Efficacy of autologous leukocyte-reduced platelet-rich plasma therapy for patellar tendinopathy in a rat treadmill model

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Summary

Background: An autologous platelet-rich plasma (PRP) therapy has currently been applied for the tendinopathy; however, its efficacy and an optimal platelets concentration in PRP were uncertain. We analyzed them in an animal model prepared using a repetitive running exercise.

Methods: We made the tendinopathy rat model of patellar tendon using a rodent treadmill machine. Rats with tendinopathy were injected with leukocyte-reduced PRP at the platelets concentration of 1.0×10⁶/µL (P10 group), PRP at the platelets concentration of 5.0×10⁵/µL (P5 group) or normal saline (control group) into the space between the patellar tendon and the fat pad bilaterally or were multiply dry-needled at the tibial insertion site (MN group) at once. To assess the pain-relieving effect, the spontaneous locomotor activities at night (12 h) were measured every day. Histological sections of the patellar tendon stained with hematoxylin-eosin or prepared by TdT-mediated dUTP nick end labeling were microscopically analyzed.

Results: The numbers of spontaneous locomotor activities in the P10 group were significantly larger than those in the P5, MN or control groups and they recovered up to a healthy level. On histological examinations, the numbers of microtears, laminations, or apoptotic cells in the patellar tendons in the P10 or P5 groups were significantly lower than those in the MN or control groups, although no significant differences were observed between the P10 and P5 groups.

Conclusions: The injections of an autologous leukocyte-reduced PRP were effective for pain relief and for partial restoration of the patellar tendon in the tendinopathy rat model. The injections of a PRP at the platelets concentration of 1.0×10⁶/µL completely relieved the pain and were more effective than those at the platelets concentration of 5.0×10⁵/µL whereas there was no difference for the effect of histological restoration or apoptosis inhibition between them.

KEY WORDS: platelet-rich plasma, rat, tendinopathy.

Introduction

Tendinopathy is the most common tendon disorder characterized by activity-related pain, focal tendon tenderness and decreased strength and movement in the affected area. Tendinopathy can occur in almost any tendon and if developed, it impairs active performance in athletes and workers of many occupations that involve repetitive movements. Some of the most common tendinopathies are tennis elbow, jumper’s knee and Achilles tendinopathy1. In the current decade, a therapy of autologous platelet-rich plasma (PRP) has been clinically applied for the treatment of the tendinopathy as tennis elbow, jumper’s knee, Achilles tendinopathy, plantar fasciitis and other tendinopathies. Significant therapeutic effects of PRP were showed for lateral or medial epicondylitis in elbows, jumper’s knee and Achilles tendinopathy compared to the conventional therapies as the injection of corticosteroid or a focused shock-wave therapy2-12. On the other hand, there were some clinical examinations which showed no significant effects of PRP for the treatment of Achilles tendinopathy13, 14. However, there were few high-level literature as randomized controlled trials regarding the application of PRP in the management of patellar and Achilles tendinopathy. An efficacy and effectiveness of PRP therapy were still uncertain for the treatment of patellar or Achilles tendinopathy at the present.

The majority of the basic research has demonstrated a beneficial effect of PRP on the proliferation of tenocytes, collagen synthesis and the tendon healing process15. In the tendinopathy animal model, Dallaudière demonstrated that a mono-injection of PRP improved tendon healing with no local toxicity in a rat collagenase-induced tendinopathy of patellar or...
Achilles tendon\textsuperscript{18} and Chen demonstrated that a PRP treatment resulted in better healing of injured tendons with improved histological outcomes and biomechanical functions in a rat collagenase-induced tendinopathy of Achilles tendon\textsuperscript{17}. As the animal model for the patellar tendinopathy, it was suggested that a model prepared using a repetitive running exercise on a rodent treadmill machine pathophysiologically reproduced well an actual tendinopathy compared to the collagenase-induced model, since there was no inflammation reaction in/around the tendon in actual tendinopathy although the inflammation was observed in/around the tendon in the collagenase-induced tendinopathy and the clinical tendinopathy was pathogenetically induced with the over-use and/or over-load stress like a repetitive running exercise for a long distance\textsuperscript{18-21}. The previous basic researches described above and also the current animal study\textsuperscript{22} were performed using the collagenase-induced model and there was no examination, which examined the therapeutic effect of PRP for the tendinopathy in an animal model prepared using a repetitive running exercise on a rodent treadmill machine to our knowledge at the present. We have experienced to make this animal model and have examined the study using the rat tendinopathy model prepared by a repetitive running exercise\textsuperscript{23}.

