Hyperoxia affects the lung tissue: A porcine histopathological and metabolite study using five hours of apneic oxygenation

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Article history:
Received 3 July 2019
Received in revised form 12 September 2019
Accepted 12 September 2019
Available online 14 September 2019

Keywords:
Apneic oxygenation
Hyperoxia
Lung tissue
Histopathology
Porcine model
Metabolites

ABSTRACT

Background: Oxygen is a liberally dosed medicine; however, too much oxygen can be harmful. In certain situations, treatment with high oxygen concentration is necessary, e.g. after cardiopulmonary resuscitation. The amount of oxygen and duration of hyperoxia causing pulmonary damage is not fully elucidated. The aim of this study was to investigate pathophysiological and metabolite changes in lung tissue during hyperoxia while the lungs were kept open under constant low pressure.

Methods: Seven pigs were exposed to 100% oxygen for five hours, using an apneic oxygenation technique with one long uninterrupted inspiration, while carbon dioxide was removed with an interventional lung assist. Arterial blood samples were collected every 30 minutes. Lung biopsies were obtained before and after hyperoxia. Microscopy and high-resolution magic angle spinning nuclear magnetic resonance spectroscopy were used to detect possible pathological and metabolite changes, respectively. Unsupervised multivariate analysis of variance and paired sample tests were performed. A two-tailed p-value ≤ 0.05 was considered significant.

Results: No significant changes in arterial pH, and partial pressure of carbon dioxide, and no clear histopathological changes were observed after hyperoxia. While blood glucose and lactate levels changed to a minor degree, their levels dropped significantly in the lung after hyperoxia (p < 0.04). Reduced levels of antioxidants (p < 0.05), tricarboxylic acid cycle and energy (p < 0.04) metabolites and increased levels of several amino acids (p ≤ 0.05) were also detected.

Conclusion: Despite no histological changes, tissue metabolites were altered, indicating that exposure to hyperoxia affects lung tissue matrix on a molecular basis.

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1. Introduction

Oxygen is one of the most widely used therapeutic agents in the world and in conditions with life threatening hypoxia it can be lifesaving. It is often liberally dosed, particularly in critically ill patients requiring mechanical ventilation, leading to supraphysiological partial pressures of oxygen (PaO2) [1–3]. Growing evidence indicates that too much oxygen may be harmful [4–6]. The organs most affected by normobaric hyperoxia are the lungs and eyes [4,7–9]. Since the lungs are exposed directly to oxygen in

https://doi.org/10.1016/j.metop.2019.100018
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the alveoli and indirectly by the blood in the pulmonary vessels, hyperoxia may cause more damage to the lung tissue compared to other tissues. Exposure time, atmospheric pressure, and fraction of inspired oxygen (FiO2) determine the cumulative oxygen amount leading to toxicity [6]. The severity of the hyperoxic lung tissue injury is directly proportional to the partial pressure of oxygen (PaO2), particularly above 60 kPa or a FiO2 approaching 1.0, and exposure time [4,6]. The pathologic changes induced by hyperoxia are not specific and resemble pathologic changes seen in other acute pulmonary diseases, such as the acute respiratory distress syndrome [10,11]. The earliest changes seen microscopically are capillary congestion, intra-alveolar hemorrhages, and interstitial and alveolar edema [12]. Inflammation of inflammatory cells from the blood to the lung tissue, such as lymphocytes, macrophages and plasma cells, as well as mild endothelial injury have also been reported [11]. These changes eventually lead to impaired gas exchange and multiorgan dysfunction [13].

