Muscle Mitochondrial Dysfunction in Horses Affected by Acute Laminitis

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Abstract

Laminitis is a common and debilitating disease affecting horses and ponies. It often leads to the demise of the animal. Energy deficiency is suspected to entail the disruption of the hemidesmosomes leading to the failure of the dermal-epidermal interface. The aim of this study was to measure the muscle mitochondrial function by high resolution respirometry. Muscle micro-biopsies were obtained from 11 horses affected by acute metabolic laminitis, 6 horses affected by acute laminitis resulting from a systemic inflammation response syndrome and 28 healthy horses distributed in 2 control groups: 17 horses with a body condition score (BSC) ranging from 0 (emaciated) to 5 (obese)) of 2 to 3 and 11 horses with a BSC of 4 to 5. During the acute phase of laminitis, a significant reduction of the muscle mitochondrial respiration was observed. The muscle mitochondrial dysfunction occurred independently of the etiology (metabolic disorder or systemic inflammation) leading to laminitis. The reduction of the oxidative phosphorylation and of the maximal respiratory capacity (after uncoupling) may induce depletion of the cell’s ATP content. If the same mitochondrial alteration occurs in the foot lamina, mitochondria targeting should be considered for the future, not only to better understand the physiopathology of the disease but also to maintain and to support the mitochondrial function before reaching the « mitochondrial dysfunction threshold » that may lead to the failure of the dermal-epidermal interface.

Keywords: Horse; Laminitis; Mitochondria; Electron transfer system; Energy metabolism dysfunction; Oxidative phosphorylation; Inflammation; Reactive oxygen species

Introduction

Laminitis is a common and debilitating disease affecting horses and ponies. In its severe form it may necessitate the demise of the animal [1]. Laminitis is characterized by the failure of the laminar dermal-epidermal interface resulting in severe lameness.

Despite ongoing research, laminitis remains a severely damaging condition of which the physiopathology is poorly understood. Currently, the primary hypotheses explaining the pathogenesis of acute laminitis include inflammation and extracellular matrix degradation, metabolic abnormalities and endothelial/vascular dysfunction [2].

Recent epidemiological studies have identified a variety of risk factors for the development of laminitis, such as weight gain, season of the year, hours of sunshine on the pasture, recent access to grass, box rest, owner-reported history of laminitis, lameness or foot sorenessness after shoeing/trimming, and existing endocrinopathic and metabolic disease [3,4]. Endotoxemia and sepsis have been recognized in a follow-up of hospitalized horses as the only significant risk factor for developing laminitis [5]. Inflammation seems to play a central role in the pathogenesis of laminitis, and this independently of the key factors involved [6]. In cases of sepsis, local infection is accompanied by systemic neutrophil activation. Systemic neutrophil activation is also encountered in equine laminitis, as demonstrated by the up-regulation of cytokine expression, the dynamic changes in blood neutrophil phenotype, the formation of neutrophil-platelet aggregates and the infiltration of inflammatory cells [7-10]. However, other factors than endotoxins are responsible for the changes in laminar tissue gene expression that occurs during the development of acute equine laminitis [11,12].

The systemic and the local inflammatory responses and the release of cytokines, enzymes such as myeloperoxidase or elastase, have largely been described in previously developed laminitis models such as Carbohydrate Overload (CHO), Oligofructose (OF) and Black Walnut Heartwood Extract (BWHE). This underlines the major role of neutrophil activation [13-16]. The hypothesis of an involvement of neutrophil-mediated mitochondrial dysfunction was previously observed in muscle micro-biopsies from horses after strenuous exercise [17]. Moreover, in cultured muscle cell lines, it was demonstrated that the potent neutrophil-derived oxidative enzyme myeloperoxidase could enter the cell and disturb the electron transfer system resulting in altered ATP production [18].

