Abstract. The aim of the present study was to investigate the effect of one week dehydration heat exposure on thoracic aorta reactivity in rats. Eighteen Male Sprague-Dawley rats were randomly divided into 3 groups (n=6 each group): Control group (CN), heat exposure group (HE), dehydration heat exposure group (DHE). The CN group was exposed to a room temperature of 24˚C, while the HE and DHE groups were exposed to a heat temperature of 32˚C. After 7 days of heat exposure, the heart rate and blood pressure of the rats were measured, and the noradrenaline (NA)-induced contraction on the aorta rings was measured by tension recording. The average contents of malondialdehyde (MDA) and superoxide dismutase (SOD) in serum were detected using ELISA. The expression of apoptotic genes in the thoracic aorta was measured using RT-PCR. Compared with CN, the heart rate in the HE and DHE groups had a tendency to become retarded, but there was no significant difference (P>0.05). In the HE group, the systolic blood pressure (SBP), diastolic blood pressure (DBP) and mean arterial pressure (MAP) of the rats were significantly higher than that of the CN (P<0.05). In the DHE group, the SBP of rats was significantly higher than that of the CN (P<0.05), while the SBP, DBP, and MAP of the rats were decreased compared to the rats in the HE group, although there was no statistical significance (P>0.05). In the HE and DHE groups, the NA-induced contraction on the rats thoracic aorta ring was larger than that of the CN (P<0.05), albeit there was no significant difference between the HE and DHE groups (P>0.05). The serum SOD content decreased in the HE and DHE groups, however, the reduction was significant only in the DHE group (P<0.05). The content of MDA in serum was significantly increased in the DHE group (P<0.05). The expression of BAX was significantly upregulated whereas Bcl2 expression was decreased in the DHE group (P<0.05). The results showed that a high temperature was harmful to the body, especially in the case of lack of food and water. Additionally, the heat exposure elevated blood pressure, and increased arterial reactivity, which were related to the elevated production of MDA, led to the impaired production of SOD, and an increase of cell apoptosis. These findings are useful to understand the influence of dehydrated heat exposure on the vascular function, and they provide certain theoretical and experimental guidance for protection under high temperature.

Introduction

High temperature has a severe influence on an individual's work, life and body, and it is easy for the individual to become exhausted, irritable and exasperation. A high temperature constitutes a risk factor for the occurrence of cerebrovascular, heart and respiratory diseases; consequently, the death rate increases correspondingly, particularly among the elderly (1-3). At present, the effect of a high temperature especially the dehydration of thermal on the physiological function of human, is lacking in terms of comprehensive knowledge and understanding.

On the basis of heat stress, if the water intake is stopped, the extracellular fluid is reduced more rapidly and becomes DHE (4). For some workers such as firefighters, the damage caused by high temperature is inevitable. Recent findings have shown that dehydrated heat exposure elevated blood pressure over a long period of time increased the viscera index of spleen, heart, thymus gland, hypothalamus and pituitary to alleviate the harmful effects of high temperature on the body (5). However, the physiopathological mechanism involved in dehydration heat exposure on the cardiovascular system remains unknown. Thus, the present study focused on whether the influence of dehydrated heat exposure on the function of thoracic aorta is useful to understand the altered function of blood vessels. This study discussed the influence of dehydrated heat exposure on the thoracic artery of rats by studying the blood pressure, arterial reactivity, superoxide dismutase (SOD) and malondialdehyde (MDA) serums and apoptotic function.
Materials and methods

Animals and heat exposure protocol. Eighteen male Sprague-Dawley rats, weighing 180-200 g, were purchased from the Laboratory Animal Center of Ningxia Medical University (Ningxia, China). The experimental procedures of the present study were approved by the Animal Ethics Committee of Ningxia Medical University and Use Committee, in accordance with the guidelines of the Council of the Physiological Society of China.

Eighteen rats were randomly divided into control group (CN), heat exposure group (HE) and dehydration heat exposure group (DEF) (n=6/group). Rats in the CN group were fed at room temperature (25±1°C) throughout the study and were provided with food and water ad libitum. Rats in the HE and DHE groups received a fixed 8 h (9:00-17:00) heat exposure process per day, and in the DHE group, the rats were fasted during the exposed time. Exposure was finished inside the artificial climate chamber with a temperature of 32°C (relative humidity of 60±5%), after exposure, and the rats were kept at room temperature (25±1°C). The behavior of the animals was observed in the process of the whole experiment.

Heart rate and blood pressure. Heart rate and blood pressure were collected using blood pressure monitor (BP-2010A; Softron Beijing Biotechnology Co., Ltd., Beijing, China), and the data were obtained directly from the machine. After the heat exposure, we measured the heart rate and blood pressure that of the thoracic aorta.

