Obstructive jaundice induces early depression of mitochondrial respiration in rat hepatocytes

A icterícia obstrutiva induz depressão precoce da respiração mitocondrial em hepatócitos de ratos

Riad Naim Younes, Renato Sergio Poggetti, Belchor Fontes, Mario Matsuo Itinoshe, Vanda Mitie Yoshida, Maria Elisa A. Carvalho, Dario Birolini

1. Research performed at Laboratory of Medical Investigation (LIM-62) and Experimental Research Division (ERD) of the Heart Institute (HI), University of Sao Paulo School of Medicine (USP-SM), Sao Paulo, Brazil.
2. PhD, Assistant Surgeon, Trauma Division, USP-SM, Sao Paulo, Brazil.
3. PhD, Associate Professor, USP-SM, Sao Paulo, Brazil.
4. PhD, Department of Surgery, USP-SM, Sao Paulo, Brazil.
5. Biologist, Researcher at the LIM-62, USP-SM, Sao Paulo, Brazil.
6. Pharmacist, Researcher at the ERD, HI, USP-SM, Sao Paulo, Brazil.
7. MD, Researcher at the ERD, HI, USP-SM, Sao Paulo, Brazil.
8. Full Professor of Surgery, Trauma Division, USP-SM, Sao Paulo, Brazil.

ABSTRACT

Introduction: Oxidative phosphorylation dysfunction of hepatocyte mitochondria is involved in the pathophysiology of organ dysfunction following obstructive jaundice (OJ). However the time period from biliary occlusion to the occurrence of the dysfunction has not been determined decisively. Purpose: To evaluate the early effects (1 d and 7 d) of OJ on liver mitochondria respiratory function in rats. Methods: Male Wistar rats (200-250 g) were randomly divided into the following 3 groups: laparotomy plus OJ for 24 h (1d group) (n = 10); laparotomy plus OJ for 7 d (7d group) (n = 10); sham control procedure (CTR group) (n = 12). At the end of OJ periods, total serum bilirubin level, hepatic enzyme activity levels (GOT, GTP, Gama-GT, ALP), mitochondrial respiration phases S3 and S4, as well as the respiratory control ratio (RC = S3/S4), and ADP consumption/oxygen consumption (ADP/O) ratio, were determined. Results: Total serum bilirubin, activity of most hepatic enzymes, and O2 consumption during basal (S4) respiration were increased in the 1d and 7d groups (ANOVA, p = 0.05 vs. CTR). After ADP addition, the O2 consumption rate (S3) in the 1d group remained similar to the CTR rate (ANOV A p > .05), while the RC rate was reduced (ANOV A, p = 0.001) vs. CTR. The effects observed on mitochondrial respiration in the 1d group were exacerbated in the 7d group. Conclusion: These results indicate that OJ induces early (24 h) depression of liver mitochondria respiration, and thus may lead to early reduction in the production of high energy bonds.

Key words: Jaundice, Obstructive. Mitochondria, Liver. Oxidative Phosphorylation.

RESUMO

Introdução: A disfunção da fosforilação oxidativa das mitocôndrias do hepatócito está envolvida na fisiopatologia da disfunção orgânica subseqüente à icterícia obstrutiva (IO). Entretanto, a precocidade da ocorrência desta disfunção permanece obscura. Objetivo: Avaliar o efeito precoce da IO na função respiratória mitocondrial em ratos. Métodos: Ratos Wistar machos (200 a 250g) foram randomizados em 3 grupos que foram submetidos a laparotomia mais: IO por 24 h (grupo 1d)(n=10); IO por 7 dias (grupo 7d)(n=10); procedimento simulado (grupo CTR)(n=12). Ao final dos períodos de IO, foram determinados: bilirrubina sérica total, atividade de enzimas hepáticas (TGO, TGP, Gama-GT, ALP), respiração mitocondrial fases S3 e S4, assí como a razão do controle respiratório (RC = S3/S4), e ADP consumo/oxigênio consumo (ADP/O) ratio, foram determinados. Resultados: Total serum bilirrubin, activity of most hepatic enzymes, and O2 consumption during basal (S4) respiration were increased in the 1d and 7d groups (ANOVA, p = 0.05 vs. CTR). After ADP addition, the O2 consumption rate (S3) in the 1d group remained similar to the CTR rate (ANOVA p > .05), while the RC rate was reduced (ANOVA, p = 0.001) vs. CTR. The effects observed on mitochondrial respiration in the 1d group were exacerbated in the 7d group. Conclusão: Estes resultados indicam que a icterícia obstrutiva induz depressão precoce (24h) da respiração mitocondrial, e pode assim levar à redução da produção de ligações de alta energia.

