Effects of hypobaric hypoxia on adenine nucleotide pools, adenine nucleotide transporter activity and protein expression in rat liver

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INTRODUCTION

The liver is the largest metabolic organ in the body. It performs a number of important and complex biological functions that are essential for survival. It also plays important roles in metabolism of carbohydrates, proteins, lipids, drugs, as well as in bile formation and secretion. Energy metabolism is closely related to these normal functions of liver. Mitochondria are the “energy factory” of cells. The adenine nucleotide transporter (ANT) is the most integral protein in inner mitochondrial membrane and consists of two identical subunits of 32 KD. It catalyzes the transporter of cytosolic ADP and mitochondrial ATP in the process of phosphorylation. Consequently, ANT is an important link between the cytosolic energy consumption and mitochondrial energy process yield. Hypoxia could influence mitochondrial oxygenation respiration function and F0-F1 ATPase activity of rat brain. However, little is known about the relationship between the effects of hypobaric hypoxia on mitochondrial energy metabolism changes and ANT function in rat liver exposed to hypoxia. We therefore used...
HPLC, isotopic assay and Western blot to examine the inner- and extra-mitochondria adenine nucleotide pool, ANT activity and its total protein level of rat liver exposed to hypobaric hypoxia.

**MATERIALS AND METHODS**

**Chemicals**

[2,8-³H] ADP was obtained from Perkin-Elmer. Atractyloside (ATR), adenosine-5'-diphosphoric acid (ADP), albumin bovine serum (BSA), nitroblue tetrazolium chloride (NBT), 5-bromo-4-chloro-3-indole-phosphate (BCIP) and mouse anti-goat IgG-alkaline phosphatase (IgG-AP) were supplied by Sigma. Goat anti-human ANT polyclonal antibody was purchased from Santa Cruz.

**Animals and treatments**

Adult male Wistar rats (150-200 g) were used in the experiments. The rats were exposed to a hypobaric chamber simulating 5000 m high altitude for 23 h every day for 0 (H0), 1 (H1), 5 (H5), 15 (H15) and 30 d (H30) respectively. Animals were fed with laboratory chow and tap water ad libitum. Rats were sacrificed by decapitation at sea level (H0 group) and hypobaric chamber simulating 4000 m high altitude (H1, H5, H15, H30 groups). After decapitation the liver tissues were immediately excised, and part of it was frozen in liquid nitrogen, and then manually homogenized by 15 up and down strokes with Teflon glass pestle. The liver mitochondria were isolated by centrifugation as described above. The results were expressed as nmol/mg tissue.

**Assay of mitochondria adenine nucleotide pool**

Adenine nucleotides were separated and quantitated by HPLC[1,2]. Briefly, 500 mmol/L ice-cold HClO₄ was added to 200 μL liver mitochondria suspension and after 5 min incubation at 4 °C, the liquid was centrifuged (12 000 r/min, 20 min) and the supernatant was saved and neutralized with 1 mol/L K₂CO₃ to pH 6.5-7.0. The liquid was centrifuged (12 000 r/min, 10 min) at 4 °C again and maintained frozen at -70 °C until supernatant was analyzed by HPLC. The results were expressed as pmol/minute per milligram mitochondrial protein.

**Assay of liver tissue adenine nucleotide pool**

The liver tissues stored at -70 °C were rapidly transferred into ice-cold 0.5 mol/L HClO₄, chopped finely with scissors, and then manually homogenized by 10 up and down strokes with Teflon glass pestle. The homogenate was placed at 4°C for 5 min, centrifuged (12 000 r/min, 20 min) at 4 °C. The supernatant was saved and neutralized with 1 mol/L K₂CO₃ to pH 6.5-7.0. The liquid was centrifuged (12 000 r/min, 10 min) at 4 °C again and maintained frozen at -70 °C until supernatant was analyzed by HPLC as described above. The results were expressed as nmol/mg tissue.

