associated with a rise in lactate concentration above baseline was designated as the power output corresponding to LT. This always occurred just before the curvilinear increase in blood lactate observed with subsequent exercise intensities. A lactate increase of at least 0.2 mM (the error associated with the lactate analyser) was required for LT determination. OBLA, on the other hand, was defined as the power output corresponding to a blood lactate concentration of 4.0 mmol/L.

EMG
Electrode placement
Surface EMG recordings were taken from the vastus lateralis and rectus femoris (at sites respectively approximately one third and one half of the perpendicular distance from the superior border of the patella to the greater trochanter). Pairs of surface electrodes (Blue-sensor Medicotest Ag/AgCl electrodes; Rugmarken, Denmark) were attached to the skin with a 4 cm interelectrode distance. The electrodes were placed longitudinally with respect to the underlying muscle fibre arrangement. For those subjects who performed the test twice (reliability study), the skin was tattooed using ink in order to place the electrodes on the same site on the two tests. Two reference electrodes were placed over the anterior superior spine of the iliac crest. Before electrode application, the skin was shaved and abraded using sandpaper and cleaned with alcohol to minimise the source impedance. A saline EMG electrode gel was placed between the electrode and the underlying skin to enhance signal conductivity. The cables connected to the electrodes to measure myoelectrical activity were firmly attached with tape to minimise artefacts from leg movements.

EMG instrumentation and procedures
Myoelectrical activity was recorded using a ME3000P analyser (ME3000P; Mega Electronics Ltd, Kuopio, Finland). The measurement sensitivity of the instrument is ±1 μV and its range for bipolar EMG signals is ±5000 μV. The raw EMG signals were band-pass filtered between 20 and 480 Hz, amplified, and converted from analogue to digital at a sampling rate of 1 kHz. An EMG power spectral density was then computed for two second sampling periods, at fixed intervals throughout the tests, and the root mean square voltage (rms-EMG) contained in each two second
Determination of EMG breakpoint

Previous pilot studies conducted in our laboratory suggested the existence of a two threshold response in elite cyclists, with a second EMG breakpoint occurring at near to maximum intensities, which is easy to detect visually. To establish objective criteria for the determination of one or two breakpoints in the EMG power output response, we used a computer algorithm (Centro de Proceso de Datos, Complutense University of Madrid) that models rms-EMG response to exercise using multisegment linear regression.

With this method, a single linear regression is initially fitted to all data points and is used for later statistical comparisons. A brute force method is then used to fit two lines to the data points. The program calculates regression lines for all possible divisions of the data into two contiguous groups, and the pair of lines yielding the least pooled residual sum of squares is chosen as representing the best fit. The intersection point between these two lines occurred near the end of the test in all the subjects.

Thereafter the program attempts to fit a third line to the data in order to detect another breakpoint in the EMG data. The third middle segment is obtained by methodically adding points on the left side of the two line regression intersection point. The new regression line is then calculated and extended in the direction that yielded the lower sum of squares.

Finally, an analysis of variance determines whether a significant (p<0.05) reduction in the total sum of squares is achieved by the addition of a third line segment. The first (EMG$_{T1}$) and second (EMG$_{T2}$) EMG thresholds are then reported as the first and second intersection points respectively of the computerised model.

Results

MAXIMAL VALUES

Maximal values of $\dot{V}O_2$ and power output averaged 69.9 (6.4) ml/kg/min (range 60.0–82.3) and 432.8 (49.1) W (range 364–518) at the end of exercise.

PATTERN OF EMG RESPONSE

A two threshold response was detected in 90% of subjects in both vastus lateralis and rectus femoris, and the two breakpoints EMG$_{T1}$ and EMG$_{T2}$ occurred around 60–70% and 80–90% of $\dot{V}O_2$MAX. In 10% of the cases, EMG$_{T2}$ could not be detected in either of the two muscles studied, whereas EMG$_{T1}$ was found in all 28 subjects.

Figure 1 shows an example of an EMG response in one subject.

RELIABILITY

No significant differences existed between mean values of either EMG$_{T1}$ or EMG$_{T2}$ obtained in both tests (p<0.05) (table 1).

Intraclass correlation coefficients (r) between repeated measurements were significant (p<0.05) and high (table 1).

VALIDITY

Comparison between VT$_1$, LT, and EMG$_{T1}$

Using the methodologies described above, VT$_1$ and LT could be detected in 100% of subjects. Average values of LT occurred at a blood lactate concentration of 1.9 (0.5) mM. Table 2 presents mean values of VT$_1$, LT, and EMG$_{T1}$, expressed in $\dot{V}O_2$ (ml/kg/min), % of $\dot{V}O_2$MAX, and W. No significant differences (p>0.05) were found between means.
Electromyographic analysis in elite cyclists

Table 1 Reliability of measurements of the electromyographic thresholds EMGT1 and EMGT2.