Currently, the amount of oxygen causing pulmonary damage is still not fully elucidated. Numerous human and animal model studies have been performed with varying exposure time, FiO2 and with or without mechanical ventilation [10,14–22]. Exposure of the lung tissue with a FiO2 of 0.75–1.0 for 90 minutes has been reported to cause interstitial edema with subsequent intrusion of inflammatory and red blood cells into the alveoli, along with thickening of alveolar septa in Wistar rats [23]. In healthy humans, exposure to a FiO2 of 0.95 elicited hyperoxia-induced alveolar capillary leak after 17 hours. These changes were reversible and disappeared within two weeks after the exposure [15]. Baboons exposed to 100% oxygen first showed signs of alveolar septal thickening after 80 hours of exposure [21], whereas in young pigs exposed to a FiO2 above 0.80, capillary congestion and interstitial infiltration could be detected with light microscopy after 48 hours of exposure [24]. In-vitro research has demonstrated that human lung endothelial cells exposed to hyperoxia for 3–12 hours had enhanced nicotinamide adenine dinucleotide phosphate (NADPH) and nicotinamide adenine dinucleotide (NADH)-dependent reactive oxygen species (ROS) production [25]. In ovine lung endothelial cells exposed to hyperoxia for 30 minutes, free radicals were produced via the mitochondrial electron transport chain without the induction of cell death [26].

Ventilator treatment itself can cause damage to the lungs. The mechanisms involved are barotrauma, volutrauma and atelectrauma [27]. Barotrauma is due to high inflation pressure-mediated lung injury [28]. Volutrauma occurs because of overdistension-mediated lung injury [27]. Using low volume ventilation can also cause injury through multiple factors, such as opening and closing of lung units, impact on surfactant and regional hypoxia. These injuries are called atelectrauma [27]. It is therefore recommended to keep tidal volume low, peak pressure low and applying positive end expiratory pressure (PEEP) to reduce the risk of ventilator induced lung injury [29–31].

We hypothesize that hyperoxic-induced lung tissue injury affects the molecular balance and that these changes can be detected at an earlier stage compared to the characteristic histopathological changes. The application of apneic oxygenation enables the assessment of the effect of hyperoxia on lung tissue, minimizing interference from possible ventilator-induced injuries. During apneic oxygenation, pure oxygen is administered with low pressure to the cuffed endotracheal tube as one long uninterrupted inspiration, while carbon dioxide (CO2) is removed with interventional lung assist (iLA). Hence, the aim of this study was to identify possible histopathological and molecular changes in lung tissue exposed to a very high oxygen tension. Due to the anatomical and immunological similarities between the porcine and human lung, pigs make an excellent model for human respiratory disease and pathologies [24,32]. Two lung biopsies were collected from seven pigs before induction of hyperoxia and after five hours of exposure to a FiO2 of 1.0 and histopathological and metabolite analyses were performed.

2. Materials and methods

2.1. Animal care and anesthesia protocol

Seven 45 kg female Danish Landrace pigs were enrolled in this study after ethical approval by The Danish National Animal Ethics Committee (No. 2016–15–0201–00930).

The animals were allowed to acclimatize for six days before the experiments, with free access to water and food. Prior to the experimental procedure, overnight fasting was required. Anesthesia was induced with intramuscular injection of Zoletil®Vet, a mixture of ketamine (8.33 mg/ml) and tiletamine (8.33 mg/ml), benzodiazepine zolazepam (8.33 mg/ml), synthetic opioid butorphanol (1.67 mg/ml), and xylazine (8.33 mg/ml). The trachea was intubated (Portex tube, ID 6.5 mm, Smiths Medical, UK) and the lungs were mechanically ventilated (Dameca DREAM, Roedovre, Denmark) with tidal volume 6–8 ml/kg, FiO2 0.23, inspiratory/expiratory ratio 1:2 and positive end-expiratory pressure 5 cm H2O. Respiratory rate was adjusted to keep end-tidal carbon dioxide (ETCO2) at 4.5–5.5 kPa. Anesthesia was maintained with continuous intravenous infusion of propofol (4 mg/kg/h) and fentanyl (5 μg/kg/h); the infusion rates were adjusted to maintain sufficient depth of anesthesia. Monitoring with electrocardiogram and peripheral oxygen saturation (SpO2) was established. A bladder catheter with thermosensor (Degania Silicone Ltd. Degania Bet 15130, Israel) was inserted.

2.2. Experimental protocol

A catheter was inserted into the left external jugular vein for medicine and saline infusion, and another catheter was inserted into the femoral artery for blood sampling. A third catheter was inserted into the femoral vein for arterial gas analyses. Blood gases were analyzed using an automatic blood gas analyzer (ABL 800 flex, Radiometer, Bronshoj, Denmark).

