Circulating exosome levels in the diagnosis of steroid-induced osteonecrosis of the femoral head

H-Y. Zhu, Y-C. Gao, Y. Wang, C-Q. Zhang
Shanghai Jiao Tong University Affiliated Sixth People’s Hospital, Shanghai, China

Objective
Circulating exosomes represent novel biomarkers for multiple diseases. In this study, we investigated whether circulating exosome levels could be used as a diagnostic biomarker for steroid-induced osteonecrosis of the femoral head (ONFH).

Methods
We assessed the serum exosome level of 85 patients with steroid-induced ONFH and 115 healthy donors by Nanosight detection. We then assessed the diagnostic accuracy of serum exosomes by receiver operating characteristic curve analysis.

Results
The circulating exosome level of the ONFH group was significantly lower than that of control group. The area under the curve was 0.72, suggesting that the level of serum exosomes has moderate diagnostic accuracy for steroid-induced ONFH.

Conclusion
Circulating exosome levels are valuable in the diagnosis of steroid-induced ONFH.

Keywords: Exosome; Osteonecrosis; Steroid

Article focus
To develop an economical and convenient method for the diagnosis of steroid-induced osteonecrosis of the femoral head (ONFH).

Key messages
- The circulating exosome levels of the ONFH group were significantly lower than that of the control group.
- Level of serum exosomes has moderate diagnostic accuracy for steroid-induced ONFH.

Strengths and limitations
- Strength: the current study is the first study to reveal the value of serum exosome levels as a diagnostic marker for steroid-induced ONFH.
- Limitation: Further studies should be conducted with larger sample sizes in different ethnic groups to validate and extend our findings.

Introduction
Steroid-induced osteonecrosis of the femoral head (ONFH) involves the necrosis of cancellous bone which ultimately leads to the collapse of the femoral head.1,2 The incidence of osteonecrosis ranges from 4.6% to 37% in patients receiving steroid treatment.1,3-6 Intervention at an early stage is the key to successful treatment of steroid-induced ONFH. However, early-stage ONFH could only be definitively diagnosed with MRI scanning which is expensive and inconvenient. It is important to find an economical and convenient method for screening ONFH development in patients receiving steroid therapy. After initial screening, MRI scanning can then be conducted for further confirmation.

Exosomes are 30 nm to 120 nm, cell-secreted vesicles that are present in all biological fluids, including blood, urine and cultured medium of cell cultures.7,8 Exosomes carrying miRNA, mRNA and proteins are important vehicles for intercellular communication
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Bone & Joint Research

Table I. Demographic features of patients with osteonecrosis of the femoral head and healthy donors

<table>
<thead>
<tr>
<th>Demographic characteristics</th>
<th>ONFH group (n = 85)</th>
<th>Control group (n = 115)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>44</td>
<td>60</td>
</tr>
<tr>
<td>Female</td>
<td>41</td>
<td>55</td>
</tr>
<tr>
<td>Age (yrs)</td>
<td>56.3 ± 7.9</td>
<td>55.9 ± 8.3</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>167.2 ± 11.4</td>
<td>166.8 ± 11.7</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>67.9 ± 8.9</td>
<td>67.3 ± 9.2</td>
</tr>
</tbody>
</table>

Statistical analysis: Student’s t test or chi-squared test
SD, standard deviation

Table II. Area under the curve (AUC) for serum exosome level in the diagnosis of steroid-induced osteonecrosis of the femoral head (ONFH)

<table>
<thead>
<tr>
<th>AUC</th>
<th>Cut-off</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PPV (%)</th>
<th>NPV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.72</td>
<td>8.4 × 10^13/ml</td>
<td>65.2</td>
<td>77.6</td>
<td>63.5</td>
<td>79.8</td>
</tr>
</tbody>
</table>

Receiver operating characteristic (ROC) curve analysis was used to calculate the AUC of serum exosome level for diagnosing steroid-induced ONFH.

Recent progress indicated that exosomes in biological fluids are promising disease-related biomarkers for chronic kidney disease, pancreatic cancer, and acute coronary syndrome. In this study, we report for the first time that the circulating exosome level in steroid-induced ONFH patients was significantly lower than the level found in a healthy population.

