The difference of organism redox state in AIDS combined with tuberculosis and tuberculosis only patients.

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Abstract

Background: To compare the differences of organism redox state in AIDS patients complicated with tuberculosis and tuberculosis patients.

Methods: Subjects were divided into two groups, AIDS and TB group (n=40) and TB group (n=40) with 30 healthy subjects as the control group. Based on the measurement of reduced Glutathione (GSH) and oxidized glutathione (GSSG) as well as NADPH and NADP, the ratios of GSH/GSSG and NADPH/NADP were used to describe the redox state of patients and compare the organism redox state in two groups.

Results: The levels of GSH, GSSG, GSH/GSSG, NADPH, NADP⁺, NADPH/NADP⁺ in AIDS and TB group, TB group and control group were different (F=115.0, P<0.01, F=3944.0, P<0.01, F=1957.0, P<0.01, F=318.0, P<0.01, F=318.0, P<0.01, F=12.0, P<0.01, F=104.0, P<0.01).

Conclusion: There is a difference in the redox state of AIDS patients complicated with tuberculosis and tuberculosis patients.

Keywords: AIDS, Tuberculosis, AIDS combined with tuberculosis, Organism redox state.

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Introduction

Tuberculosis is still an infectious disease threatening the health of the people. In 2015, 10.4 million new tuberculosis patients were reported, of which 1.2 million were tuberculosis patients with HIV infection, accounting for 11% of all TB patients [1]. Tuberculosis was the most common opportunistic infections in AIDS patients with the mortality rate of 30% [2]. Therefore, the new approach to the treatment of AIDS combined with tuberculosis has become a new hot spot in clinical research. Studies have shown the presence of oxidative stress in patients with tuberculosis [3]. Mycobacterium tuberculosis (Mtb) suffered from exogenous oxidative stress [4]. In the course of infection, mycobacteria were affected by lots of redox stresses, such as Reactive Oxygen Species (ROS), Reactive Nitrogen Species (RNS) and hypoxia [5]. Toxicological effect induced by ROS was further compounded in Mtb result from the absence of important DNA repair pathways [6,7]. In addition, ROS and RNS produced by host was the key to control Mtb [8,9]. Moreover, our preliminary study also showed that the redox state of AIDS patients was biased toward the oxidation direction. Some researchers suggested that oxidative stress-induced free radicals could lead to cellular damage in HIV [10,11]. And Pocernich et al. [12] showed that AIDS dementia was related to increased oxidative stress. Furthermore, TB is the main infectious cause of death in HIV co-infected patients [13]. However, there were little relevant reports on the difference between the redox state of AIDS patients with tuberculosis and tuberculosis patients. In this study, 40 cases of AIDS patient with tuberculosis and 40 cases of tuberculosis patients were tested on redox state to compare the difference.

Materials and Methods

Instruments

Refrigerated centrifuge (20PR-52D, Japan HI-TACHI); ultra-low temperature storage box (MDF-382E, Japan Sanyo); centrifuge (TDL-5-A, Shanghai Anting); high-speed refrigerated centrifuge (1-15K, United States Sigma).

Reagents

Reduced glutathione (GSH) and oxidized glutathione (GSSG) kit (Wuhan Biofavor, product code S0053); Nicotinamide adenine dinucleotide phosphate (NADP⁺) and NADPH kit (Wuhan Biofavor). In accordance with the requirements of the kit instructions, related reagents were prepared.

Object and grouping

The subjects of AIDS and TB group were selected from patients treated in Beijing Medical Center from June 2010 to October 2016 (A total of 40 patients, aged from 18 to 59 y old, participated in the group, which had 30 males and 10 females with the average age of 42.5 y old). AIDS diagnostic criteria referred to Guidelines for diagnosis and treatment of HIV/AIDS in China [14]. Anti-HIV antibody test was conducted by
Wuhan City Center for Disease Control and Prevention. Tuberculosis diagnosis referred to tuberculosis-diagnosis, management, prevention, and control guidance [15]. The subjects of TB group (TB group were selected from patients treated in Beijing Medical Center from June 2010 to October 2016 (Methods 40 subjects, 30 males and 10 females, aged from 18 to 60 y old with the average age of 43.4, were included in the this group). Thirty healthy subjects were selected as the control group.