**¹H-ADP label and liquid scintillation**

The activity of ANT was determined at 4°C by isotopic technique[3,4]. ¹H-ADP label was stopped by atractyloside, a specific inhibitor of ANT. Fifty microliter mitochondrial suspension solution was diluted by 150 μL ice-cold isolation medium. Twenty microliter ¹H-ADP (specific activity 33.9 Ci/mmol ADP) 0.3 μmol/L were added. After 10 s incubation at 4 °C, the reaction was inhibited by 50 μL 3.2 nmol/L ATR, then centrifuged (12 000 r/min, 20 min) at 4 °C. The precipitation was dissolved with 1 ml ice-cold isolation medium, centrifuged (12 000 r/min, 20 min) at 4 °C again for 20 min and the process was repeated three times. The pellet was digested at 70 °C with 7 mol/L HClO₄·200 μL, 8.8 mol/L H₂O·400 μL for 40 min. The 200 μL sample was dissolved in 1 ml scintillator (5 g PPO was dissolved by 700 mL dimethylbenzene and 300 mL anhydro-ethanol) and measured in liquid scintillation counter. ANT activity was calculated from the radioactivities of ¹H-ADP (count per min (cpm)). Nonspecific binding of ¹H-ADP to mitochondria was determined by incubation of mitochondrial samples with 50 μL 3.2 nmol/L ATR prior to addition of 0.3 μmol/L ¹H-ADP. The results were expressed as pmol ADP/min per milligram mitochondrial protein.

**Western blot**

The rat liver mitochondrial ANT protein level was determined by Western blot[5,6]. The mitochondria sample was solubilized in sample buffer and supplemented with 5 μL β-mercaptoethanol. Following that, it was heated for 10 min at 100 °C and supplemented with 5 μL β-mercaptoethanol again. Electrophoresis was performed on a 12% polyacrylamide slab gel with Tris/glycine running buffer. The staking gel contained 5% polyacrylamide. The lanes were loaded with 30 μL aliquots of solubilized mitochondria (10 μg). After separation the protein bands were stained with Commassie blue for 4 h or heated for 10 min at 100 °C and transferred to polyvinylidene difluoride (PVDF) sheet for 90 min. The PVDF sheet was washed twice with Tris-buffered saline (TBS) containing 0.1% Tween-20 (0.1% TBST), pH 7.4 at room temperature. ANT polyclonal antibody at a 1:1000 dilution was used. Incubation was performed in 0.1% TBST, pH 7.4 for at least 12 h at 4 °C. The sheet was washed six times with 0.1% TBST at room temperature. Mouse anti-goat IgG coupled with alkaline phosphatase was used as secondary antibody. Incubation was performed in 0.1% TBST, pH 7.4 for 1 h at room temperature. The sheet was washed six times with 0.1% TBST again. Antigen was visualized by luminescence (NBT/BCIP). Signals were quantified with Smartview (Furu...
Table 1 Changes of total adenine nucleotide pool in liver tissue during hypoxia exposure (nmol/mg tissue, mean ± SD)

| Group | n | AMP      | ADP      | ATP       |
|-------|---|----------|----------|-----------|
| H0    | 6 | 9.32 ± 1.63 | 9.02 ± 0.87 | 4.00 ± 0.26 |
| H1    | 6 | 7.14 ± 1.41  | 5.53 ± 0.71   | 2.40 ± 0.29   |
| H5    | 6 | 4.90 ± 0.74   | 4.57 ± 0.48   | 2.41 ± 0.17   |
| H15   | 6 | 6.08 ± 1.57   | 5.40 ± 1.31   | 3.11 ± 0.25   |
| H30   | 6 | 6.38 ± 1.19   | 6.21 ± 1.36   | 3.27 ± 0.30   |

*P < 0.05, *P < 0.01 vs H0 group.

Table 2 Changes of intra-mitochondrial adenine nucleotide pool of rat liver during hypoxia exposure (nmol/mg protein, mean ± SD)

| Group | n | AMP      | ADP      | ATP       |
|-------|---|----------|----------|-----------|
| H0    | 6 | 8.64 ± 1.57 | 11.34 ± 1.97 | 5.47 ± 0.54 |
| H1    | 6 | 3.94 ± 0.84  | 5.07 ± 1.05  | 4.50 ± 0.35  |
| H5    | 6 | 3.77 ± 0.67   | 8.70 ± 2.11   | 4.72 ± 0.60   |

*P < 0.05, *P < 0.01 vs H0 group.