Laminitis has also been associated with horses diagnosed with metabolic disorders, such as obesity, pituitary pars intermedia dysfunction or equine metabolic syndrome. The primary characteristic of these pathologies is the development of insulin resistance, characterized by hyperinsulinemia with eu- or hyperglycemia and subsequent chronic pro-inflammatory state [19,20]. While several aspects of the relationship between insulin resistance and laminitis have been investigated, the underlying pathological pathway has not yet been identified. An experimental model of laminitis showed that healthy Standard bred horses subjected to prolonged...
hyperinsulinemia develops laminitis within 48 h [21]. Furthermore, natural cases of equine endocrinopathic laminitis are clearly associated with hyperinsulinemia [22].

Recently, Gauff et al. showed that short-term hyperinsulinemia in the isolated, perfused extracorporeal equine digit leads to a marked increase in vascular resistance and an increase in endothelin-1 expression [23].

Several studies confirmed a mitochondrial dysfunction in human patients suffering from metabolic disorders related to insulin resistance [24,25]. Indeed, glucose and lipid metabolism are largely dependent on energy generated by mitochondria and several experimental protocols have demonstrated an effect of hyperinsulinemia on mitochondrial function [26,27].

In this work, we hypothesized that muscle mitochondrial function is altered in horses suffering from laminitis. As the mitochondrial function is frequently studied in muscles, we decided to investigate this using High-Resolution Respirometry (HRR) on muscular micro-biopsies [28]. High-resolution respirometry allows the determination of Oxidative Phosphorylation (OXPHOS) and Electron Transfer System (ETS) capacities in small samples (2 mg) of permeabilized tissues [28]. Multiple Substrate-Uncoupler-Inhibitor Titration (SUIT) protocols have been recently developed to screen the mitochondrial function of equine muscles [29]. In the present study, this technique was used to study mitochondrial function in laminitis affected horses.

Material and Method

Horses

All procedures were approved by the Animal Ethic Commission of the University of Liege (agreement n°07–629) and informed owner consent was obtained when appropriate. Muscle micro-biopsy is performed on healthy horses (n=28) and on clinical cases of acute laminitis observed in the Equine Clinic of the University of Liege between 2011 and 2013 (n=17). The Body Condition Score (BSC) was determined based on palpation of fat deposits in five specific body areas and by visual assessment of seven anatomic sites [30]. Each evaluation receives a notation and the average defines the BSC ranging from 0 (emaciated) to 5 (obese). All the horses were categorized in two groups: score 0 for a BSC between 2 to 3 (n=23) and score 1 for a BSC between 4 to 5 (n=22).

The healthy horses were divided in two control groups: control obese (contob) and control non obese (cont fit). Muscle micro-biopsies were obtained from 2 distinct groups of acute laminitic horses. The first group was assigned as “metabolic laminitis” (Met Lam). These horses were showing symptoms of acute laminitis and were diagnosed as suffering from metabolic syndrome [31]. The second group was assigned as “inflammatory laminitis” (Infl Lam) and was composed of horses affected by acute laminitis resulting from a systemic inflammatory response syndrome.

So, 4 groups of horses were established: healthy non obese (Cont fit: n=17), healthy obese (Contob: n=11), laminitic obese (Met Lam, n=11) and laminitic non obese (Infl Lam: n=6). All data obtained from the horses are reported in the Table 1.

Muscle micro-biopsy

Approximately 20 mg of muscle have been collected from the triceps brachii muscle using a 14 G biopsy needle mounted on an automatic instrument as earlier described (Pro-Mag™ Ultra Biopsy Instrument, Angiotech, Gainesville, FL, USA) [29]. Briefly, the sampling site was shaved and desensitized by subcutaneous injection of 0.5 ml of mepivacain (Scandicaine 2%, AstraZeneca, Brussels, Belgium), and aseptically prepared.