Two lung biopsies were taken with a stapler (Endo GIF™ Auto SutureTM, Covidien, USA) from the left middle lung lobe, through a small thoracotomy, ensuring that the lung could be inflated afterwards.

A 17- and 19- french catheter (Novaport Vascular Access, Novalung, Germany) were inserted into the femoral artery and vein, respectively. These catheters were connected to an interventional lung assist device (iLA Membrane Ventilator, Novalung Gmbh, Germany) ensuring CO2 removal from the circulation. The device was primed with isotonic saline and the extracorporeal circuit was opened. The blood flow in the extracorporeal circuit was monitored with an ultrasonic flow meter (Novaflow s, Novalung GmbH), placed on the tube from the arterial catheter. Before cannulation 15.000 IE heparin was given for anticoagulation, and then 5.000 IE heparin was given every second hour to reduce the risk of clotting. The gas flow to the iLA was 100% oxygen at 10 L/min. throughout the whole experiment. Apneic oxygenation was established via tracheal gas insufflation through a catheter, connected to the orotracheal tube at one end and connected to a y-piece at the other end. The y-piece was connected to the oxygen supply and to a water manometer ensuring continuous pressure of 20 cm H2O in the system (Fig. 1). The oxygen flow (1–5 L/min) was regulated to keep the water in the manometer bubbling [33,34].

After the preparation, when a steady state had been obtained, a set of baseline values were registered after which the mechanical
ventilation was stopped and the iLA started. Arterial blood gas samples were taken every 30 minutes. A noradrenaline infusion was started (0.5 mg/h) if the systolic blood pressure dropped below 90 mmHg, to ensure a sufficient flow in the iLA. All pigs received noradrenaline infusion during the experimental period. The animals were kept on apneic oxygenation for 5 hours, two biopsies were then taken from the lung, using the same thoracotomy as before. The animals were then euthanized with an intravenous injection of pentobarbitone.

2.3. Sample preparation

Lung biopsy specimens were obtained for histological and metabolite examination, one for each examination, before and after the experiment. The samples taken before the experiment were defined as the baseline time point and the samples taken after the hyperoxia exposure were taken at 300 minutes (Table 1). The specimens for histopathology were initially immersion fixed in 4% buffered formalin and subsequently embedded in paraffin (PAS). This yielded six sections per pig, three from before oxygen treatment and three from after treatment. Furthermore, immunostains for T and B lymphocytes (CD3 and CD20), epithelial markers (CK7), and endothelial markers (CD 34 and Factor VIII related antigen) were applied. Light microscopy was performed by an investigator unaware of which section was from before and after oxygen treatment. The other half of the lung biopsy specimens were flushed with saline and frozen in –80 °C freezer, for subsequent metabolomic experiments. High-resolution magic angle spinning (HR-MAS) nuclear magnetic resonance (NMR) spectra were recorded on a Bruker Avance III 600 MHz NMR spectrometer (Bruker Biospin GmbH, Rheinstetten, Germany) equipped with a triple resonance (1H, 13C, 31P) HR-MAS probehead. Disposable Kel-F inserts with sodium formate solution in D2O (25 mM, 5 μL) and freshly thawed tissue samples (9–16 mg), were placed in 4 mm zirconia MAS rotors. The rotors were spun at 5.000 Hz at 5 °C during recordings. One-dimensional Carr-Purcell-Meiboom-Gill (CPMG) spectra were recorded with 72k data points, 256 scans, 20 ppm spectral width and a total echo time of 77 ms. Water presaturation (25 Hz) was applied during the relaxation delay (4 s). The CPMG NMR experiment is inherently non-quantitative, meaning that the individual metabolite signals are attenuated differently by the T2 filter, according to their transverse relaxation constants. In addition, the longitudinal relaxation (T1) for some nuclei will be too slow for complete relaxation during the combined acquisition time and relaxation delay, including the signal from the formate. However, given the presumption that relaxation properties are similar for all samples in the cohort, CPMG is useful for determining relative differences of metabolite levels.