Introduction

The high morbidity and mortality rates associated with the surgical treatment of patients with obstructive jaundice (OJ) have frequently been attributed to metabolic disorders, specifically disorders of the oxidation-reduction function of hepatocyte mitochondria.1-2 It was shown in rats in the 1960s, that elevated bilirubin levels were associated with altered oxidative phosphorylation of liver mitochondria. This phosphorylation disturbance was attributed to a possible affinity of bilirubin for lipid binding sites in the mitochondrial membrane.3 Mitochondria isolated from the hepatic parenchyma of jaundiced rats were found to exhibit greater dysfunction than mitochondria extracted from hepatocytes, implicating the involvement of extramitochondrial mediators in this dysfunction.4 Other potential factors for this mitochondrial dysfunction include biliary infection,5,6 degradation of mitochondrial phospholipids caused by the increased biliary salts in OJ,7 and the increased expression of uncoupling protein 2.8 In the 1980s it was observed that alterations in hepatocyte mitochondrial respiratory function in jaundiced rats were amplified with increased duration of biliary occlusion,9 such that ATP production was reduced by 25% in the first and second weeks, and by 42% in the third week after biliary occlusion. This progressive depression in ATP production was accompanied by an increase in the serum concentrations of liver enzymes (AST, ALT, GGT, ALP).10 Several independent research groups have reported evidence of mitochondrial dysfunction onset as early as 3 d after biliary occlusion. However, it has not yet been clearly established how early after biliary occlusion that this dysfunction first becomes detectable. The objective of the present study was to examine the effects of OJ on mitochondrial function as well as on serum bilirubin and liver enzyme levels in rats submitted to bile duct occlusion at a very early post-occlusion time point (24 h).

Methods

All procedures were performed in accordance with the norms of the Institution Ethical Committee for animal care and use in experimental research. The rat is the most widely used animal in studies on liver mitochondrial dysfunction, and therefore was used in this study. Male Wistar rats, weighing 250-300 g, were submitted to anesthesia with intraperitoneal diazepam + ketamine injection and divided into three groups: 1) control group (CTR) (n = 12) subjected to simulated procedures only; 2) 1d group, subjected to OJ for 24 h (n = 10); 3) 7d group, subjected to OJ for 7 d (n = 10). The anterior abdomens of Group 1d and 7d animals were prepared for surgery by shaving and cleaning with iodopovidone. Rats were then subjected to a median laparotomy, during which the common bile duct was identified, isolated, and then ligated at two adjacent points using a 4-0 cotton thread. The duct was then sectioned between the ligatures, in order to provoke OJ. The OJ animals were then maintained under observation in individual cages, given standard laboratory rat chow and water ad libitum for 24 h (group 1d) or 7 d (group 7d), at which time the animals were sacrificed by injection of an overdose of Sodium pentobarbital. The rats livers were then immediately removed for analysis of hepatocyte mitochondrial function, and blood samples were collected for biochemical assessments (total bilirubin, AST, ALT, GGT, ALP). The animals of the CTR group were submitted to the same procedures as group 1d and 7d animals, except for the ligation and sectioning of the common bile duct. Dissected livers were immediately transferred to a Potter homogenizer, in a homogenizing solution (220 mM mannitol; 70 mM sucrose; 5 mM MES pH 7.4; 2 mM EGTA and 0.2% BSA) on ice, and homogenized. Lowry’s method12 was used to determine protein content in the mitochondrial fraction. After centrifugation at 1,500 x g of the homogenate, the supernatant was centrifuged at 10,000 x g in a centrifuge refrigerated to 0°C, and the resulting pellet was suspended in a solution of the following composition: 220 mM mannitol, 70 mM sucrose, 5 mM MOPS pH 7.4. Mitochondrial function was evaluated by polarography, using an O2 monitor (Yellow Spring Instr.) at 28°C. The standard variables of mitochondrial respiration assessed were the ATP synthesis, the state 4 (S4) and state 3 (S3) of mitochondrial respiration, the S3/S4 respiratory control (RC) ratio, and the ADP to oxygen consumption (ADP/O) ratio. Incubation medium contained 80 mM KCl, 50 mM MOPS pH 7.4, 4.5 mM potassium phosphate, 0.1% BSA in a 2 mL total volume. Glutamate-malate was the substrate employed. Oxidative phosphorylation (S3) was started by addition of adenosine diphosphate (ADP, 280 nmol/mL). S3 and S4 phases and respiratory control ratio RC = S3/S4, as well as the ADP consumption per consumed oxygen ratio (ADP/O) were determined. Total bilirubin and hepatic enzyme levels were measured by means of standard laboratory tests.13