Muscle micro-biopsy specimens were taken at 40 mm depth in the long head of the triceps brachii through a skin incision. Two to three muscle samples were obtained via the same skin opening and transferred immediately into ice-cold relaxing solution BIOPS containing 10 mM CaK2-EGTA, 7.23 mM K2-EGTA, 20 mM imidazole, 20 mM taurine, 50 mM K-MES, 0.5 mM dithiothreitol, 6.56 mM MgCl2, 5.77 mM ATP and 15 mM phosphocreatine adjusted to pH 7.1. Fibers were kept at 4°C until further preparation.
10 mM) for convergent electron flow through complex I and II (CI+II) electron flow through complex I (CI) was supported by the NADH-respiration medium (MiR0 limitation of respiration. As recommended, respiratory flux was High resolution respirometry

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Table 1: Characteristics of horses used in the study and their respiratory parameters from substrate-uncoupler-inhibitor titration protocol I (SUIT1) applied on equine permeabilized muscle fibers of triceps brachii: Complex I (CI), Complex I + II (CI+II), maximal respiratory (ETSmax) and Complex II (CII) capacities.

| Group                     | Mean (± SD) | Mean (± SD) | Mean (± SD) | Mean (± SD) |
|---------------------------|-------------|-------------|-------------|-------------|
| Control obese (n=11)      | 13 ± 4      | 522 ± 122   | 4.5 ± 17.11 | 82.38 ± 33.23 | 108.65 ± 44.58 | 68.31 ± 31.17 |
| Inflammatory laminitis    | 12 ± 5      | nd          | 4.5 ± 17.11 | 102.76 ± 23.45 | 119.79 ± 24.27 | 81.85 ± 16.79 |
| Hol1                      | 10          | 625         | 3           | 27.00        | 65.50          | 70.40          | 54.60          |
| Hol2                      | 2           | 470         | 2           | 52.37        | 77.78          | 103.16         | 56.21          |
| Hol3                      | 3           | 700         | 3           | 27.02        | 48.17          | 73.15          | 42.67          |
| Hol4                      | 21          | 650         | 3           | 48.64        | 71.50          | 97.82          | 56.27          |
| Hol5                      | 11          | 580         | 2           | 78.28        | 118.38         | 151.27         | 78.80          |
| Hol6                      | 4           | 700         | 3           | 35.36        | 67.31          | 67.07          | 57.05          |
| Control non obese (n=17)  | 9 ± 7       | 604 ± 89    | 2-3         | 44.78 ± 19.57 | 74.77 ± 23.54 | 97.15 ± 29.52 | 57.60 ± 11.70 |
| mean (±SD)                | 8±3         | nd          | 2-3         | 58.79 ± 12.6 | 116.37 ± 18.9 | 139.66 ± 19.38 | 96.13 ± 12.82 |

High resolution respirometry

Connective tissue and other visible tissues than muscle were removed and muscle fibers were genetically separated using two pairs of forceps with sharp tips. Complete permeabilization of the plasma membrane was obtained by gentle agitation for 30 min at 4°C in 2 ml of BIOPS solution containing 50 µg/ml saponin. The fiber bundles were rinsed by agitation for 10 min in ice-cold mitochondrial respiration medium (MiR0: 0.5 mM EGTA, 3 mM MgCl2, 60 mM K-lactobionate, 20 mM taurine, 10 mM KH2PO4, 20 mM Heps, 110 mM sucrose and 1 g/l BSA essentially fatty acid free) adjusted to pH 7.1. The permeabilized muscle fibers (Pfi) were immediately used for HRR.

One to 2.5 mg (Microbalance, Mettler Toledo, Zaventem, Belgium) of Pfi were added to each Oxgraph-2k chamber (Oroboros Instruments, Innsbruck, Austria) containing 2 ml of MiR0 at 37°C. Oxygen concentration (µM), and oxygen flux per muscle mass (pmol O2/sec*mg) were recorded online using DatLab software (Oroboros Instruments, Innsbruck, Austria). After calibration of the oxygen sensors at air saturation, a few ml of oxygen were introduced into the gas phase above the stirred aqueous phase in the partially closed system (ETS) over the phosphorylation system [28]. This was tested by stepwise addition of the uncoupler FCCP (Carbonyl cyanide 4-(trifluoromethoxy)phenylhydrazone; 0.05 µM, followed by several 0.025 µM additions (steps) until maximal oxygen flux was reached), obtaining the ETS capacity, ETSmax with convergent electron flow through CI+II. Electron input into the Q-junction through complex II (CII) alone was subsequently induced by inhibition of CI by rotenone (Rot; 0.5 µM). Finally, residual oxygen consumption (ROX) was obtained by addition of antimycinA (2.5 µM) to block electron transfer through complex III (CIII). Oxygen fluxes were corrected by subtracting ROX from each measured mitochondrial steady-state.