Regarding the optimal concentration of the platelets in PRP for the tendon healing, some researchers reported \textit{in vitro} studies. McCarrell demonstrated that the PRP at the platelets concentration of 1.2x10\textsuperscript{5}/µL led to a paradoxical effect on collagen synthesis on the culture of horse tendons compared to the PRP at the intermediate concentration of the platelets, while also presenting an increase in inflammatory mediators\textsuperscript{24}. Boswell also demonstrated that the PRP at the high platelets concentration reduced tendon metabolism and collagen synthesis in equine tendon culture\textsuperscript{25}. Moreover, Giusti demonstrated that the PRP at the platelets concentration between 5.0x10\textsuperscript{5}/µL and 1.0x10\textsuperscript{6}/µL stimulated cell proliferation and migration as well as collagen production in human tenocyte cultures while higher platelet concentrations have an inhibitive effect both in terms of cell metabolism and collagen synthesis\textsuperscript{26}. From those information it seemed that the platelets concentration between 5.0x10\textsuperscript{5}/µL and 1.0x10\textsuperscript{6}/µL in PRP was optimal for the tendon healing. There was no examination which analyzed the difference for the efficacy of a PRP therapy at the different platelets concentration in the tendinopathy animal model. The optimal concentration of the platelets in PRP was still uncertain for the therapy of the tendinopathy.

Regarding the influence of the leukocytes in PRP against the metabolism of tenocytes, Dragoo demonstrated that PRPs rich in leucocytes led to a significant inflammatory cellular reaction in rabbit tendons at 5 days after PRP infiltration\textsuperscript{27}. McCarrell demonstrated that the PRP with a high concentration of leucocytes led to higher expression of pro-inflammatory cytokines and catabolic mediators as well as lower collagen synthesis\textsuperscript{24}. It appeared that the leucocyte inhibited the restoration process for the treatment of the tendinopathy since the inflammation damaged the extracellular matrix and a lower synthesis of the collagen molecules suppressed the conformation of the tendon matrix. Hence, we used a leucocyte-reduced PRP for the present study.

We investigated the therapeutic effect of an autologous leucocyte-reduced PRP for tendinopathy in a rat model prepared using a repetitive running exercise on a rodent treadmill machine in the present study. An efficacy for a pain-relieving or a histological tendon tissue-repair of PRP was scientifically analyzed. Moreover, we evaluated the therapeutic effects of an autologous PRP at the different concentration of the platelets between 5.0x10\textsuperscript{5}/µL and 1.0x10\textsuperscript{6}/µL in the same manner. Furthermore, we analyzed the effectiveness of a multiple dry-needling performed at the tendinopathy site in order to know its contribution for the treatment of the tendinopathy. The present study was designed and underwent according to the journal’s editorial\textsuperscript{28}.

### Materials and methods

#### Reagents and materials

The autologous PRP preparation system, MyCells, was purchased from Kaylight, Tel Aviv, Israel. The normal saline for injections was from Otsuka Pharmaceutical Factory, Inc., Naruto, Japan. A rodent treadmill, KN-73, was purchased from Natsume Seisakusho, Co., Ltd., Tokyo, Japan. The Supermex system for measurement of spontaneous locomotor activities of rats was from Muromachi Kikai, Tokyo, Japan. The ApopTag Peroxidase In Situ Apoptosis Detection Kit was from EMD Millipore Corporation, Billerica, MA, USA.

#### Animals

Male Wistar rats (16-18 weeks of age, weighing 350-400 g) were purchased from Nippon SLC, Hamamatsu, Shizuoka, Japan, and served as subjects in the study. Rats were housed in an environmentally controlled animal facility on a 12:12 light/dark cycle with food and water available \textit{ad libitum} in the laboratory animal section.