Thoracic aorta reactivity. After anesthesia, the chest of the rats was immediately opened, and the thoracic aorta was removed and placed in a paraffin plate filled with physiological saline. Connective tissues were excised carefully, and vascular rings were made (4-5 mm wide). The vascular ring was quickly hung in the organization bath systems with the presentation of 10 ml Krebs solution [ingredients (mmol/l): 5.6 glucose, 10 NaCl, 24.8 NaHCO₃, 4.6 KCl, 2.5 CaCl₂, as well as 1.2 MgSO₄ and KH₂PO₄, respectively]. The system was perfuse with 5% CO₂ and 95% O₂ continguously and maintained a constant temperature of 37°C. Resting tension was adjusted to 1 g, and the ring was balanced for 40 min with Krebs changed every 15 min. The maximal contraction was induced by the addition of 60 mM KCl. After resting tension was stabilized, the Krebs fluid was replaced and basal tension was returned to 1 g, and cumulative concentrations of norepinephrine (10⁻¹⁰-10⁻⁵ M) was added to the bath system. The rates of the vascular tension range induced by noradrenaline (NA) (10⁻¹⁰-10⁻⁵ M) were expressed as percentages of the maximum contraction tension range (100%) induced by KCl (60 mM).

Detection of the serum SOD and MDA. Blood was collected from the left atrium and centrifuged at 3,500 x g for 15 min and the serum was segregated for further detection. The serum SOD was detected using a commercially available sandwich ELISA kit (Chenglin Biotechnology, Beijing, China). The MDA was detected by TBA kit (Nanjing Jiancheng Bioengineering Research Institute, Jiangsu, China). All the detections were tested in accordance with the kit’s specifications. Absorbance was read at 450 nm (Bio-Rad 680; Bio-Rad, Hercules, CA, USA). The quantity of SOD in the serum was estimated from a calibration curve.

Detection of Bcl-2 and BAX gene expression. Vascular thoracic aortas were quickly removed and placed into liquid nitrogen. Frozen samples were reserved at -80°C for further analysis. The thoracic aorta (50 mg) was homogenized using glass-Teflon[]. Total RNA was prepared using TRIzol® Reagent (Invitrogen, Thermo Fisher Scientific, Inc., Waltham, MA, USA) according to the manufacturer’s instructions. Complementary DNA (cDNA) was synthesized with a First Strand cDNA Synthesis kit (Thermo Fisher Scientific, Inc., Beijing, China). RT-PCR was carried out using a Maxima SYBR-Green PCR kit (Thermo Fisher Scientific, Inc.) with indicated primers. After an initial 10 min at 95°C, the PCR program was finished as follows: 95°C for 15 sec, 60°C for 30 sec, and extension at 72°C for 30 sec, for 40 cycles. At the end of the reaction, melting curve analysis was performed to ensure the specificity of the reaction. β-actin was used as an internal control. Primers used for the PCR are shown in Table I.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Sequence (5'-&gt;3')</th>
<th>bp</th>
<th>Tm/°C</th>
<th>GenBank</th>
</tr>
</thead>
<tbody>
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<td>Bcl2</td>
<td>Forward: AGCCTGAGAGCAACCGAAC Reverse: AGCGAGAGAGAAGTCAATCC</td>
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<td>60</td>
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<tr>
<td>Bax</td>
<td>Forward: TTGCTACAGGGTTTCCATCCAG Reverse: TGTGTGTGTCAGTTCATCG</td>
<td>145</td>
<td>60</td>
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<tr>
<td>β-actin</td>
<td>Forward: CACCGCGAGTACAACCTTC Reverse: CCCATACCCACCACCTACACC</td>
<td>207</td>
<td>60</td>
<td>NM_031144</td>
</tr>
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</table>

Statistical analysis. Data were analyzed by SPSS, Inc. 21.0 (Chicago, IL, USA), and the results were presented as mean ± SD. Statistical difference was evaluated using the t-test. P<0.05 was considered to indicate a statistically significant difference.

Results

Behavior observation in rats. Before heat exposure, all the rats had good appetite, activity and agile reaction. Their fur was...
tight, clean and smooth. The rats were restless in the evening. During thermal exposure, the rats’ limbs, nasal and testicular filled with blood, which was more obvious in the DHE group. Concerning mental state, the rats were tired and listless. Their surface fur was damp, the response to external stimuli was significantly reduced, and the rats hid their body in bedding material.

**Measurement of heart rate and blood pressure.** Compared with CN, the heart rate in the HE and DHE groups was retarded, albeit there was no significant difference (P>0.05) (Fig. 1). In the HE group, SBP, DBP and the mean arterial pressure (MAP) of rats were significantly higher than the CN (P<0.05). In the DHE group, the SBP of rats was significantly higher than that of the CN (P<0.05), albeit SBP, DBP and MAP
of rats were lower than the rats in the HE group, although there was no statistical significance (P>0.05) (Fig. 2).