2.4. Data analysis

CPMG spectra were manually phased and baseline corrected, calibrated to the alanine doublet at 1.48 ppm in TopSpin 3.1 (Bruker BioSpin, Rheinstetten, Germany) and reduced to integrals of 0.001 ppm bin width, excluding the water region (4.5–5.1 ppm) in AMIX (Analysis of MXtures v. 3.9.10, Bruker BioSpin, Karlsruhe, Germany). Data was exported to MATLAB (R2018a, MathWork) and was subsequently normalized (probabilistic quotient normalization) and mean centered.

Unsupervised principal component analysis (PCA) was carried out on processed spectra in PLS Toolbox 6.5 (Eigenvector Research, Wenatchee, WA, USA) to identify possible outliers and trends in the data. Data was visualized by plotting the scores of the first two principal components (PC1 and PC2). Metabolites were assigned based on our previous study [35] and the Human Metabolite Database [36], while relative quantification was determined using the integrals produced by the line shape peak fitting option in MNova (Mestrelab Research v.12) relative to the known formate concentration and tissue weight.

Table 1

|                      | Baseline 0 min | 30 min | 60 min | 90 min | 120 min | 150 min | 180 min | 210 min | 240 min | 270 min | 300 min | p-value (time) | p-value (Baseline vs. 300 min) |
|----------------------|----------------|--------|--------|--------|---------|---------|---------|---------|---------|---------|---------|----------|--------------------------|
| pH                   | 7.45           | 7.44   | 7.36   | 7.35   | 7.34    | 7.32    | 7.31    | 7.30    | 7.29    | 7.28    | 7.27    | 0.01     | 0.03                     |
| PCO_{2} (kPa)        | Mean 5.35      | 5.34   | 5.17   | 5.13   | 5.42    | 5.64    | 6.61    | 6.47    | 6.02    | 6.70    | 6.68    | 6.86     | 0.89                     | 0.15                     |
|                      | SD 0.03        | 0.07   | 0.12   | 0.12   | 0.09    | 0.08    | 0.07    | 0.09    | 0.08    | 0.07    | 0.07    |          |                          |
| PO_{2} (kPa)        | Mean 22.1      | 21.6   | 21.5   | 19.1   | 19.9    | 19.1    | 19.1    | 15.1    | 15.1    | 14.8    | 15.1    | 15.4     | 0.006                    | 0.003                    |
|                      | SD 13.1        | 16.4   | 18.8   | 13.4   | 16.6    | 14.4    | 16.0    | 15.5    | 14.8    | 16.1    | 15.9    | 13.9     |                          |                          |
| Lactate (mM)        | Mean 1.1       | 1.2    | 1.5    | 1.6    | 1.6     | 1.8     | 1.8     | 1.7     | 1.5     | 1.4     | 1.3     | 1.2      | 0.81                     | 0.68                     |
|                      | SD 0.2         | 0.6    | 0.6    | 0.5    | 0.7     | 0.6     | 0.6     | 0.6     | 0.6     | 0.6     | 0.5     | 0.5      |                          |                          |
| Glucose (mM)        | Mean 6.2       | 5.7    | 6.5    | 6.2    | 6.4     | 6.2     | 6.2     | 5.3     | 5.1     | 4.9     | 4.8     | 4.7      | 0.43                     | 0.055                    |
|                      | SD 2.3         | 2.4    | 3.4    | 1.9    | 2.3     | 2.0     | 1.9     | 2.1     | 2.1     | 1.9     | 1.6     | 1.7      |                          |                          |
The statistical analysis was performed in the IBM® SPSS Statistics software (v. 25, SPSS Inc., Armonk, NY, USA). To be able to determine differences between paired samples collected before and after the exposure, we first employed a Shapiro-Wilk test to verify if the data followed a normal distribution. Accordingly, a paired t-test or its nonparametric equivalent Mann-Whitney-Wilcoxon rank test was applied. A calculated two-tailed p-value of less than or equal to 0.05 was considered statistically significant. Data was summarized using mean and standard deviation (SD), while scatter plots were used for visualization purposes.