#### The animal model

All experiments were conducted in accordance with the institutional guidelines for the care and use of experimental animals after an institutional approval for the study was obtained. A rat model of patellar tendinopathy was made according to the method of Flatow et al.\textsuperscript{18-21, 23}. A running exercise was performed on a rodent treadmill with 10\% inclination of the running protocol. The rats were acclimated to the treadmill by gradually increasing running speed and
time for 2 weeks before the running exercise. Running distance reached up to 40 km during 40 days and the mean running speed was 0.5 km/h. The spontaneous locomotor activities at night were measured for 12 h with the Supermex system on the day before the running exercise and on the day after exercise. The reduction in spontaneous locomotor activity after exercise was different among rats. The rats showing > 25% reduction (25-33%) in spontaneous locomotor activity after exercise were then selected as the tendinopathy rats for later study. Approximately 85% of the rats ran the whole distance and approximately 90% of them revealed > 25% reduction in locomotor activities. In pre-study, we made histological sections of the patellar tendons or rotator cuffs in the shoulders in ten rats, which revealed > 25% reduction in locomotor activities or in five rats which revealed no reduction in activities after running exercise. Then, the tendinopathy-specific histopathological changes were observed at the insertion site of the patellar tendon with light microscopy in all ten activity-reduced rats (Fig. 3) whereas no tendinopathy-specific histopathological changes were observed in five activity-non-reduced rats. Moreover, less tendinopathy-specific histopathological changes were observed at the insertion site of the rotator cuffs in shoulders in both rats. We finally made 30 tendinopathy rats for the present study from 50 rats.

**Group division**

The rats with tendinopathy were divided into four groups. The rats in the P10 group, the P5 group or the control group were injected with autologous leukocyte-reduced PRP at the platelets concentration of 1.0×10^6/μL, autologous leukocyte-reduced PRP at 5.0×10^5/μL or normal saline, respectively, for further examinations. The MN group consisted of eight rats. Rats in fourth, the multiple dry-needling (MN) group, were pin-pricked ten times using a 27G injection needle at the tibial insertional site of the patellar tendon without injections. The MN group consisted of six rats.

**Observation of general status and body weight measurements**

The general status was observed once a day from the day before the running exercise started to the end of the study. Body weight was measured at every 5 days during the study period.

**Preparation of autologous platelet-rich plasma**

We used an autologous PRP preparation system, MyCells, to obtain a rat autologous leukocyte-reduced PRP since the system could prepare the leukocyte-reduced PRP at high platelet concentration and it was hard to purchase the other PRP preparation kits like Biomet GPSIII, Arteriocyte Magellan or PRGF in our nation. A whole blood of 10 mL was aspirated from the hearts in 5 rats and a leukocyte-reduced PRP of 2.0 mL was obtained by using a PRP preparation system according to the manufacture’s instruction. Briefly, a whole blood of 10 mL was inserted into the separation slit of the system and it was centrifuged for 7 minutes at 2000 x g. After aspiration of the supernatant plasma, the residual plasma of 2.0 mL was pipetting well in order to dissolve the platelets which were precipitated on the separation gel. Then, the filter column was inserted into the separation slit to exclude the debris and a filtered non-activated PRP of 2.0 mL was obtained by aspiration from the filtration column. A hundred μL of a prepared PRP was mixed in the normal saline at the final volume of 1.0 mL and the number of the platelets or leukocytes was counted. We obtained a rat autologous PRP at the platelets concentration of 1.3×10^6/μL containing the leukocytes at the approximate half concentration of the one in whole blood and no erythrocytes. An obtained PRP was classified as type 3 in Mishra’s classification and was classified as pure PRP in PAW classification since we did not activate a PRP before injection. Then, we made a PRP of 1.0 mL at the platelets concentration of 1.0×10^6/μL or 5.0×10^5/μL, respectively, for further examinations.

