Osteonecrosis in patients with systemic lupus erythematosus develops very early after starting high dose corticosteroid treatment

K Oinuma, Y Harada, Y Nawata, K Takabayashi, I Abe, K Kamikawa, H Moriya

Abstract

Objectives—To investigate the actual time of onset of osteonecrosis (ON) after high dose corticosteroid treatment in systemic lupus erythematosus (SLE).

Methods—72 patients with active SLE, who received high dose corticosteroid for the first time, for the development of ON at hips and knees were monitored by magnetic resonance imaging for at least 12 months.

Results—ON lesions were detected in 32/72 patients (44%) between one and five months (3.1 months on average) after starting high dose corticosteroid treatment. No osteonecrotic lesion was newly detected from the sixth month of treatment until the end of the follow up period.

Conclusion—The findings suggested that the actual time of onset of ON in SLE is within the first month of high dose corticosteroid treatment.

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It is well known that osteonecrosis (ON) frequently develops in patients with systemic lupus erythematosus (SLE) with a prevalence ranging from 10% to 52%1–6 and that the development of ON in SLE is closely related to the high doses of corticosteroids received. Previous clinical studies have provided conflicting results about the possible association of corticosteroid treatment with the development of ON. The total treatment period,1 the highest daily dose,2 a continuous high dose,4 5 and the cumulative dose of corticosteroid6 have been reported as risk factors of ON in SLE. These conflicting results seem to be mainly due to the difficulties in detecting early ON, which occurs silently and cannot be based on conventional radiographic images. Magnetic resonance imaging (MRI) has proved to be more sensitive than conventional methods, such as radiography, computed tomography, and skeletal scintigraphy, for the detection of very early, asymptomatic ON.7 8

We investigated the actual time of onset of ON in SLE after starting high dose corticosteroid treatment using serial MR images of both hips and both knees obtained every three months, during one year.

Patients and methods

Patients

We evaluated 72 new patients with active SLE who were treated with a high dose of corticosteroid (40 mg/day or more equivalent prednisolone) at Chiba University Hospital. Thirty-five of the 72 patients were also given methylprednisolone pulse treatment (MPPT). All the patients fulfilled the revised criteria for SLE established by the American Rheumatism Association.9 The study group comprised 68 women and four men with a mean age of 34.8 years (range 13–66). The disease activity of SLE was evaluated using the SLE Disease Activity Index (SLEDAI).10

We excluded patients with a history of corticosteroid use and alcoholism because it is well known that osteonecrotic lesions already exist in these patients.

MRI study

MRI of both hips and both knees was carried out at 1, 3, 6, and 12 months after starting corticosteroid treatment in all patients, irrespective of joint symptoms, or as close to these intervals as possible. At the end of the study follow up period, patients were divided into the ON group and the non-ON group based on the presence/absence of abnormal MRI findings. Abnormal MRI findings indicating osteonecrotic lesions were well demarcated, band-like zones of decreased signal intensity on T1 weighted, spin echo (SE) images and band-like zones of increased signal intensity on short T1 inversion recovery (STIR) images.7 8 MRI was conducted using a 0.5 Tesla superconductive unit (MRF-50; Toshiba, Japan). T1 weighted, SE images were obtained with repetition times (TR) of 300 to 400 ms and echo times (TE) of 18 to 40 ms. STIR images were obtained with a TR of 1500 to 3000 ms, inversion times (TI) of 100 to 150 ms, and a TE of 30 to 42 ms.

Statistical analysis

To compare the clinical variables between the ON and non-ON groups, the Mann-Whitney U test or χ² test was performed. Differences at p<0.05 were regarded as significant.