3. Results

One pig had to be excluded after three hours of hyperoxia due to hemodynamic instability; hence, six pigs were included in this study, (weight; mean ± SD, 45.6 ± 7.9 kg).

3.1. Blood gas measurements

Arterial blood gas measurements performed at different time points before and during the exposure confirmed the presence of hyperoxia (Table 1). At the end of the experiment, pH levels dropped slightly in all animals. The partial pressure of carbon dioxide presented minor increases across the experimental period; however, these changes were within the normal range. As expected, the partial pressure of oxygen was above physiological values during exposure to hyperoxia, with mean values between 52 and 64 kPa, due to the high FiO2 received. The increase in PaO2 was pronounced already from the first arterial blood gas drawn after initiating the iLA and remained above 50 kPa throughout the experiment. Lastly, while arterial blood glucose levels were slightly decreased after 300 minutes of hyperoxia, lactate levels remained mainly unchanged, indicating successful circulation in the pigs during the experimental procedure.

3.2. Lung histological characterization

Lung tissue biopsies were obtained before and after the exposure. Paired samples were evaluated on size, pleural, interstitial and parenchymatous morphology. Large inter- and intra-individual differences were observed in the histopathological sections across the animals, with no visible characteristic changes related to the oxygen treatment received (Fig. 2). No hyaline membranes were detected (Fig. 3). Elastic fibers, T- or B-lymphocytes and epithelial and endothelial cells did not differ between samples taken before and after the oxygen exposure, indicating that vascular permeability was not increased during the five hours of hyperoxia.

3.3. Lung metabolite profiles of hyperoxia

To analyze the metabolic response to hyperoxia, unsupervised PCA was performed on processed spectra. Samples clustered according to the experimental conditions (Fig. 4); with tissue samples collected before and after hyperoxia along the positive principal component 1 (PC1) axis, and tissue samples affected by hyperoxia along the negative PC1. Several metabolites were found to be different between the paired samples (Table 2). The most relevant...
metabolites are visualized in Fig. 5. Tissue glucose and lactate levels were reduced at the end of the experimental period, indicating deranged glycolysis with a subsequent effect on the organism’s energy status, including ATP, adenosine monophosphate (AMP) and inosine monophosphate (IMP) (Fig. 5). In line with these findings, metabolites involved in the tricarboxylic acid (TCA) cycle including citrate, succinate, fumarate, and 2-ketoglutarate were also decreased. The oxidant and antioxidant metabolites hypoxanthine, uracil, uridine, adenine, inosine, aspartic acid, glutathione, and uracil were also found to be altered, indicating an increase in oxidative stress due to imbalance in the oxidant-antioxidant status during hyperoxia. Previous studies have shown that the combination of increased ROS production together with the lack of cellular energy ultimately compromises the organism’s antioxidant defense mechanism [40,41].

We did not induce hypoxia in our research as the method we used (apneic oxygenation) cannot be done with other gasses than oxygen [33]. However, by comparing the literature describing metabolomics studies performed on hypoxic human and animal models with our hyperoxia result, we have found that several metabolite changes were quite contrary, strengthening our findings. For example, our results show that hyperoxia is characterized by energy depletion, imbalance in the oxidant-antioxidant defense, and cell membrane and surfactant dysfunction. During hypoxia however, Heerlein and co-workers have found a reduced energy demand due to a decreased activity in ATP-consuming pathways [42]. Secondly, the levels of glutamate and glutathione were found diminishing after 5 hours of hyperoxia whereas they increased levels of glucose, ATP, AMP and IMP, suppressed TCA cycle, and increased production of ROS. Although the study investigates changes in a pig model, the findings can improve understanding of the biochemical and molecular alterations the lungs undergo during hyperoxic conditions, an event often encountered at intensive care units across the world. Metabolites are the end products of gene, transcript, and protein regulation, and are strongly influenced by environmental changes such as disease progression [35,37] and treatment administered [38,39]. Therefore, changes in their levels should reflect an overall molecular window of events occurring during hyperoxia. Previous studies have shown that the combination of increased ROS production together with the lack of cellular energy ultimately compromises the organism’s antioxidant defense mechanism [40,41].