Table 3 Changes of adenine nucleotide pool in extra-mitochondria of rat liver during hypoxia exposure (nmol/mg tissue, mean ± SD)

| Group | n | AMP      | ADP      | ATP       |
|-------|---|----------|----------|-----------|
| H0    | 6 | 8.64 ± 1.58 | 8.08 ± 0.93  | 3.45 ± 0.22 |
| H1    | 6 | 6.77 ± 1.40  | 5.07 ± 0.74  | 2.01 ± 0.27  |
| H5    | 6 | 4.63 ± 0.82   | 4.09 ± 0.60   | 2.01 ± 0.15   |
| H15   | 6 | 5.68 ± 1.60   | 4.90 ± 1.34   | 2.67 ± 0.23   |
| H30   | 6 | 6.21 ± 2.01   | 5.37 ± 1.53   | 2.80 ± 0.33   |

Science and Technology Co., Ltd., Shanghai, China).

Statistical analysis
Results were expressed as mean ± SD. Significant difference was determined by one-way ANOVA followed by LSD test between different groups. Statistical analyses were performed using SPSS 12.0 software.

RESULTS

Changes of adenine nucleotide pool in rat liver tissue during hypoxia exposure

Compared with H0 group, liver tissue ATP and ADP content decreased significantly in all hypoxia groups (P < 0.01). Tissue AMP in groups H5, H15, and H30 was significantly lower than in H0 group (P < 0.05 and P < 0.01), while H1 group had no marked change compared to H0 group (Table 1).

Changes of intra-mitochondrial adenine nucleotide pool of rat liver during hypoxia exposure

Compared with H0 group, intra-mitochondrial ATP content decreased in all hypoxia groups, which in H1 group was 70.6% of that in H0 group (P < 0.01) reaching the lowest point. Intra-mitochondrial ADP and AMP levels showed the same change and were significantly lower in H0 group than that in all hypoxia groups (P < 0.01) (Table 2).

Changes of adenine nucleotide pool in extra-mitochondria of rat liver during hypoxia exposure

Compared with H0 group, extra-mitochondrial ATP and ADP content decreased significantly in all hypoxia groups (P < 0.05 and P < 0.01). Extra-mitochondrial AMP in groups H5, H15, and H30 was significantly lower than that in H0 group (P < 0.01), while in H1 group it had no marked change compared with H0 group (Table 3).

Change of mitochondrial ANT activity of rat liver during hypoxia exposure

Compared with H0 group, the activity of ANT decreased significantly in all hypoxia groups, which in H5 group was 55.7% of that in H0 group (P < 0.01), being the lowest. Activity in H30 group was higher than that in H15 group (P < 0.01), but was still lower than in H0 group (P < 0.01) (Figure 1).

Effect of hypoxia on mitochondrial ANT protein expression

ANT protein expression in H5, H15, and H30 groups, compared with H0 group, decreased significantly, which in H5 group was the lowest point (27.1%) (P < 0.01). The expression in H30 group was higher than that in H15 group (P < 0.01), but was still lower than in H0 group (P < 0.01) (Figures 2 and 3).

DISCUSSION

ATP is the direct energy for cell usage. Mitochondrial ATP level is influenced by two factors. First, mitochondrial ATP is produced by oxidative phosphorylation. Mitochondria oxidative respiration and phosphorylation states are the main factors that affect the ATP level. Second, mitochondrial ATP provides energy for cytoplasm as well as for its own demand such as the synthesis of mitochondrial DNA, RNA and proteins, etc. Our previous work showed that during hypoxia exposure, mitochondrial...
ATP content and Fo−F1 ATPase activity of rat brain decreased significantly compared with control [4,7]. This indicates that hypoxia could influence ATP production and then the mitochondrial ATP level. However, there were no reports about the hypobaric hypoxia effect on rat liver mitochondrial adenine nucleotide pool.