Results

As summarized in Table 1, there was a significant increase in O2 consumption during basal respiration (S4) in mitochondria preparations of the livers from OJ rats (ANOVA, p = 0.009). Following stimulation with ADP, the velocity of O2 consumption (S3) in OJ cells was not different from that of CTR cells (ANOVA, p = 0.126), but the RC value was significantly lower than that observed in the CTR group (ANOVA, p ≤ 0.001).
Production of energy constitutes one of the main functions of the hepatocyte, which, through the mitochondrial respiratory chain, performs oxidative phosphorylation and ATP synthesis. Reduction in this energy production is a basic component of the pathophysiology of severe OJ.10 In the present study, we found that total serum bilirubin and hepatic enzyme activity levels were elevated 24 h after biliary obstruction. This is 2-3 d earlier than observations reported previously.2,6,7,10,14 Table 1 displays the mitochondrial respiratory dysfunction in jaundiced rats hepatocytes.

**TABLE 1 - Mitochondrial respiratory dysfunction in jaundiced rats hepatocytes**

<table>
<thead>
<tr>
<th>GROUP</th>
<th>CTR 1d OJ</th>
<th>7d OJ</th>
</tr>
</thead>
<tbody>
<tr>
<td>S3</td>
<td>197±9.1</td>
<td>190±9.2</td>
</tr>
<tr>
<td>S4</td>
<td>24±1.0</td>
<td>28±0.9*</td>
</tr>
<tr>
<td>RC</td>
<td>8.0±0.2</td>
<td>6.6±0.1*</td>
</tr>
<tr>
<td>ADP/O</td>
<td>2.7±0.2</td>
<td>3.1±0.2</td>
</tr>
</tbody>
</table>

S4: basal conditions; S3: in the presence of ADP; RC: respiratory control (S3/S4); ADP/O: ADP consumption/oxygen consumption rate. Data were determined for day 1 (ICT 24h), day 7 (ICT 7d), and control (CTR) group. *p < 0.05 for increase of S4; or decrease of S3, RC, and ADP/O, compared to corresponding CTR values.

The ADP/O ratio was not reduced in the 1d group. All evaluated parameters of the respiratory chain (S3, S4, RC and ADP/O ratio) were significantly altered by 7 d after biliary occlusion. Total serum bilirubin concentration significantly increased in the first 24 h after biliary obstruction, relative to the CTR group, and was similar to that observed on the seventh day of biliary obstruction (Table 2). Furthermore, hepatic enzymes activity levels were significantly increased 24 h after induction of biliary obstruction relative to the CTR group (Table 2).

**TABLE 2 - Disturbances induced by OJ on total bilirubin and enzymes activity**

<table>
<thead>
<tr>
<th>GROUPS</th>
<th>CTR</th>
<th>1d</th>
<th>7d</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bilirubin (mg/dL)</td>
<td>0.56±0.42</td>
<td>4.2±1.7*</td>
<td>4.5±3.1*</td>
</tr>
<tr>
<td>AST (TGO) (U/L)</td>
<td>60.9±20.4</td>
<td>557.3±153.1*</td>
<td>223.7±225.9*</td>
</tr>
<tr>
<td>ALT (TGP) (U/L)</td>
<td>43.0±11.8</td>
<td>399.0±188.2*</td>
<td>87.1±51.7***</td>
</tr>
<tr>
<td>GGT (U/L)</td>
<td>2.1±1.7</td>
<td>12.4±11.6**</td>
<td>18.9±24.6</td>
</tr>
<tr>
<td>ALP (U/L)</td>
<td>319.9±76.5</td>
<td>1066.3±224.9*</td>
<td>696.0±159.4*</td>
</tr>
</tbody>
</table>