Integrity of the outer mitochondrial membrane was tested by adding 10 µM cytochrome c after ADP in the presence of substrates feeding electrons into CI. Injury of the outer mitochondrial membrane leads to loss of cytochrome c from the mitochondria, and to significant stimulation of respiration following addition of exogenous cytochrome c to the respiration medium.

Oxygen flux was expressed as tissue mass-specific respiration (per mg), or as Flux Control Ratios (FCR), with internal normalization for maximum ETS capacity.

To obtain extended information on coupling and substrate control parameters, the SUIT protocol was applied on Pfi. In this protocol, electron flow through complex I (CI) was supported by the NADH-linked substrates glutamate + malate (GM; 10 and 2 mM) with subsequent addition of ADP (2.5 mM). ADP-stimulated respiration represents OXPHOS capacity (State P). Then, we added succinate (S; 10 mM) for convergent electron flow through complex I and II (CI+II) into the Q-junction (supported by GMS). The capacity of the phosphorylation system (adenine nucleotide translocase, inorganic phosphate transporter, and ATP synthase) may limit OXPHOS capacity with an apparent excess capacity of the electron transport system (ETS) over the phosphorylation system [28].

Statistical analysis

Respirometry measurements were performed at least in duplicate for each protocol. The oxygen flux and FCR are presented as mean ± SD for the 4 groups, i.e. Cont fit, Contob, Met Lam and Infl Lam. A levene’s test was used for all the dependent variables to test the equality of variance then a two-way analysis of variance was used for each dependent variable taking into account two qualitative factors:
score and laminitis. Furthermore, a Student T test were used when the variances are equal to compare the mean oxygen flux values of CI, CI +II, ETS\textsubscript{max} and CII alone between the Cont fit group and the Infl Lam group. The SUIT protocol allowed us to also establish the FCR for each group of horses (Figure 3). Cont fit horses used 68.99 (± 7.21) % of their maximal ETS capacity via respectively CI and CII alone. When substrates for CI and CII are available (CI+II), they used 83.45 (± 9.46) % of their maximal ETS capacity. Similar values are observed in the control obese group (CI: 43.11 ± 6.9; CII: 68.87 ± 9.13; CI+II: 86.16 ± 12.32). The C1+CII values represented the maximal proportion of their respiratory capacity that mitochondria are effectively able to use for producing ATP, and thus, energy. Notwithstanding, via CI, Met Lam or Infl Lam horses did not use a significant greater proportion of their respective maximal ETS capacity than Control horses. On the contrary, the 2 pathological groups used a significant lower part of their maximal ETS capacity via CII: 61.72 (± 7.02) % of their maximal ETS capacity via CII alone. When FCR was applied only the CII was significantly different between the horses affected or not by the disease. The T-test analysis comparing the ContOb group to the Met Lam group showed a significant decrease of the oxygen flux for CI +CII and CII alone. Compared to Cont Fit horses, the Infl Lam group showed a significant decrease of the oxygen flux for CI+CII, ETS capacities and CII alone.

The two-way analysis of variance to test the effect of the two qualitative factors (Score and laminitis) revealed that any dependent variable was significantly affected by the body score index. However, a significant effect of the factor “laminitis” was observed for CI, CI+II, ETS capacities, CII alone. Otherwise, when FCR was applied only the CII was significantly different between the horses affected or not by the disease. The T-test analysis comparing the ContOb group to the Met Lam group showed a significant decrease of the oxygen flux for CI +CII and CII alone. Compared to Cont Fit horses, the Infl Lam group showed a significant decrease of the oxygen flux for CI+CII, ETS capacities and CII alone.