**Injections of autologous PRP or normal saline into hind paws**

After general anesthesia with 2.5% isoflurane, the fur at the knee joints was shaved and 25 μL of autologous leukocyte-reduced PRP or normal saline was injected into the space between the patellar tendon and the fat pad tissues at the distal area of the patellar tendon under ultrasonography (US) guidance using a 0.5 mL injection syringe and a 27G injection needle from lateral side of the knees, respectively, in the P10, P5 or control groups. Before injection the platelets in PRP were not activated. Since there was a possibility that the direct injection into the tendon produced the tears or laminations in the tendon, we avoided the direct injection into the tendon and selected a space between tendon and fat pad as a target for injection. In pre-study, we injected 30 μL of PRP or saline containing the indigo carmine dye into the space between patellar tendon and fat pad in ten rats and we confirmed that there was no damage in the tendon and no leakage into the cavity of knees in ten rats by the direct inspection after surgical open of the knees. Therefore, we injected 25 μL of PRP or saline for more prevention of the leakage. Then, one technician practiced the injection of PRP or normal saline containing the indigo carmine dye for twenty times under ultrasonography (US) guidance followed by confirmation whether there was a leakage or not with the direct inspection after surgical open of the knees. After the technician mastered the way of the injection, the injections were performed once at the next day following completion of the running exercise (Fig. 1).
Measurements of spontaneous locomotor activities

Spontaneous motor activity of each rat was examined using the Supermex system for 12 h each night for 17 days after injections of PRP or saline or multiple dry-needling in all groups. The Supermex system could quantify the locomotor activities, which reflected the degree of the pain in limbs of rats. Thus, the degree of pain-relief in the knee could be quantified by the measurement of the locomotor activities. The system was used in many animal studies, in which the degree of the pain-relief was quantitatively measured, and the reliability of the system was highly evaluated comparing to other methods in the literatures23, 30-33. Details of spontaneous locomotor activity measurement have been described previously23, 30-33. Briefly, spontaneous locomotor activities were determined by detecting the movement of infrared radiation emitted from each animal. The activities were measured as a single count when an animal moved from one region of the measurement area divided optically by multiple lenses, to a neighboring region. Total counts were calculated by summing up all counts of 10-min periods. The infrared filter with 25% signal reduction was used to remove the noise of the grooming motion. The relative number of motor activities in each group was calculated and the mean of the relative changes was plotted on a graph in order to evaluate the relative changes for motor activities of rats during the experiment period, since the absolute values for motor activities were different among rats, which were 9787±1982 (6703-14191) counts for 50 healthy rats. A statistical analysis was performed to examine the outcome differences between each group.

Histological examinations

The knee tissue samples were obtained from 6 healthy normal rats and 6 tendinopathy rats at day 1 (controls), and from 6 rats in the P10, the P5, the MN or the control groups, respectively, at day 18. Following inhalation with 5% isoflurane, the rats were killed by immediate blood draw. Whole knee joints were fixed in 4% paraformaldehyde at pH 7.4 for 3 days, and decalcified in 20% EDTA solution for 21 days at 4 °C, and then embedded in paraffin wax. Whole knee joints were sectioned in the sagittal plane with a section width of 5 μm. The sections were stained with hematoxylin-eosin (H&E) for histology. They were assessed with a histopathological score determined by counting the number of microtears and laminations located at the insertion site of the patellar tendon as a point in high-power (magnification 200x) fields in each five sections of tendinopathy rats or P10, P5, MN or control groups. A point in the score was doubled in cases where the length of tears or laminations were longer than half the diameter of the visual high-power field (magnification 200x) on the microscope23. The histological sections were also prepared by TdT-mediated dUTP nick end labeling using the ApopTag Peroxidase In Situ Apoptosis Detection Kit according to the manufacturer’s instructions. Tendon cells for apoptosis were quantified by counting the number of TUNEL-positive cells in high-power (magnification 400x) fields at the joint side of the tibial insertion site for the patellar tendon in each five sections of tendinopathy rats or P10, P5, MN or control groups. Three researchers counted the histopathological score in H&E sections and the number of apoptosis cells in all samples respectively. The standard deviations of these scores or numbers in each histological sample among three researchers were within 5%. The data values were expressed as the mean of the differences among groups after statistical analysis23.

Statistical analysis

Comparative series were considered independent of each other. A mean value and standard deviation was calculated for each locomotor activity in each group or for the histopathological score and the cell number of apoptosis in each group. We compared the daily locomotor activities, the histopathological scores and the numbers of apoptosis among groups. Statistical analysis was done using the Wil-
coxon’s matched-pairs signed-rank test. A probability value of < 0.05 was considered statistically significant.