<table>
<thead>
<tr>
<th></th>
<th>First test</th>
<th>Second test</th>
<th>Correlation (r)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vastus lateralis</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>EMGT1 V0 (ml/kg/min)</td>
<td>45.5 (2.5)</td>
<td>43.9 (3.0)</td>
<td>0.76</td>
</tr>
<tr>
<td>% V0_MAX</td>
<td>68.4 (4.0)</td>
<td>63.3 (4.2)</td>
<td>0.82</td>
</tr>
<tr>
<td>W</td>
<td>252.7 (17.6)</td>
<td>238.2 (17.9)</td>
<td>0.83</td>
</tr>
<tr>
<td>EMGT2 V0 (ml/kg/min)</td>
<td>59.1 (1.4)</td>
<td>62.6 (1.4)</td>
<td>0.96</td>
</tr>
<tr>
<td>% V0_MAX</td>
<td>90.2 (1.3)</td>
<td>90.2 (2.3)</td>
<td>0.87</td>
</tr>
<tr>
<td>W</td>
<td>367.6 (15.4)</td>
<td>372.6 (14.7)</td>
<td>0.86</td>
</tr>
<tr>
<td>Rectus femoris</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>EMGT1 V0 (ml/kg/min)</td>
<td>48.7 (2.5)</td>
<td>47.3 (1.7)</td>
<td>0.89</td>
</tr>
<tr>
<td>% V0_MAX</td>
<td>70.4 (3.0)</td>
<td>68.4 (2.9)</td>
<td>0.85</td>
</tr>
<tr>
<td>W</td>
<td>280.2 (22.4)</td>
<td>268.2 (20.8)</td>
<td>0.96</td>
</tr>
<tr>
<td>EMGT2 V0 (ml/kg/min)</td>
<td>59.0 (2.1)</td>
<td>59.8 (1.5)</td>
<td>0.73</td>
</tr>
<tr>
<td>% V0_MAX</td>
<td>89.0 (2.0)</td>
<td>86.2 (1.8)</td>
<td>0.75</td>
</tr>
<tr>
<td>W</td>
<td>374.2 (20.7)</td>
<td>360.8 (20.0)</td>
<td>0.87</td>
</tr>
</tbody>
</table>

All values are expressed as mean (SEM). No significant differences existed between means (p>0.05). All correlation coefficients were significant (p<0.05).

Table 2 Comparison between mean values of the first electromyographic threshold (EMGT1), the first ventilatory threshold (VT1), and the lactate threshold (LT).

<table>
<thead>
<tr>
<th></th>
<th>Vastus lateralis</th>
<th>Rectus femoris</th>
<th>VT1</th>
<th>LT</th>
</tr>
</thead>
<tbody>
<tr>
<td>EMGT1 V0 (ml/kg/min)</td>
<td>43.9 (1.7)</td>
<td>47.0 (1.5)</td>
<td>45.1 (1.2)</td>
<td>46.1 (1.6)</td>
</tr>
<tr>
<td>% V0_MAX</td>
<td>62.8 (2.8)</td>
<td>69.0 (2.1)</td>
<td>64.6 (1.1)</td>
<td>68.7 (1.7)</td>
</tr>
<tr>
<td>W</td>
<td>240.3 (9.8)</td>
<td>270.8 (13.8)</td>
<td>257.8 (10.0)</td>
<td>269.8 (12.9)</td>
</tr>
</tbody>
</table>

All values are expressed as mean (SEM). No significant differences existed between means (p>0.05).

Table 3 Comparison between mean values of the second electromyographic threshold (EMGT2), the second ventilatory threshold (VT2), and the onset of blood lactate accumulation (OBLA).

<table>
<thead>
<tr>
<th></th>
<th>Vastus lateralis</th>
<th>Rectus femoris</th>
<th>VT2</th>
<th>OBLA</th>
</tr>
</thead>
<tbody>
<tr>
<td>EMGT2 V0 (ml/kg/min)</td>
<td>61.5 (1.1)</td>
<td>59.7 (1.3)</td>
<td>59.0 (1.1)</td>
<td>60.7 (1.5)</td>
</tr>
<tr>
<td>% V0_MAX</td>
<td>86.9 (1.5)</td>
<td>88.0 (1.4)</td>
<td>84.6 (6.5)</td>
<td>87.7 (1.3)</td>
</tr>
<tr>
<td>W</td>
<td>371.1 (9.2)</td>
<td>367.5 (14.1)</td>
<td>352.8 (11.4)</td>
<td>377.6 (13.0)</td>
</tr>
</tbody>
</table>

All values are expressed as mean (SEM). No significant differences existed between means (p>0.05).