The current results showed that ATP level in rat liver mitochondria reduced during hypoxia exposure, which in H1 group was the lowest point, 70.6% of control (P < 0.01). The decrease of mitochondrial ATP content may be related to the following factors. First, mitochondrial oxidative respiratory function was inhibited. Our previous study showed that hypobaric hypoxia inhibited the oxidative respiratory function of mitochondria in rat brain[1,6]. Our results (data not shown) also revealed that hypoxia significantly decreased the mitochondrial three state oxygen consumption and respiratory control rate in rat liver, while significantly increased the rat liver mitochondrial four state oxygen consumption. Second, hypoxia decreased the mitochondrial membrane potential (MMP). Our results (data not shown) demonstrated that hypoxia inhibited the MMP of rat hepatocytes. However, MMP is the motive power of mitochondrial ATP synthesis. Thirdly, hypoxia lowered the mitochondrial Fo−F1 ATPase activity[14,15]. Fourthly, the lower mitochondrial ATP content may be also related to the reduction of intra-mitochondrial ADP concentration[1], intra-mitochondrial Ca2+ content[8] and extra-mitochondrial adenine nucleotide pool[10,11].

ANT is the most integral protein in inner mitochondrial membrane and consists of two identical subunits of 32 KD[9]. It is a key energy link between the mitochondrial and cytoplasm since it catalyses the transmembrane exchange between ATP synthesized by the F1−Fo ATP synthase inside mitochondria and ADP generated by the metabolism in cytoplasm[16,17]. The ADP/ATP exchange follows the Michaelis-Menten kinetics and ANT activity is moderate, 1500−2000 molecules per min[14]. Schonfeld et al reported that matrix adenine nucleotides and the ANT protein content are associated with the changes of the ANT activity in rat heart mitochondr[9]. Rufis et al also reported that matrix adenine nucleotide concentration influenced the ANT activity in rabbit liver mitochondr[13]. However, little is known about the effect of hypobaric hypoxia on mitochondrial ANT activity in rat liver.

Our study showed that the activity of ANT after hypoxia exposure decreased significantly. The ANT activity was the lowest point, 55.7% of control after hypoxia exposure for 5 d (P < 0.01), while after hypoxia exposure for 30 d it was higher than after 15 d exposure (P < 0.01), but was still lower than control. This indicates that hypoxia could inhibit the mitochondrial ANT activity in rat liver. The decrease of ANT activity may be related to the following factors. First, the ANT protein level and content was the main factor. Our results showed that ANT protein expression decreased significantly in H5, H15, H30 groups, which in H5 group was 27.1% of that in H0 group. Second, MMP also influenced the ANT activity[3]. Passarella et al reported that helium neon laser increased the rate of ADP/ATP exchange through increasing the MMP in rat liver[15]. In the presence of an MMP of about 100 mV positive inside, the rates of [14C] ATPp,ATPm exchanges were stimulated[9]. All these indicate that MMP is one of the most important factors that affect the ANT activity. The mechanism that decrease of MMP reduced ANT activity is not clear. Thirdly, the change of ANT conformation also influenced its activity. ANT has two conformational states, cytosolic conformation (c-conformation) and matrix conformation (m-conformation)[16,17]. ANT is not a pore, which opens or closes simply as a response to stimuli. Conformational changes have to occur to release nucleotides to the matrix (and the reverse) without creating leakage in the membrane. Fourthly, it was reported that the size of mitochondrial adenine nucleotide pool influenced the ANT activity. Previous studies showed that the postnatal increase in the matrix adenine nucleotides concentration contributed to the increase of ANT activity in rat liver[13] and heart[9].

The decrease of extra-mitochondrial ATP level influences the mitochondrial carrier family including ANT synthesis and transport. Extra-mitochondrial ATP level is mainly determined by the ANT activity. However, the lower ANT protein level has an identical role in influencing ANT activity during hypoxia. So the ANT activity−ANT protein level− ATP content form the vicious cycle and aggravate the dysfunction of cell energy metabolism during hypoxia exposure.

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