Results

General status and body weight

There were no abnormalities observed in any of the groups during the experiment period. The change of body weight during the injection period is shown in Table I. All groups showed a similar trend of body weight changes compared to the control group.

Spontaneous locomotor activities

The counts for the spontaneous locomotor activities in the P10 group increased up to approximate 1.4-fold at day 3 and were maintained, not decreased, for 15 days until the end of the study without any additional PRP injections. Spontaneous motor activities in the P10 group recovered up to the original level as before commencement of the running exercise. On the other hand, the counts for the spontaneous locomotor activities in the P5 group increased up to approximate 1.2-fold at day 3 and were maintained, not decreased, for 15 days until the end of the study without any additional PRP injections. There were significantly more counts for the spontaneous locomotor activities in the P10 group than in the P5, the MN or the control groups (p < 0.05) and there were significantly more counts for the spontaneous locomotor activities in the P5 group than in the MN or the control groups (p < 0.05) during 16 days of the experimental periods from day 2 until the end of the study. The counts of the locomotor activities in the MN and the control groups did not significantly increase and remained in the range of 80 to 120% of the initial activities. No significant differences were observed for the counts of locomotor activities between the MN group and the control group. The counts for the spontaneous locomotor activities in all groups displayed a trend to gradually increase up along with the days during experimental periods, although no statistical significances were observed (Fig. 2).

Histological findings

Two major pathological findings were found at the tibial insertion sites of the patellar tendons in tendinopathy rats in all the histological sections stained with hematoxylin-eosin. They were microtears in the tendon body and lamination, i.e., several longitudinal disruptions between the fiber bundle layers (Fig. 3). The same pathological findings were found at the tibial insertion sites of the patellar tendon at day 18 in the P10, P5, MN or control groups in the histological knee sections stained with hematoxylin-eosin (Fig. 4). However, the histopathological scores determined by the number and length of the microtears and laminations in five rats of the P10 or P5 groups were significantly lower than those in five rats of the MN or control groups (p < 0.05) (Tab. II, Fig. 4), indicating that the reactions of tendon repair in the P10 or P5 groups were progressed compared to those in the MN or control groups. No significant differences were observed in the histopathological scores between the P10 group and the P5 group or among tendinopathy rats, MN group and control group (Tab. II). TUNEL-positive tendon cells were observed at the joint side of the tibial insertion site for the patellar tendon in all tendinopathy rats. The number of TUNEL-positive cells at the same area in five rats of the P10 or P5 groups was significantly lower than those in five rats of the tendinopathy rats, MN group or control group at day 18 (p < 0.05) (Tab. III, Fig. 5). No significant differences were observed in the number of TUNEL-positive cells between the P10 group and the P5 group or among tendinopathy rats, MN group and control group (Tab. III, Fig. 5).

Discussion

In the present study, we created an insertional tendinopathy model of the patellar tendon in rats and examined the therapeutic effects of autologous leukocyte-reduced PRP injections. The following findings were obtained: (1) the injections of autologous leukocyte-reduced PRP into the space between the patellar tendon and the fat pad tissue significantly increased the number of spontaneous locomotor activi-
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Figure 2. Relative values for spontaneous locomotor activities in the P10, P5, MN or control groups. The spontaneous locomotor activities in rats were examined using a Supermex system at each night for 12 h, every day for 17 days, after completion of the running exercise and after commencement of the injection protocol. The relative number of locomotor activities in each group was calculated and the means or the SD were plotted on a graph.

Figure 3. Histological sections of the tibial insertion sites of the patellar tendons stained with H&E in a healthy rat (a) or in the tendinopathy rat (b) (×100). Two major pathological findings were found with light microscopy at the tibial insertion site of the patellar tendons in the tendinopathy rats: (1) microtears in the body of the tendon and (2) several longitudinal disruptions between the layers of the fiber bundles of the tendon body, so-called laminations.
Platelet-rich plasma injection for tendinopathy

![Image](https://example.com/image)

Figure 4. Histological sections of the tibial insertion sites of the patellar tendons stained with H&E in the P10 group (a), in the P5 group (b), in the MN group (c) or in the control group (d) (x100). The number of microtears or laminations found in 5 rats of the P10 or the P5 groups was significantly lower than those in 5 rats of the MN or control groups (P < 0.05). No significant difference was observed in the number of microtears or laminations between the P10 group and the P5 group or among tendinopathy rats, MN group and control group.