Table 2

| Metabolites                        | Before (pmol/mg) | After (pmol/mg) | Pair t-/Wilcoxon Signed Ranks Test Sig. (2-tailed) |
|------------------------------------|-----------------|-----------------|---------------------------------------------------|
| Mean SD                            | Mean SD         |                 |                                                   |
| Nicotinate                         | 680 169         | 415 110         | 0.02                                              |
| Adenosine                          | 350 60          | 282 63          | 0.03                                              |
| Inosine                            | 83 8            | 72 13           | 0.06                                              |
| Hypoxanthine                       | 527 60          | 652 158         | 0.03                                              |
| Histidine                          | 22 27           | 262 157         | 0.03                                              |
| Uracil                             | 60 46           | 93 57           | 0.13                                              |
| Phosphatidyl choline               | 158 62          | 337 216         | 0.11                                              |
| Sphingomyelin and Choline          | 302 49          | 522 226         | 0.07                                              |
| Tyrosine                           | 136 36          | 236 149         | 0.07                                              |
| Fumarate                           | 30 8            | 12 7            | 0.004                                             |
| ATP                                | 279 124         | 164 101         | 0.13                                              |
| AMP & IMP                          | 272 49          | 214 65          | 0.01                                              |
| Fatty acyl chain                   | 438 187         | 705 290         | 0.08                                              |
| a-Glucose                          | 1512 374        | 1269 439        | 0.04                                              |
| Glycerocephosphocholine            | 797 189         | 634 186         | 0.01                                              |
| Lactate                            | 1643 364        | 1353 454        | 0.017                                             |
| Myo-inositol                       | 6856 766        | 4792 1220       | 0.03                                              |
| Creatinine                         | 1678 302        | 1353 278        | 0.001                                             |
| Creatine                           | 941 244         | 797 225         | 0.04                                              |
| Glycine                            | 9844 1333       | 8823 2419       | 0.23                                              |
| Taurine                            | 8298 1897       | 7025 2419       | 0.07                                              |
| Trimethylamine-N-oxide             | 2308 304        | 1769 413        | 0.02                                              |
| 2-ketoglutarate                    | 5196 625        | 4492 1039       | 0.12                                              |
| Aspartic acid                      | 631 141         | 488 125         | 0.02                                              |
| Glutathione                        | 842 140         | 664 198         | 0.004                                             |
| Citrate                            | 419 99          | 261 35          | 0.02                                              |
| Succinate                          | 639 58          | 566 108         | 0.04                                              |
| Glutamate                          | 5925 1163       | 4567 1121       | 0.005                                             |
| Glutamine                          | 4340 616        | 3520 730        | 0.002                                             |
| Acetate                            | 338 65          | 416 82          | 0.03                                              |
| Proline & Arginine                 | 725 149         | 952 132         | 0.07                                              |
| Alanine                            | 1405 217        | 2337 411        | 0.002                                             |
| Valine                             | 306 31          | 495 171         | 0.03                                              |
| Isoleucine                         | 180 24          | 219 50          | 0.20                                              |
| Leucine                            | 330 44          | 465 91          | 0.04                                              |

4. Discussion

To our knowledge, this is the first study investigating lung metabolite profiles of hyperoxia. This study demonstrates that despite the relatively unchanged lung histology after a five-hour exposure to hyperoxia, there were clear changes in the small metabolite levels. The general picture that emerges from our metabolomics approach is one of energy depletion, with decreased

levels of oxygen. However, Heerlein and co-workers have found a reduced energy demand due to a decreased activity in ATP-consuming pathways [42]. Secondly, the levels of glutamate and glutathione were found diminishing after 5 hours of hyperoxia whereas they increased...
during hypoxia [43,44]. Moreover, in our study, tricarboxylic acid cycle (TCA) intermediates were found reduced while their levels increased in patients suffering from hypoxemia [43]. Lastly, hypoxanthine and histidine have been found reduced in patients suffering from hypoxemia [35]. In contrast, in our hyperoxia study, these metabolites were found elevated after 5 hours exposure of oxygen. Altogether, the contrasts in these metabolites levels indicate that the presence or lack of oxygen, as is the case of hyperoxia and hypoxia, respectively, influences same metabolite pathways but with opposing effect on metabolite levels.