Results

Characteristics of patients with or without osteonecrotic lesions

Table 1 shows the clinical profiles of patients in the ON group and non-ON group. The ON group comprised 32 patients (29 women, three men) with a mean age of 34.1 (range 16–66). Abnormal MRI findings were found in the hip, knee, or both. The non-ON group comprised 40 patients (39 women, one man) with a mean age of 35.1 (range 13–63). The initial mean (SD) corticosteroid dose was 58.2 (10.1) mg/day in the ON group and 58.6 (16.6) mg/day in the non-ON group. Seventeen patients in the ON group and 18 in the
Table 1  Baseline characteristics of the groups with and without osteonecrosis (ON)

<table>
<thead>
<tr>
<th></th>
<th>ON group</th>
<th>Non-ON group</th>
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</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>34.1</td>
<td>35.1</td>
</tr>
<tr>
<td>Mean</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>16–66</td>
<td>13–63</td>
</tr>
<tr>
<td>Sex</td>
<td>Male</td>
<td>Female</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>39</td>
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<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Initial corticosteroid dose†</td>
<td>58.2 (10.1)</td>
<td>58.6 (16.6)</td>
</tr>
<tr>
<td>(mg/day equivalent prednisolone)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>SLEDAI*†</td>
<td>27.2 (12.3)</td>
<td>25.1 (9.8)</td>
</tr>
<tr>
<td>MPPT*</td>
<td>18</td>
<td>17</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td></td>
</tr>
<tr>
<td></td>
<td>34.1</td>
<td>35.1</td>
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</table>

†Mean (SD).

* SLEDAI = SLE Disease Activity Index; MPPT = methylprednisolone pulse treatment.

Discussion

We believe this is the first study to investigate the actual time of onset of ON after high dose corticosteroid treatment in SLE using MRI. In this study, ON lesions, which were seen as well demarcated band-like zones, were detected by MRI at an average of 3.1 months after starting corticosteroid treatment. In a recent MRI study of patients who had received a renal transplant, ON lesions were detected at 1–4 months after starting high dose corticosteroid treatment.11 Although this study showed the presence of ON at the time of MRI, the actual time of onset of ON was not clarified.

In this paper we showed that ON lesions appeared on MRI early after starting corticosteroid treatment—namely, between 39.6 and 100.2 days of treatment. Our observations indicated that early development of ON in SLE...
was related to an event just after high dose corticosteroid treatment, and was not related to the total treatment period, the highest daily dose, a continuous high dose, or the cumulative dose of corticosteroid. Although corticosteroid is a well known cause of ON in SLE, clinical studies attempting to demonstrate an association of corticosteroid treatment with the development of ON in SLE have yielded conflicting results.1–6 This conflict seems to be mainly caused by the employment of conventional methods, such as x ray examination and bone scintigraphy, in many of the previous studies.78 Many patients with early ON, which might have been detected by MRI, were probably missed by these conventional methods. MRI has proved to be the most useful method to detect early stage ON lesions that remain occult, clinically, radiologically, and scintigraphically.78

Our study suggested that the actual onset of ON in SLE may be within the first month after starting high dose corticosteroid treatment because the first MR images in 14 patients were obtained significantly later than those of the 18 remaining patients in the ON group (92.6 (22.0) days v 39.6 (19.9) days, p<0.0001).

We speculate that high dose corticosteroid treatment may enhance the development of ON as a result of microvascular ischaemia in patients with SLE who have vasculitis. It has been reported that corticosteroids decrease bone blood flow, thereby enhancing ischaemia owing to increased bone marrow pressure produced by intramedullary lipocytes hypertrophy.14 On the other hand, it was also reported that patients with SLE present haemostatic abnormalities, resulting from vascular endothelial damage,15 therefore, measurement of haemostatic variables in patients with SLE in the early period after starting high dose corticosteroid treatment may predict the subsequent development of ON. In addition, the presence of antiphospholipid antibody (aPL) has been proposed to predispose towards ON, but its role remains controversial.2 4–6 10–16 Recently, some large scale studies failed to show any association between aPL and ON.16–18 We also measured aPL for 45 patients in this study and found no positive relation between ON and the presense of aPL positivity.

In this prospective MRI study, hip and knee ON lesions occurred in 44% of patients with SLE, a percentage which was within the range reported by most other studies (10–52%).1,6 All our patients with ON lesions were asymptomatic during the follow up period of 12 months, but in other reports some patients...
were symptomatic and needed surgical treatment. Segmental collapse is reported to be mainly a matter of time and depends on the size of the necrotic lesion.19 Although new osteonecrotic lesions appear after the follow up period only rarely, further follow up is recommended.

In conclusion, we found that in patients with SLE ON lesions developed very early (1–2 months) after starting high dose corticosteroid treatment. We think that MRI is useful not only in detecting early development of osteonecrotic lesions but also in examining the risk factors for their development.

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