81.85 (± 16.79) pmol O\textsubscript{2}/sec. Compared to the cont fit horses, the contob horses showed a significant decrease of ETS capacities and CII alone.

The comparison of the BSC measured in each group showed significant differences between the Cont fit and the Contob groups and between the Infl Lam and the Met Lam groups.

**Results**

The comparison of the BSC measured in each group showed significant differences between the Cont fit and the Contob groups and between the Infl Lam and the Met Lam groups.

The figure 1 illustrates oxygraphic records obtained on one healthy horse and two laminitic horses (Metabolic Laminitis, Inflammatory Laminitis).

**Figure 1:** Example of HRR curves. Illustration of the curves obtained via the Oroboros oxygraph software [DatLab 4.0, OROROBOS DatLab software, Innsbruck, Austria] for one horse of each group [Control, Metabolic laminitis and Inflammatory laminitis]. The blue line represents the oxygen concentration in the medium [pmol/ml] and the red line represents the slope of the blue line, thus the oxygen flux expressed as tissue mass-specific respiration [pmol oxygen/sec*mg of permeabilized fibers]. The vertical lines represent the different substrates/inhibitors/uncouplers added to the oxygraph chambers. The black rectangles represent the selected time laps during which CI [Complex I], CI + II [Complex 1 and 2], maximal respiratory [ETSmax] and CII [Complex II] capacities were calculated by DatLab™.

The Cont fit horses showed a mean tissue mass-specific respiration (per mg of muscle) of 58.79 (± 12.56) pmol O\textsubscript{2}/sec with CI substrates (Table 1). Their maximalOXPHOS and ETS capacities with physiological CI+II substrates were 116.37 (± 18.89) and 139.65 (±19.38) pmol O\textsubscript{2}/sec respectively. For the CII alone, the tissue mass-specific respiration was 96.13 (± 12.82) pmol O\textsubscript{2}/sec. The ContOb horses showed a mean tissue mass-specific respiration (per mg of muscle) of 51.60 (±13.98) pmol O\textsubscript{2}/sec with CI substrates (Table 1). Their maximalOXPHOS and ETS capacities with physiological CI+II substrates were 102.76 (± 23.46) and 119.76 (± 24.27) pmol O\textsubscript{2}/sec respectively. For the CII alone, the tissue mass-specific respiration was 68.99 (± 7.21) % of their maximal ETS capacity via respectively CI and CII alone. When substrates for CI and CII are available (CI+II), they used 83.45 (± 9.46) % of their maximal ETS capacity. Similar values are observed in the control obese group (CI: 43.11 ± 6.9; CII: 68.87 ± 9.13; CI+II: 86.16 ± 12.32). The CI+CII values represented the maximal proportion of their respiratory capacity that mitochondria are effectively able to use for producing ATP, and thus, energy. Notwithstanding, via CI, Met Lam or Infl Lam horses did not use a significant greater proportion of their respective maximal ETS capacity than Control horses. On the contrary, the 2 pathological groups used a significant lower part of their maximal ETS capacity via CII: 61.72 (± 4.85) % for Met Lam and only 59.29 (± 9.33) % for Infl Lam. However, when regarding the proportion of ETS\textsubscript{max} used via CI+II, a significant reduction was only observed for Met Lam horses as compared to the Contob group. Thus, they were able to use only 75.94 (± 8.1) % of their maximal respiratory capacity. A similar non-significant trend was also observed in the Infl Lam group.
Discussion

The results of the present study confirm the hypothesis that significant reduction of muscle mitochondrial respiration is observed during the acute phase of laminitis. The reduction of OXPHOS may result in a depletion of the cells’ ATP content. This is especially true if the glycolytic pathway cannot compensate for the deficient ATP. The mitochondrial function may be measured directly on lamina biopsies obtained immediately following euthanasia when the horses were trained was realized for the normal BSC groups, in order to obtain an adequate control group for the inflammatory laminitis group.