4.1. Arterial blood gas changes during hyperoxia

Arterial blood gas measurements showed pronounced hyperoxia already from the first measurement after the apneic oxygenation was initiated (Table 1). The partial pressure of carbon dioxide increased during the experimental period and the pH levels dropped accordingly. Even though these changes were within normal ranges, they might have been avoided by increasing the sweep gas flow up to 15 L/min, which is the maximum recommended gas flow for the iLA device [45].

4.2. Lung tissue histological changes during hyperoxia

Healthy humans exposed to a FiO2 of 0.95 at normobaric atmospheric pressure will have a latent period of 4–22 hours before experiencing any symptoms [15]. This latent period is generally accepted as clinically safe for hyperoxia treatment [46]. The first
histological findings after normobaric hyperoxia can be identified after 12–24 hours of exposure, the main findings being capillary congestion and interstitial and intra-alveolar edema [11]. This is consistent with our results from the lung tissue histopathology, where changes related to hyperoxia could not be detected, not even the appearance of a hyaline membrane, which is one of the hallmarks of the acute phase of pulmonary injury. Despite negative results in the levels of inflammatory cells or the epithelial and endothelial cells, in-vitro studies performed on ovine pulmonary artery endothelial cells have shown that within 30 minutes of 100% oxygen exposure there is already increased production of superoxide anions and other radicals [26]. Parinandi and co-workers [25] found increased formation of H2O2 in human pulmonary artery endothelial cells exposed to 100% oxygen for 3 hours.

4.3. Lung metabolite profile of hyperoxia

Glucose is the most important metabolite in glycolysis, involved in the synthesis of ATP and NADH for cellular energy. In this study, glucose levels decreased both in blood and lung tissue samples. Our results confirm previous findings conducted on mouse lung tissue, where glycolytic activity was reduced [40]. The energy metabolites ATP, AMP and IMP were also decreased following hyperoxia, indicating an overall derangement of energy production in the lung tissue.

Imbalance in the levels of oxidants and antioxidants is thought to be one the key players in the pathogenesis of hyperoxic injuries. Since TCA can regulate the production and decomposition of antioxidants [47,48], it is interesting to observe that citrate, fumarate, succinate, and 2-ketogluutaric acid were decreased following the 5 hour period of hyperoxia. Chambellans and Lemires demonstrated that the cell’s immediate response to hyperoxia is downregulating the antioxidant defenses resulting in protective cell cycle arrest to minimize the effects of hyperoxia [49,50], which is in line with our findings. An overall reduction was observed in the levels of the antioxidants nicotinate, inosine, and adenosine, and of the detoxification metabolite glutathione, the most abundant antioxidants present in cells [48], indicating increased consumption of antioxidants, increased production of ROS, or both. These findings have been previously confirmed in studies conducted on human and ovine pulmonary endothelial cells [25,26]. The precursors of glutathione, glutamine and glutamate, were also found to be decreased.

The levels of several amino acids were increased at the end of the hyperoxic period, possibly indicating augmented proteolysis, as suggested previously by Oury and co-workers [20] caused by the severe energy depletion occurring during hyperoxia [40].

Lastly, the lung tissue levels of choline and sphingomyelin, glycerophosphocholine, and phosphatidyl choline were lower after hyperoxia. These metabolites are known to be the main components of the pulmonary surfactants [51] and act as signaling molecules embedded in the cellular membrane [52], hence, our results possibly indicate dysfunctionality in either surfactant production, cell membrane integrity, or both.

