The timing and extent of intraosseous hypoxia in the oxidative stress-induced rat osteonecrosis model*

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ABSTRACT

Using a rat oxidative stress-induced femoral head osteonecrosis model, we determined the presence/absence and timing of the generation of hypoxia in the femoral head. DL-Buthionine-(S,R)-sulfoximine (BSO) 500 mg/kg was administered intraperitoneally to male Wistar rats. The rats were killed at 1, 3, 6, 12 hours, and 1, 3, 5 days after BSO administration, and the bilateral femora were removed. A group not administered BSO (control group) was also studied (each group n = 5). In the femoral heads of each group, the expression of hypoxia-inducible factor-1 alpha (HIF-1α) as an index of hypoxia was confirmed by the Western blot method, and quantified using analytical software. In the femoral head increased HIF-1α expression was found in all groups from 1 hour after BSO administration (p < 0.05). In particular, in all specimens of the group 3 hours after BSO administration the most intense expression of HIF-1α amounting to about 13-fold of that of control group was noted (p < 0.001). The present results suggested that in the extremely short period of 3 hours after BSO administration hypoxia severe enough to cause osteonecrosis was induced by oxidative stress in the rat femoral head.

Keywords: Osteonecrosis; Oxidative Stress; Buthionine Sulfoximine; Hypoxia-Inducible Factor-1α

1. INTRODUCTION

A broad consensus has been reached implicating intraos-
Medical University. All rats were housed under standard laboratory conditions (temperature 24°C, 12-hour light/dark cycle) and were given food and water ad libitum. They were given a single intraperitoneal injection of BSO (500 mg/kg) and were killed 1 hour (B1h group), 3 hours (B3h group), 6 hours (B6h group), 12 hours (B12h group), 1 day (B1d group), 3 days (B3d group) or 5 days (B5d group) after administration. Untreated rats were compared as a control group (group N). Each group contained 5 rats. This study was conducted in accordance with the guidelines of the Animal Research Committee of Kanazawa Medical University.

2.2. Tissue Preparation

All rats were killed using intraperitoneally injected sodium pentobarbital. Promptly after sacrifice the bilateral femoral heads were extracted and cryopreserved at −80°C after the bilateral femora were removed.

2.3. Western Blot Assay

Immunoblotting for HIF-1α was performed on the femoral head of BSO administered rats, as well as non-treated rats. Proteins were extracted using lysis buffer (50 mM Tris-HCl, pH 7.6, 10% glycerol, 5 mM magnesium acetate, 0.2 mM ethylenediamine tetraacetic acid, 1 mM phenylmethylsulfonyl fluoride, and 1% sodium dodecyl-sulfate). Extracted protein (20 mg) was applied to and electrophoresed on a 10% polyacrylamide gel, and transferred to a nitrocellulose membrane (Atoh, Tokyo, Japan). The membranes were reacted overnight at 4°C with anti HIF-1α polyclonal antibody (Millipore) at a concentration of 1:200 dilution. After incubation with peroxidase-labeled goat anti-rabbit IgG antibody (Dako Cytomation) for 1 hour at room temperature and vigorous washing, the nitrocellulose membrane was incubated with ECL-pulse (GE Medical, USA) and photographed digitally using LAS4000 (FUJIFILM, Tokyo, Japan). All samples were standardized by immunoblot using anti-β-actin mouse monoclonal antibody (Sigma Chemical Co., St. Louis, MO).

2.4. Quantification of the Western Blot Results

Protein quantification was done using analytical software “Multi Gauge” (FUJIFILM, Tokyo, Japan). β-actin was used as the internal control for all specimens in each group. All values were expressed as the mean ± SD, and Arbitrary Units (AU) were used at the time of measurement.

2.5. Statistical Analysis

Statistical analysis was performed using analysis of variance with Dunnett’s multiple comparison test. By using this test we were able to determine the point in time after administration of BSO that HIF-1α level showed a significant change, as compared with the values in N group. p values less than 0.05 were considered significant. The statistical analysis was performed using StatView J-5.0 software (SAS Institute).

