Original Article

Pathomorphological changes of bone marrow adipocytes in process of steroid-associated osteonecrosis

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Abstract: Objective: The role of extravascular fat deposition in pathogenesis of steroid-associated osteonecrosis (ON) still remains unclear. This study aimed to explore the pathomorphological changes of bone marrow adipocytes over time in a rabbit ON model. Methods: Thirty-two adult rabbits were divided into control group (n=16) and steroid group (n=16). Rabbits in the steroid group were injected with venous lipopolysaccharide once and intramuscular methylprednisolone trice to induce ON. Rabbits in the control group were treated with normal saline of equal volume. 2 weeks (early stage; n=8) and 4 weeks (late stage; n=8) after the last steroid injection, animals were sacrificed, and bilateral femora were harvested. The density, diameter and area of bone marrow adipocytes were determined by histomorphometry, and ON was evaluated histopathologically. Results: The adipocyte density in steroid group increased by 67.1% and 54.4% at week 2 and week 4, respectively, when compared with control group, but there was no significant difference between week 2 and week 4. The adipocyte diameter in the steroid group at week 4 was significantly larger than that in the control group, but the adipocyte diameter in the steroid group at week 2 was slightly smaller than that in the control group. The adipocyte area in the steroid group increased by 44% and 83.4% at week 2 and week 4, respectively, when compared with control group, and the adipocyte area in the steroid group at week 4 was markedly larger than that at week 2. In the control group, there were a largest number of adipocytes with 40-50 μm in diameter. When compared with the control group, most of increased adipocytes in the steroid group at week 2 were 30-40 μm in diameter, and those at week 4 were 50-60 μm in diameter. In the steroid group, histopathological examination showed ON was found in 25% (2/8) of rabbits at week 2 and 87.5% (7/8) of rabbits at week 4. Conclusion: In the process of ON, extravascular fat deposition is characterized by increased small adipocytes at the early stage and hypertrophy of adipocytes at the late stage.

Keywords: Steroid-associated osteonecrosis, adipocyte, adipogenesis, fat transport

Introduction

Glucocorticoids have been widely applied in clinical treatment of inflammation and autoimmune diseases, but glucocorticoid treatment brings a severe complication: osteonecrosis (ON). Recently, many study showed high incidence of ON and high collapse rate in patients recovering from SARS after receiving glucocorticoid treatment, which makes steroid associated ON becoming a hot topic [1, 2]. There are several theories for steroid associated ON, such as theory of vascular compression due to extravascular fat deposition, theory of hypercoagulability and theory of fat embolism [2-5]. Among these theories, vascular compression due to extravascular fat deposition becomes a widely accepted theory. In patients with ON, histopathological examination has confirmed fat deposition in bone marrow. However, how the extravascular fat deposition involves in the pathogenesis of steroid associated ON is still poorly understood [6, 7], and the changes of bone marrow adipocytes in steroid associated ON are also unclear [7, 8]. In the present study, steroid associated ON was introduced to rab-
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bits according to previously described (this has a high incidence of ON) [7], and histomorphometry and classic methods were employed to evaluate ON. This study aimed to investigate the dynamic change in bone marrow adipocytes in steroid associated ON. Our results may provide evidence on the relationship between extravascular fat deposition and steroid associated ON.

Materials and methods

Animal model and grouping

A total of 32 rabbits aged 28-32 weeks were randomly assigned into control group (n=16) and steroid group (n=16). The steroid associated ON was introduced with methods developed in our department [7]. In brief, rabbits in the steroid group were intravenously injected with endotoxin (LPS; 10 μg/kg, Escherichia coli 0111:B4, Sigma-Aldrich) once and intramuscularly with methylprednisolone (20 mg/kg, Pharmacia and Upjohn, USA) thrice with an interval of 24 h. Animals in the control group were treated with normal saline of equal volume. Rabbits were housed in a standard environment and given ad libitum access to food and water without activity restriction. 2 weeks (early stage) and 4 weeks (late stage) after last steroid injection, animals were sacrificed (n=8 at each time point), and bilateral femora were collected for examination.

Sample collection and treatment

After anesthesia, animals were sacrificed using an overdose of sodium pentobarbital, and bilateral femora were collected and fixed in 10% formalin for 72 h and then in 75% ethanol for 1
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week. Decalcification was done in 9% formic acid for 4 weeks followed by dehydration in a series of ethanol solution and subsequent embedding in paraffin. The proximal 1/3 of femora was cut into 4-μm sections followed by mounting with resin and H&E staining.