4.4. Strength and limitations

The application of both histopathological and NMR studies to estimate the effect of hyperoxia on lung tissue is new and to our knowledge has not been attempted before. The interaction of oxygen with other pathological conditions cannot be evaluated without better knowledge of the injuries caused by oxygen alone. By using an apneic oxygenation technique during the experiments, any effect of the respirator treatment on the development of lung tissue damages could be minimized, hence, the changes observed are indeed caused by hyperoxia. Each pig served as its own control, minimizing the possible individual variation that can add confounding to the data. However, several practical limitations must be acknowledged. Although our study is the first of its kind published to date, sample size is small and the pigs used were young and healthy, whereas patients exposed to hyperoxia are critically ill and of varying ages. Furthermore, we did not have a set of blood samples to investigate metabolite changes as a consequence of hyperoxia. This information would have helped characterize the effect of hyperoxia on the whole metabolome. Additional studies are needed to investigate the whole body response to this stress factor. Increasing evidence shows that liberal oxygen therapy increases mortality in acutely ill adult [53]. With liberal oxygen therapy, the lungs are exposed to high concentrations of oxygen, which increases the risk of acute lung injury. This present porcine study visualizes significant metabolic changes in the lung tissues following exposure to 100% oxygen. In the clinical settings, lung biopsies are not possible; however, the results are important as no specific biomarkers for lung tissue injury exits. New studies using bronchoalveolar lavage and exhaled breath condensate will be needed to confirm the results in acutely ill patients.

5. Conclusion

Despite no histopathological changes being detected in the lung tissue after a period of five hours of hyperoxia, significant changes were observed in the levels of lung tissue metabolites. Hyperoxia is characterized by energy depletion, imbalance in the oxidant-antioxidant defense, possible proteolysis, and cell membrane and surfactant dysfunction. These results clearly indicate that providing FiO2 of 1.0, and hence a high PaO2, affects the lung tissue. Thus, this study emphasizes the need for further metabolite studies in critically ill patients. If the same metabolite pattern can be recognized, it might call for changes in targeting FiO2, and thereby PaO2, in order to minimize the risk of oxygen’s adverse effects.

Author contribution

B.S.R. and B.K. came up with the main research idea and supervised the project. S.O.M. and B.K executed the surgeries together, collected biopsies, and performed blood gas analyses. H.V. prepared the histopathological samples. I.H.B. and U.B. examined the histopathological samples. T.A. and T.F.B. prepared and performed the NMR experiments. R.G.M processed and performed data analysis. R.G.M. and S.O.M. interpreted the results. S.O.M. wrote the first draft of the paper. All authors assisted in drafting the manuscript and approved the final document.

Funding

This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgement

We would like to thank the staff at Biomedical Research Laboratory from Aalborg University Hospital for their support. The HR MAS analyses were performed at the MR Core Facility, Norwegian University of Science and Technology (NTNU). The MR Core Facility is funded by the Faculty of Medicine at NTNU and Central Norway.
relationship connecting oxidative stress with endometriosis. BioMed Res Int 2013;2013. https://doi.org/10.1155/2013/329058.

[49] Chambellan A, Cruickshank PJ, McKenzie P, Cannady SB, Szabo K, Comhair SAA, et al. Gene expression profile of human airway epithelium induced by hyperoxia in vivo. Am J Respir Cell Mol Biol 2006;35:424–35. https://doi.org/10.1165/rcmb.2005-0251OC.

[50] Lemire J, Milandu Y, Auger C, Bignucolo A, Appanna VP, Appanna VD. Histidine is a source of the antioxidant, α-ketoglutarate, in Pseudomonas fluorescens challenged by oxidative stress. FEMS Microbiol Lett 2010;309:170–7. https://doi.org/10.1111/j.1574-6968.2010.02034.x.

[51] Agassandian M, Mallampalli RK. Surfactant phospholipid metabolism. Biochim Biophys Acta 2013;1831:612–25. https://doi.org/10.1016/j.bbalip.2012.09.010.

[52] Ghidoni R, Caretti A, Signorelli P. Role of sphingolipids in the pathobiology of lung inflammation. Mediat Inflamm 2015;2015. https://doi.org/10.1155/2015/487508.

[53] Chu DK, Kim LH-Y, Young PJ, Zamiri N, Almenawer SA, Jaeschke R, et al. Mortality and morbidity in acutely ill adults treated with liberal versus conservative oxygen therapy (IOTA): a systematic review and meta-analysis. Lancet Lond Engl 2018;391:1693–705. https://doi.org/10.1016/S0140-6736(18)30479-3.