**Research methods**

The fasting venous blood samples (3 ml) were collected and stored in a cryopreservation chamber after low-temperature centrifugal in heparin anticoagulant tube at -70°C for further NADPH/NADP⁰ and GSH/GSSG measurement. All samples were re-dissolved at room temperature before the determination. According to the requirements of kit, GSH and GSSG levels were measured by colorimetry. NADPH and NADP⁰ were measured by enzyme-linked immunosorbent assay.

**Statistical processing**

SPSS 11.5 statistical analysis software package was used for statistical analysis of data, and measurement data was recorded as \( \bar{x} \pm s. \) T test was used to compare between groups. A statistical significance was defined when \( p<0.05. \)

**Results**

**Redox state of plasma glutathione**

The levels of GSH in AIDS and TB group, TB group and Control group were different (\( F=115.0, P<0.01. \) Compared with control group, the GSH levels of AIDS and TB group and TB group were significantly lower (\( P<0.01. \) The levels of GSSG in AIDS & TB group, TB group and control group were different (\( F=3944.0, P<0.01. \) Compared with control group, the GSSG levels in AIDS and TB group and TB group were significantly lower (\( P<0.01. \) Compared with TB group, the GSSG level of AIDS and TB group was significantly increased (\( P<0.01. \) The GSH/GSSG values of AIDS and TB group, TB group and Control group were different (\( F=1957.0, P<0.01. \) Compared with control group, the GSH/GSSG values of AIDS and TB group and TB group were significantly decreased (\( P<0.01. \) Compared with TB group, the GSH/GSSG value of AIDS and TB group was significantly decreased (\( P<0.01, \) Figure 1).

**Redox state of plasma coenzyme II**

The levels of NADPH in AIDS and TB group, TB group and Control group were different (\( F=318.0, P<0.01. \) Compared with control group, the levels of NADPH in AIDS and TB group and TB group were significantly decreased (\( P<0.01. \) The levels of NADP⁰ in AIDS and TB group, TB group and control group were different (\( F=12.0, P<0.01. \) Compared with the control group, the levels of NADP⁰ in AIDS and TB group and TB group were significantly decreased (\( P<0.01. \) Compared with TB group, the level of NADP⁰ in AIDS and TB group was significantly decreased (\( P<0.01. \) The NADPH/NADP⁰ ratio of AIDS and TB group, TB group and control group was different (\( F=104.0, P<0.01. \) Compared with the control group, the ratios of NADPH/NADP⁰ in AIDS and TB group and TB group were significantly decreased (\( P<0.01. \) Compared with TB group, the ratio of NADPH/NADP⁰ in AIDS and TB group was significantly decreased (\( P<0.01, \) Figure 2).

**Discussion**

HIV infection is the most important risk factor for tuberculosis. Tuberculosis is the most common opportunistic infection of HIV/AIDS, which can lead to a significant increase in the probability and mortality of other opportunistic infections in AIDS patients. Studies have found that the redox states of HIV/AIDS patients were biased toward the oxidation direction [16,17], and its degree of bias was positively correlated with disease progression. GSH is the main defensive mechanisms against toxic agents and oxidant-mediated injury [18,19]. And the GSH/GSSG ratio is a useful measure of cellular redox status [20]. In addition, NADPH/NADP⁰ oxidation and reduction played an important role in maintaining GSH/GSSG equilibrium [21]. With Intracellular GSH depletion, the incidence of tuberculosis in AIDS patients was much higher than healthy individuals. The study showed that the GSH levels of peripheral monocytes and erythrocytes in tuberculosis patients were significantly decreased [22]. Therefore, no matter tuberculosis or AIDS patients, the occurrence and development of disease were related to the body redox situation. This study showed that, based on GSH or NAD-PH concentration, redox state were biased toward the oxidation direction. And patients with AIDS combined tuberculosis showed more obvious bias than tuberculosis patients, resulting from overlay effect of AIDS and tuberculosis. The change of redox state was one of the important mechanisms of AIDS and had a certain relevance.
to AIDS development. This study provided a basis for the treatment of AIDS patients with tuberculosis from the perspective of changing the body's redox state.

References


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