Control group (CTR), Day 1 (ICT 24h – 1d) and Day 7 (ICT 7d). (ANOVA, p<0.05) *p < 0.05 ANOVA vs. CTR, **p=0.084 ANOVA vs. CTR, ***p=0.082 ANOVA vs CTR.

Discussion

Production of energy constitutes one of the main functions of the hepatocyte. The mitochondrial respiratory chain, which, through oxidative phosphorylation and ATP synthesis. Reduction in this energy production is a basic component of the pathophysiology of severe OJ.10 In the present study, we found that total serum bilirubin and hepatic enzyme activity levels were elevated 24 h after biliary occlusion. This is 2-3 d earlier than observations reported previously.2,6,7,10,14 In basal conditions S4 represents the O2 consumption rate after ATP synthesis, and reflects the degree of coupling of mitochondrial respiration to ATP production. Meanwhile S3 represents the O2 consumption rate in the presence of ADP and ADP/O consumption/oxygen consumption rate. Intact mitochondria in a milieu containing NADH, O2 and Pi, and a respiratory substrate (e.g. malate) will cease ATP synthesis as soon as endogenous ADP is exhausted. Addition of ADP rapidly restores the process, such that the S3 state is restored and ATP synthesis is resumed. Thus the S3/S4 ratio termed RC reflects the degree of coupling of mitochondrial respiration to ATP production. Meanwhile the ADP/O ratio indicates the amount of synthesized ATP per consumed oxygen (mol ATP/mol O2) and can reveal O2 inefficiency in oxidative phosphorylation.215 We observed a significant increase in O2 consumption during basal respiration (S4) after 24 h of biliary occlusion. The observed decrement in the RC ratio indicates that although the rate of O2 consumption by mitochondria was maintained during stimulation with ADP (S3), the utilization of O2 for ATP generation was compromised in the cholestatic liver tissue of the 1d group. Although S3 depression has been reported in diverse studies of liver mitochondrial dysfunction in rats7 Nagano et al reported mitochondrial dysfunction from the 3rd to the 21st day of biliary obstruction evidenced by an S4 increase and an RC reduction, in the absence of an alteration in S3.6 Thus our 1d group findings here are consistent with Nagano et al’s findings of increased S4 with an unchanged S3. In the present study, the alteration in mitochondrial function observed on the 1st day of biliary obstruction became more marked on the 7th day as evidenced by the persistence of RC and S4 alterations, and the addition of decreases in S3 and ADP/O ratio. Worsening of alterations in mitochondrial function with long-term biliary obstruction is commonly reported in the literature.6,7,9,16 Increased O2 consumption during S4 with decreased RC is indicative of respiratory chain depression, which ultimately leads to reduction in the production of high energy bonds (ATP). Our observation of increased S4 before an alteration in S3 could be observed seems to indicate a certain degree of uncoupling of the mitochondrial chain, potentially attributable to the presence of uncoupling agents. Uncoupling agents (e.g. 2,4-dinitrophenol) allow continuation of NADH oxidation and O2 reduction at high levels, but do not support ATP synthesis, thereby altering mitochondrial function.15,17
Conclusion

The results of this experimental study in rats show that 24 h after its induction, OJ provokes increased O₂ consumption during the S4 phase of liver mitochondria respiration, as well as decreased RC, indicative of respiratory chain depression. This early respiratory chain depression may lead to early reduction in the production of high energy bonds (ATP) by liver mitochondria. This early alteration in liver mitochondrial function was concomitant with increased serum bilirubin levels and increased hepatic enzyme activity levels.

References


Correspondence:
Belchor Fontes
Rua Itacema, 246/11
04530-050 São Paulo – SP Brazil
Phone: (55 11)8109-0033 / 3079-3048
belchor@uol.com.br

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