Comparison between VT2, OBLA, and EMGT2. Using the methodologies described above, VT2, OBLA, and EMGT2 could be detected in 100% of subjects. Table 3 presents mean values of VT2 and OBLA.

Discussion

To our knowledge, this is the first report to determine the validity of the EMG method for analysing the aerobic-anaerobic transition phase in top level athletes (professional cyclists) during cycle ergometry using a ramp test. Previous studies on EMG response during such protocols have been conducted with subjects of considerably lower fitness levels (in such protocols have been conducted with subjects of considerably lower fitness levels (in most studies, mean V0_MAX <65 ml/kg/min, and in all of them mean peak power output <360 W) than those selected for our investigation (V0_MAX 69.9 (6.4) ml/kg/min; peak power output 432.8 (49.1) W). To date, no conclusion could be inferred from previous research.
on the EMG response of elite cyclists reaching high power outputs during exercise testing. In addition, no previous study has assessed the reliability of the EMG method during a ramp protocol. As reliability is an integral part of validity, no previous report has assessed the validity of such a method for analysing the aerobic-anaerobic transition.

A novel finding of our study was that in elite cyclists the EMG activity of the muscles primarily involved in pedalling (vastus lateralis, rectus femoris) show a two threshold response with two distinct breakpoints, EMG\textsubscript{1} and EMG\textsubscript{2}, occurring at an exercise intensity of 60–70\% and 80–90\% of VO\textsubscript{2MAX} respectively. A second finding of our investigation was that the EMG method is both valid and reliable for analysing the aerobic-anaerobic transition during exercise. Finally, in elite cyclists it appears that muscle fatigability is not significantly affected by muscle fibre type distribution, as EMG\textsubscript{1} and EMG\textsubscript{2} occurred at similar intensities in the vastus lateralis and rectus femoris.

**METHODOLOGICAL LIMITATIONS**

Our study is not without potential limitations. Firstly, the literature on the usefulness of EMG in the determination of ventilatory or lactate thresholds is somewhat controversial. The differences in results could be attributed to difficulties in reliably following the myoelectrical activities in working muscles. In addition, during exertion the skin blood flow and temperature increase. This phenomenon could induce alterations in the electrical properties of the skin and affect the records of surface EMG. In this regard, previous research has shown no correlation between skin temperature and EMG measurements, excluding the possibility that EMG breakpoints are related to changes in electrical properties of the skin.

On the other hand, although we measured the myoelectrical activity of the muscles (vastus lateralis and rectus femoris) involved during the work (descending) period of pedalling only, no records were obtained for other muscles—for example, gastrocnemius, tibialis anterior—that are also involved during the rest (ascending) period of pedalling exercise. However, previous research has shown that a non-linear increase in EMG also occurs in the gastrocnemius muscle. In addition, previous EMG\textsuperscript{22} and biopsy studies\textsuperscript{23} suggest that the muscle most heavily involved during cycling is the vastus lateralis.

Furthermore, a single ramp increment (5 W every 12 seconds) was chosen in our protocol, whereas the results of a previous study using different ramp slopes (10, 20, 30 and 40 W/min) suggest that the metabolic state at which the EMG\textsubscript{2} occurs in untrained subjects may differ during the different ramp exercises. Further research should be conducted with elite cyclists to determine the influence of varying ramp protocols on the EMG response. Finally, the results obtained during a progressive test in a cycle ergometer using a fixed cadence (70–80 rpm), as was used in this investigation, cannot be easily extrapolated to

![Figure 2](image1.png) Average power output (W) by EMG\textsubscript{1} and VT\textsubscript{1} 

![Figure 3](image2.png) Average power output (W) by EMG\textsubscript{1} and LT 

![Figure 4](image3.png) Average power output (W) by EMG\textsubscript{2} and VT\textsubscript{2} 

![Figure 5](image4.png) Average power output (W) by EMG\textsubscript{2} and OBLA
Electromyographic analysis in elite cyclists

In these studies, the aerobic-anaerobic transition was expressed using ventilatory or lactate thresholds. Therefore, with increasing work intensity, the muscular power required in tests performed at a fixed cadence is considerably higher than that required by tests at increasing cadence—that is, field tests—and under actual cycling conditions.