Nevertheless, the oxygen flux for CI+CII, CII alone were significantly reduced in both acute laminitis groups compared to healthy horses and maximal ETS capacity especially in the Infl Lam group.

The differences between the laminitis horses and the control horses having a high BSC could be related to a certain level of metabolic syndrome. Indeed, equine metabolic syndrome is characterized by obesity and local adiposity coupled with evidence of recurrent laminitis [35]. These authors observed that a marked increase in neutrophil reactive oxygen species production upon phagocytosis was observed in horses with metabolic syndrome that was closely correlated to the blood insulin concentration. Insulin resistance is currently observed in horses suffering from metabolic syndrome. Based on the results of this study, it would have been interesting to dose insulin in our patients. However, to confirm insulin resistance, it is also necessary to show abnormal glycemic and insulinemic responses to oral or IV glucose and/or insulin challenges. These tests would have been ethically unacceptable on clinical cases, taking into account the risks of aggravating the disease.

When internal normalization for maximum ETS capacity (FCR) was applied, we observed clearly that the complex II was principally affected. Mitochondrial complex II oxidizes succinate to fumarate as part of the Krebs cycle and reduces ubiquinone in the electron transport system. Succinate-driven oxidation via complex II may contribute significantly to high rates of production of Reactive Oxygen Species (ROS) by mitochondria [36,37].

A Complex II defect was accompanied by a biphasic increase of ROS and concurrent glutathione oxidation, both concomitant to a decrease of the ATP cellular level [38]. High ROS generation is considered as a key point in human patients affected by metabolic syndrome and insulin resistance [39]. The implication of ROS in the pathogenesis of laminitis has already been suggested [40,41]. Moreover, activated neutrophils generate ROS and release oxidant enzyme such as myeloperoxidase and proteases including elastase and matrix metalloproteinase 9. In a model of muscle cells in culture, it was demonstrated that myeloperoxidase increased significantly the ROS amount generated by anoxia and reoxgenation cycles [42]. Thus, the production of ROS in lamina could be explained by neutrophil activation with the presence of myeloperoxidase in the lamina and/or by the mitochondrial dysfunction.

In this study, the muscle mitochondrial dysfunction was observed independently of the key factors responsible of the laminitis [metabolic or inflammation]. If the same dysfunction occurs in the foot lamina, it might be possible that the ATP available to maintain the hemidesmosome activity becomes insufficient, once the alteration of the OXPHOS reaches a certain level, and that this would occur independently of the etiology. In other words, a «mitochondrial dysfunction threshold» would be reached to trigger the pathology.

To improve the comprehension of this pathophysiological mechanism, mitochondrial respiration and ATP levels should be studied on muscular biopsies from experimental laminitis induced by CHO, BWE, OF administration or hyperinsulinemic/euglycemic clamp model.

The mitochondrial function may be measured directly on lamina biopsies obtained immediately following euthanasia when the horses...
show an OBEL 1 grade. Mitochondrial respiration may also be measured on muscle micro-biopsies following the administration of the triggering factor.

Since samples were obtained from clinical cases, standardization was not possible and factors such as the time point of the micro-biopsy related to the onset of the laminitis, the gravity of the disease, and treatment before referral to the clinic may have influenced the results. These factors may partly explain the wide standard deviation observed.

Conclusion

This clinical study shows for the first time a significant decrease of muscle mitochondrial oxidative phosphorylation in horses affected by acute laminitis when compared to the values obtained from fit or obese healthy horses. Targeting the mitochondria in the foot should be considered as essential for a better understanding of the physiopathology and for new treatment strategies, with the aim to maintain and to support the mitochondrial function before the « mitochondrial dysfunction threshold » is reached and failure of the dermal-epidermal interface occurs.

Conflict of interest statement

The authors certify that there is no conflict of interest with any financial organization regarding the material discussed in the manuscript.

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