Materials and Methods

Patient information. The study was approved by the Ethics Committee of Shanghai Jiao Tong University Affiliated Sixth People’s Hospital. Written consent was obtained from all donors in accordance with the Declaration of Helsinki. The serum samples were obtained from the ONFH Bank of Shanghai Jiao Tong University Affiliated Sixth People’s Hospital. This study included 85 serum samples from patients with steroid-induced ONFH and 115 serum samples from healthy donors. All patients were diagnosed based on MRI and CT scanning, and had a history of receiving long-term or high-dose steroid therapy. No subject in this study had exposure to all known ONFH risk factors for at least three months before serum samples were obtained. The healthy participants were recruited from the community. All participants in the control group received a general medical examination in Shanghai Jiao Tong University Affiliated Sixth People’s Hospital and no systemic diseases such as hypertension or diabetes were detected. The primary diseases of the ONFH patients are summarised in the supplementary material. All ONFH patients were classified with the Steinberg staging system of ONFH.

Exosome concentration detection. All the serum samples were immediately frozen and stored at −80°C after isolation from donors. Before detection, randomly generated, eight-digit numbers were assigned to each sample. HZ kept the group information and diluted all the samples with phosphate-buffered saline at a ratio of 1:10. YG detected the exosome level. YW then obtained the group information from the first researcher and was responsible for data analysis. The exosome concentration was determined by the number of particles of size 30 nm to 120 nm using Nanosight (Malvern, United Kingdom). All data in this study were expressed as mean with standard deviation (SD). Receiver operating characteristics (ROC)
Results
The demographic characteristics of the patients. The demographic features of all subjects are shown in Table I. There were no significant differences in age, gender, height or weight between the oNFH and control groups.

The level of circulating exosomes in steroid-induced ONFH patients was lower than that of healthy donors. The circulating exosome level was determined by detecting the number of particles with sizes ranging from 30 nm to 120 nm (Fig. 1). The exosome level in the control group was $9.1 \times 10^{13}$/ml (sd 2.5). For the patient group, the serum exosome level was $7.3 \times 10^{13}$/ml (sd 2.0). The serum exosome level was significantly lower in the patient group than in the healthy donors ($p < 0.01$, Student’s $t$-test).

Diagnostic value of serum exosome for diagnosis of steroid-induced ONFH. The diagnostic value of circulating exosome levels for steroid-induced ONFH were then calculated by ROC curve analysis (Fig. 2). Circulating exosome levels showed moderate diagnostic accuracy for the presence of steroid-induced ONFH (Table II). We further demonstrated that the level of circulating exosomes is not predictive of the stages of steroid-induced ONFH (Table III).

Discussion
Intervention before collapse of the femoral head is the key to success in hip preservative operations. However, the steroid-induced ONFH is a latent disease and usually develops several years after intensive steroid exposure. Additionally, early-stage ONFH is usually asymptomatic, highlighting the need for a convenient laboratory test to detect potential ONFH development. The present study first evaluated the circulating exosome level in patients with steroid-induced ONFH. We demonstrated that the serum exosome level was significantly lower in patients with steroid-induced ONFH compared with healthy donors. ROC analysis demonstrated that the serum exosome level had moderate diagnostic accuracy for the detection of steroid-induced ONFH. The serological examination is more convenient and economical than traditional MRI scanning. After screening by serum exosome level detection, MRI scanning should be conducted for definitive diagnosis.

It is also possible that the change in exosome levels might even precede the alteration in MRI, such as showing the ‘double-line’ sign. Thus, it would be interesting to know whether circulating exosomes can be used as ‘super early’ diagnosis markers for steroid-induced ONFH. A prospective study on a population with intensive steroid exposure might address this question.

In conclusion, circulating exosome level is valuable in the diagnosis of steroid-induced ONFH.

Supplementary Material
A table showing a summary of the primary diseases of the ONFH patients can be found alongside this paper at http://www.bjr.boneandjoint.org.uk/

References

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**Author Contribution**
- H-Y. Zhu: Study design.
- Y-C. Gao: Collection of serum samples and patient information, Exosome level detection.
- Y. Wang: Data analysis and manuscript writing.
- C-Q. Zhang: Study design and manuscript revision.
- Y-C. Gao is co-first author.

**ICMJE conflict of interest**
- None declared

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