The EMGT has therefore been suggested as an alternative non-invasive method for determination of ventilatory and/or lactate thresholds. Our results are in overall agreement with those of previous studies, as no significant differences were observed between the exercise intensity corresponding to the EMGT and that corresponding to VT, or LT. It follows that rms-EMG may represent a complementary non-invasive indicator of the aerobic-anaerobic transition during gradual exercise tests. A second breakpoint (EMGT2), however, also occurred in our subjects, which needs further discussion.

Coincidence of EMG and Blood Lactate Responses

The reason for an abrupt increase in EMG activity above a certain exercise intensity is not fully understood, and could be at both a local (muscle fatigue) and a generalised level. With respect to the first phenomenon, it can be assumed that after LT is reached, ATP supply from slow twitch oxidative (type I) fibres through oxidative phosphorylation becomes inadequate and therefore must be supplemented by using energy reserves available through anaerobic glycolysis, leading to metabolic acidosis. Accumulation of hydrogen ions, in turn, has been shown to impair excitation-contraction coupling through impairment of the function of (a) the sodium/potassium ATPase of the sarcolemma, (b) the calcium ATPase of the sarcoplasmic reticulum, or (c) the myosin ATPase involved in actin-myosin interaction. Moreover, high lactate levels per se may disrupt excitation-contraction coupling. Thus, under these physiological conditions, in order to compensate for the deficit in contractility resulting from impairment of fatigued motor units, muscle force output must be increased during an incremental exercise test through recruitment of additional motor units, particularly those made of fast twitch oxidative glycolytic (type IIa) and fast twitch glycolytic (type IIb) muscle fibres. The observed non-linear increases in rms-EMG (EMGT1 and EMGT2) could therefore be explained by a progressive recruitment of motor units with possible participation of type IIa and IIb fibres (at the EMGT1 and EMGT2, respectively) producing larger action potentials, followed by some degree of synchronisation of motor unit potentials as these fibres may undergo progressive fatigue. In fact, our findings are in agreement with previous research with muscle biopsy samples, which has shown a 1:1 relation between the fraction of active fibres in the vastus lateralis and the intensity (determined as a percentage of \( \text{VO}_{2\text{MAX}} \)) of cycle ergometer exercise following this pattern: at about 40% of \( \text{VO}_{2\text{MAX}} \), almost only type I fibres are recruited, whereas at about 60% of \( \text{VO}_{2\text{MAX}} \) (EMGT2) both type I and IIa are activated, and during severe exercise (about 90% of \( \text{VO}_{2\text{MAX}} \) or EMGT3), fibres of type I, IIa, and IIb are recruited.

However, the underlying mechanism may not be limited to a local level (muscle fatigue induced by metabolic acidosis). Such an hypothesis is supported by previous research showing that changes in EMG activity consistent with motor unit recruitment and muscle fatigue can also be recorded from patients with McArdle's syndrome or from normal subjects under conditions of changing muscle pH. Therefore, other ions—for example, potassium, ammonia, adenosine monophosphate, and magnesium—may be responsible for altering muscle function during exercise. In addition, Airaksinen and co-workers showed that both working (vastus lateralis and gastrocnemius) and non-working (frontalis) muscles showed a shift in EMG at the same load, suggesting that the breakpoint(s) observed in muscle electrical activity may not only be attributed to peripheral fatigue. Thus the explanation could also be a change in the basic activation of muscles in general. Such a generalised response could be due to a change in the membrane function initiated not only by an
increase in lactate levels but also by other factors such as an increase in neural or hormonal activity.\textsuperscript{1}  

**COINCIDENCE OF EMG AND VENTILATORY RESPONSES**  
The correlation encountered between VT and EMG\textsubscript{T1}, on the one hand, and VT\textsubscript{2} and EMG\textsubscript{T2}, on the other, could be due to a muscle derived signal to ventilation. Indeed, previous studies provide evidence for the existence of ergoreceptors or mechanoreceptors which respond to increases in the work performed per unit muscle.\textsuperscript{20} Moreover, Morikawa and co-workers\textsuperscript{31} found that a ventilatory response occurred in normal subjects with both active and passive leg exercise, but was absent with passive exercise in patients with thoracic spine transection. It could then be hypothesised that ventilation shows a first deflection point (VT\textsubscript{1}) when muscle work is increased (at an exercise intensity corresponding to EMG\textsubscript{T1}) and a further increase (VT\textsubscript{2}) when additional motor units are recruited to maintain power output—that is, recruitment of type IIb fibres at the EMG\textsubscript{T2}.