Histomorphometry of adipocytes in bone marrow

Four fields were randomly selected from each section at 100 magnification and two sections were collected from each femur. A total of 16 fields were selected from 4 sections of each rabbit followed by quantification of adipocytes in bone marrow. The density, diameter and area of adipocytes in bone marrow were determined and analyzed with Image-Pro Plus 5.1. Adipocyte density refers to number of adipocytes in unit bone marrow area (excluding bone trabecula); adipocyte diameter refers to mean diameter of adipocytes in bone marrow; adipocyte area refers to area of bone marrow adipocytes in unit bone marrow area (excluding bone trabecula) [9].

Histopathological examination of ON

ON was defined as formation of empty lacunae or karyopyknosis of a large number of necrotic osteocytes in bone trabecula, accompanied necrosis of surrounding fat tissues [9]. Once one femur developed ON, positivity for ON was confirmed.

Statistical analysis

Statistical analysis was done with SPSS version 13.0 for Windows (SPSS Inc., Chicago, IL, USA). Density, diameter and area of adipocytes were compared with analysis of variance between two groups. Data were expressed as mean ± standard deviation (±SD). A value of P<0.05 was considered statistically significant.

Results

Density of adipocytes in bone marrow

In the control group, the density of adipocytes in bone marrow was 149±15/mm². In the steroid group, the density of adipocytes in bone marrow was 249±25/mm² at week 2 and 230±23/mm² at week 4, which were increased by 67.1% and 54.4%, respectively, when compared with the control group (P<0.01). However, the density of adipocytes in bone marrow was comparable between week 2 and week 4 (P>0.05) (Figure 1).

Diameter of adipocytes in bone marrow

The mean diameter of adipocytes in bone marrow was 42.6±3.6 μm in control group, 38.2±3.1 μm in steroid group at week 2 and 52.9±4.1 μm in steroid group at week 4. Significant difference in the mean diameter of adipocytes was noted between the control group and the steroid group at different time points (Figure 2).

Area of adipocytes in bone marrow

The area of adipocytes in bone marrow was 23.2±3.9% in the control group. In the steroid group, the area of adipocytes in bone marrow was 33.4±4.7% at week 2 and 42.6±5.3% at week 4, which were increased by 44% and 83.6%, respectively, when compared with the control group (P<0.01) (Figure 3).

Proportion of adipocytes with different sizes in bone marrow

In the control group, the majority of adipocytes in bone marrow had a diameter of 40-50 μm. In the steroid group, the majority of adipocytes in bone marrow were 30-40 μm at week 2 and 50-60 μm at week 4 (Figures 4 and 5).

Incidence of ON

At the early stage (week 2), ON was found in 2 out of 8 rabbits (25%), and the incidence of ON was 87.5% (7/8) at the late stage (week 4). In the control group, ON was not observed.

Discussion

This study for the first time explored the relationship between changes of adipocytes in bone marrow and steroid associated ON with histopathological examination. Our results showed extravascular fat deposition in bone marrow (fat area in bone marrow) deteriorated over time, and the incidence of ON increased over time in animals with steroid treatment. Our findings provide evidence on that extravascular fat deposition in bone marrow may involve in the pathogenesis of steroid associated ON.
In the present study, the area, density and diameter of adipocytes in bone marrow at week 2 (early stage) were significantly larger than those in control group. This suggests that extravascular fat deposition at the early stage of ON is attributed to the increase in the number of adipocytes. In addition, at late stage (week 4), the area of adipocytes in bone marrow was markedly larger than that in the control group and the steroid group at week 2. With the increased incidence of ON at week 4, the mean diameter of adipocytes in bone marrow in steroid group was also significantly larger than that in the control group. Of note, the number of large adipocytes in the steroid group at week 4 was larger than that in the control group. Although the density of adipocytes at week 4 was higher than that in the control group, but comparable to that at week 2 in the steroid group. This indicates that the extravascular fat deposition at the late stage is attributed to the enlargement (hypertrophy) of adipocytes and at the early stage is attributed to the increase in the number of adipocyte.

In vivo study showed glucocorticoid could promote the differentiation of bone marrow derived mesenchymal stem cells (BMSCs) into adipocytes [10]. Recently, our group also found that BMSCs from rabbits with steroid associated ON had increased potential to differentiate into adipocytes following differentiation induction [11]. This implies that the small adipocytes which increase at early stage after steroid treat-
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Adipocytes are attributed to the differentiation of BMSCs in the presence of glucocorticoids. The increased diameter of adipocytes in live animals with ON can be explained by disturbance of fat transport [12, 13]. This indicates that the increase in large adipocytes at late stage of steroid treatment may be, at least partially, associated with compromised fat transport in adipocytes. Thus, our findings provide an interventional strategy for ON: to block the signaling pathway involving in glucocorticoid induced adipogenesis at early stage and to promote fat transport in adipocytes may be feasible in prevention of ON.

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