Thoracic aorta reactivity. In the CN and HE groups, NA induced a dose-dependent contraction on the aortic ring. However, the contractive response was stronger in the HE compared with the CN group (P<0.05), but there was no significant difference between the HE and DHE groups (P>0.05) (Fig. 2).

Detection of the serum SOD and MDA. The serum SOD in the DHE group was lower than that of the CN group (P<0.05) (Fig. 4), the level of MDA was significantly higher than that of the CN group (P<0.05) (Fig. 5). Compared with the CN and DHE groups, the serum SOD and MDA in the HE group was fluctuated but no significant difference was observed (P>0.05).

Detection of Bcl-2 and Bax. Compared with CN, the expression of Bcl2 was significantly reduced (P<0.05) in the DEF group, while the Bax expression was increased (P<0.05). Compared with the CN and DHE groups, the expression of Bcl2 and Bax in the HE group had no significant difference (P>0.05) (Fig. 6).

Discussion

The present study has demonstrated that heat stress without food and water changed the rats' existence, increased blood pressure and damaged blood vessel function.

Living or working in a high temperature environment leads to not adapting to adverse reactions (6). Consequently, a lack of food and water reduces an individual's ability to survive. In a high temperature environment, the evaporation of the skin becomes the main way of cooling the body, but the high humidity limits the evaporation of perspiration. Under water shortage conditions, excessive loss of perspiration may damage the body's cardiovascular and thermoregulatory function (7-9). In response to the high temperature, the human body needs heat dissipation and expansion of body surface blood vessels, the amount of blood in the body surface increases, and the amount of blood supply for heart, cerebrovascular is relatively reduced, leading to cerebral ischemic and hypoxic reactions (10,11). The experiment results showed that, with the increase of heat exposure intensity, the rat heart rate had a tendency to slow down. Overton et al. found that hunger and high temperature were able to reduce the heart and metabolic rate (12). The study by Williams et al. showed hunger and high temperature were associated with the gradual reduction of activity (13). These indicated that, under the condition of hunger and high temperature, the rats could reduce heat production by decreasing movement and the metabolic rate. In the HE group, SBP, DBP, MAP were significantly higher than that of CN. The SBP of the rats in the DHE group was significantly higher than that of the CN. These results showed that high temperature increased blood pressure.

Vascular reactivity is a basic and direct index that reflects the state of the artery blood vessel function (14). Enhanced contractive function is the main performance of damaged blood vessels (15,16). The experiment results showed that the thoracic aorta reactivity in the HE and DHE groups was higher than that of the CN group. The environment of high temperature is harmful to blood vessels. However, it is difficult to find vascular injury; we investigated blood vessel function and attempted to identify the mechanism underlying vascular injury.

SOD is an important enzyme, which removes superoxide free radical from aerobic organisms and defense against the toxic effect of the oxygen-free radicals (17,18). MDA is the oxidative product of polyunsaturated fat in biological membrane (19). Its level can reflect the degree of lipid peroxidation and free radicals attacking the body's cells. In this experiment, SOD decreased significantly whereas the MDA level was increased in the the DHE group. This result suggested that high temperature without food and water caused increase of endogenous oxygen free radicals. Therefore, enhanced oxidative stress damaged the vascular elasticity.

Cell apoptosis was induced by stress, such as free radicals, hypoxia and blood deficiency (20,21). The present study focused on the function of Bcl2 and Bax in the regulation of apoptosis. In the process of cell apoptosis, members of the Bcl-2 family play a vital role. They have high homology, and share the conserved domain of BH1, BH2, BH3 and BH4. The Bcl2 family can be divided into two categories, one of which is anti-apoptotic, mainly containing Bcl2, Bcl-XL, Bcl-W, Mcl-1 and CED9. The other type promotes cell death, and mainly includes Bax, Bak, Bcl-XS, Bad, Bik and Bid. Increased Bax promotes cell apoptosis, whereas increased Bcl2 inhibits cell apoptosis (22). In this experiment, the expression of Bcl2 was significantly reduced in the DHE group, while Bax was increased. The present study has demonstrated that under the condition of dehydration heat exposure, the expression of Bax was markedly elevated, whereas Bcl2 was reduced. These results suggested that the cell apoptotic process was initiated by the heated environment, resulted in the altered organizational structure and affected the function of blood vessels.

The results of the present study show that a high temperature was harmful to the body, where particularly in the case of lack of food and water, the heat exposure elevated blood pressure, increased arterial reactivity, which was related to the elevated production of MDA, the impaired production of SOD, and the increase of cell apoptosis. These findings are useful in gaining a better understanding of the influence of dehydrated heat exposure on vascular function, and provide certain theoretical and experimental guidance for protection under high temperature.

Acknowledgements

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References