On the other hand, the ventilatory thresholds observed during an incremental test could be elicited by enhancement of the neural activity that originates directly from the subthalamic motor region or indirectly via α-γ coactivation of motoneurons innervating the muscle fibres of exercising limbs.\textsuperscript{32} This increase in neural activity may occur during incremental exercise in response to the need to progressively recruit additional motor units comprised of type IIA possibly at the EMG\textsubscript{T1} and IIb fibres possibly at the EMG\textsubscript{T2} respectively as the work rate is increased and individual fibres begin to fatigue.\textsuperscript{1} Indeed, numerous investigations have shown simultaneous increases in ventilation and EMG activity during incremental exercise.\textsuperscript{3 4 15 32}

The results of a recent study, however, do not support a link between motor unit recruitment and ventilation, as evidenced by the disassociation between the EMG, of the rectus femoris and the ventilatory threshold that existed in trained cyclists after glycogen depletion.\textsuperscript{5} A similar study should be conducted with professional cyclists in order to assess the effects of glycogen depletion on the rms-EMG response.

**ABSENCE OF EMG\textsubscript{T2} IN PREVIOUS RESEARCH**  
The question remains to be answered of why a second EMG threshold (EMG\textsubscript{T2}) was found only in this study, in contrast with previous research. In this regard, a possible explanation could be that the significant increases in neural activity to the exercising muscles that occur at EMG\textsubscript{T2} were not detected in previous studies because EMG activity was mainly recorded from a single muscle—that is, the vastus lateralis. This hypothesis is supported by the findings of Green and Patla,\textsuperscript{19} which showed that many different muscle groups contribute to the completion of an incremental exercise task and that the average level of activation increases differentially among the muscle groups. It may be possible that only highly trained cyclists (such as those selected for our study) are able to effectively recruit a sufficient number of motor units (especially those with fast fibres) within individual muscles (vastus lateralis or rectus femoris) at near to maximum intensities during an incremental test such as to induce a second breakpoint in the EMG response to exercise. Indeed, EMG\textsubscript{T2} occurred at a power output as high as 370 W, which in turn elicited a VO\textsubscript{2} of about 60 ml/kg/min. Both values are well above the maximal levels of power output and VO\textsubscript{2} reported in most previous EMG studies (<350 W and <55 ml/kg/min respectively). Although two studies\textsuperscript{33} have been conducted in this area using highly trained cyclists with similar values of VO\textsubscript{MAX} to those of our subjects (about 70 ml/kg/min), they attained considerably lower peak power outputs (<360 W) during incremental tests.

**SIMILAR RESPONSES IN VASTUS LATERALIS AND RECTUS FEMORIS**  
Based on the results of previous research with non-elite athletes,\textsuperscript{33–35} a disparity in the EMG/exercise intensity relation would be expected to exist between the vastus lateralis and the rectus femoris. In effect, previous research has shown distinct responses in the two muscles, such as the occurrence of the EMG fatigue threshold at lower intensities in the rectus femoris.\textsuperscript{14} This disparity could be partly attributed to kinesiological differences, as the rectus femoris (unlike the vastii muscles) is a biarticular muscle involved in both leg extension and thigh flexion.\textsuperscript{34} On the other hand, possible differences in muscle fibre composition cannot be excluded, the rectus femoris being comprised of a greater percentage of fatigable fast twitch fibres.\textsuperscript{35} In this regard, few data are available on the muscle fibre composition of leg muscles in professional cyclists. In our investigation, however, EMG\textsubscript{T1} and EMG\textsubscript{T2} occurred at similar intensities in both muscles, suggesting a similar pattern of muscle fibre recruitment in the different leg muscles of top level cyclists as an adaptation to training and competition.

**CONCLUSION**  
In elite cyclists the EMG activities of two leg muscles (vastus lateralis and rectus femoris) show similar patterns, with two distinct breakpoints, EMG\textsubscript{T1} and EMG\textsubscript{T2}, occurring at an exercise intensity of 60–70% and 80–90% of VO\textsubscript{MAX} respectively. The rms-EMG method seems to be both valid and reliable for analysing the aerobic-anerobic transition during cycle ergometer exercise.
Electromyographic analysis in elite cyclists


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**Take home message**

During incremental cycle ergometer tests performed by top level cyclists, the relation between EMG activity (rms-EMG) of exercising muscles (vastus lateralis and rectus femoris) and exercise intensity exhibits two breakpoints, at around 60–70% and 80–90% of VO2 max respectively. rms-EMG may be used as a complementary indicator of the aerobic-anaerobic transition in physiological evaluations of cyclists.