3. RESULTS

3.1. Western Blot for HIF-1α

An approximate molecular weight of around 120 kDa for HIF-1α was demonstrated by Western blot analysis. HIF-1α was overexpressed in BSO administered specimens. HIF-1α expression started to increase from B1h Group, with a marked increase found in all specimens of B3h Group. From B6h-B12h Group onwards increased HIF-1α expression was found as compared to N Group (Figure 1).

3.2. Comparison by Quantification of HIF-1α Expression by Western Blot

When quantified using analytical software, the mean value was (9.9 ± 1.0) × 10^4 AU in N Group, and (128.6 ± 12.2) × 10^4 AU in B3h group, representing a significantly increased expression of about 13-fold in the latter as compared to the former (p < 0.001). The corresponding values were (97.0 ± 37.0) × 10^4 AU in B6h group, (94.8 ± 43.2) × 10^4 AU in B12h group, (74.6 ± 31.5) × 10^4 AU in B1d group, (58.0 ± 31.2) × 10^4 AU in B3d group, and (80.9 ± 38.8) × 10^4 AU in B5d group. In all of the groups administered BSO HIF expression was significantly increased as compared to N group (p < 0.05) (Figure 2).

4. DISCUSSION

Various theories have been proposed to explain the pathogenesis of steroid-induced osteonecrosis [12-14]. As noted above, while there is general agreement that in the end osteonecrosis is induced by intraosseous ischemia, various issues such as the timing of occurrence and amount of the intraosseous hypoxia remain unclear. To elucidate the developmental mechanisms of osteonecrosis
and devise optimal prophylactic strategies it is important to grasp the changes occurring in bone from immediately after drug administration. For this reason, detailed determination of the actual severity of the ischemic event occurring after drug administration and pinpointing the stage at which it occurs, as well as confirmation of the previously described finding that only a single ischemic event is induced are required.

In the present study, our attention was drawn to the transcription factor HIF-1α as an index of tissue hypoxia. HIF-1α is particularly well suited to assess the timing and amount of hypoxia because while it is rapidly degraded in the presence of normal oxygen concentrations, in the presence of hypoxia it is not degraded and its concentration rises [10,11,15-17].

In the rat, osteocyte apoptosis has been reported to begin 12 h from the ischemic insult, with osteonecrosis observed 4 - 5 days from the onset of ischemia [8,9]. Accordingly, in this investigation we considered it important to focus on the state of the hypoxia occurring in the bone up to 5 days after BSO administration at which point the findings of osteonecrosis can be confirmed histopathologically in this model [7].

This experiment confirmed that in the model used hypoxia in the femoral head was induced by oxidative stress within the extremely short period of <3 h. Also, especially in B3h group as compared to N group about a 13-fold increase in HIF-1α expression was found. The expression of HIF-1α has been reported to increase by about ≥10-fold in hypoxia of about 2.5% - 5% of the normal oxygen concentration [15-17]. Accordingly, we concluded that in this model after BSO administration the oxygen concentration of femoral head tissues in all specimens fell to less than about 2.5% - 5% of normal within 3 h.

The results of this study are extremely important in considering the developmental mechanism of femoral head osteonecrosis. This is because if a hypoxic event is indeed the ultimate cause of osteonecrosis, all specimens in this model would be expected to develop osteonecrosis. However, since the rate of development of femoral head osteonecrosis was only 40% in this model, the possible involvement of some kind of secondary factor in the development of osteonecrosis occurring after the hypoxic event 3 h after BSO administration is also suggested. Also, from B6h group onwards as compared to N group HIF-1α expression was significantly enhanced. Accordingly, it could be concluded that the intraosseous hypoxic state induced by oxidative stress begins in bone from 1 h, with an extremely strong hypoxic state developing within 3 h, and the hypoxic state persisting from B6h group onwards albeit while showing some gradual improvement.

It has previously been reported that oxidative stress develops within 24 - 72 hours after steroid administration [1-3]. The present results showing the occurrence of severe hypoxia from 3 h after oxidative stress help to identify the mechanism underlying the development of osteonecrosis very soon after steroid administration. Accordingly, the present experimental results can be considered extremely important also from the viewpoint of devising novel preventative approaches for steroid-induced osteonecrosis.

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REFERENCES