Table II. Histopathological score for the histological sections located at the tibial insertion site of the patellar tendon.

<table>
<thead>
<tr>
<th>Group</th>
<th>Histopathological score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tendinopathy</td>
<td>16 ± 5</td>
</tr>
<tr>
<td>Control</td>
<td>15 ± 6</td>
</tr>
<tr>
<td>MN</td>
<td>17 ± 7</td>
</tr>
<tr>
<td>P10</td>
<td>6 ± 2</td>
</tr>
<tr>
<td>P5</td>
<td>7 ± 3</td>
</tr>
</tbody>
</table>

*p < 0.05

Table III. Cell number for apoptosis at the joint side of the tibial insertion site for the patellar tendon.

<table>
<thead>
<tr>
<th>Group</th>
<th>Apoptosis cell</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tendinopathy</td>
<td>24 ± 10</td>
</tr>
<tr>
<td>Control</td>
<td>23 ± 12</td>
</tr>
<tr>
<td>MN</td>
<td>25 ± 12</td>
</tr>
<tr>
<td>P10</td>
<td>6 ± 2</td>
</tr>
<tr>
<td>P5</td>
<td>7 ± 3</td>
</tr>
</tbody>
</table>

*p < 0.05

ties in the tendinopathy rats compared to those in the control or multiple needling groups; (2) the spontaneous locomotor activities in rats which were injected with autologous leukocyte-reduced PRP at the platelets concentration of 1.0x10⁶/µL recovered up to the original healthy level at 2 days after injections and were then maintained, not decreased, for the following 15 days until the end of the study without any additional injections of PRP; (3) the spontaneous locomotor activities in the rats which were injected with autologous leukocyte-reduced PRP at the platelets concentration of 1.0x10⁶/µL were significantly greater than those in the rats which were injected with PRP at the platelets concentration of 5.0x10⁶/µL during the experimental periods for 16 days from day 2 until the end of the study; (4) the multiple dry-needling against the insertional tendinopathy site of the patellar tendon did not significantly increase the sponta-
neous locomotor activities and no significant differences in the spontaneous locomotor activities were observed between the MN group and the control group; (5) the number of microtears or laminations observed in histological specimens at the tibial insertion site of the patellar tendon in tendinopathy rats decreased in the PRP group compared to those in the control or multiple needling groups at day 18; (6) the number of TUNEL-positive cells at the joint side of the tibial insertion site of the patellar tendon in the PRP group was lower than those in the control or multiple needling groups at day 18; (7) no significant differences were observed in the histopathological scores or the number of TUNEL-positive cells between the P10 group and the P5 group or among tendinopathy rats, MN group and control group.

The findings of (1) and (2) indicated that an autologous leukocyte-reduced PRP injection was effective and rapid analgesic method in tendinopathy rats, since no remarkable tendinopathy-specific histological changes were observed in the rotator cuffs of the shoulders in tendinopathy rats. The present study for the first time demonstrated the analgesic effect of the autologous leukocyte-reduced PRP for the treatment of the tendinopathy in the rat model prepared by a repetitive running exercise on a rodent treadmill machine. Regarding the mechanism for the generation of the pain in the tendinopathy, there were several studies which showed that the abnormal capillary vessels with shunts were conformed around the tendon, not intra tendon, in the tendinopathy. It was suggested that the pain was generated from the free nerve endings, which were distributed along with the abnormal capillary vessels especially conformed in the tendinopathy, since the focal embolization of the abnormal capillary vessels immediately removed the pain in tendinopathy. Therefore, it was speculated that the abnormal capillary vessels with free nerve endings were degraded by a reaction induced with an autologous leukocyte-reduced PRP, although no distinct capillary vessels were histologically found at the tibial insertional sites of the patellar tendon in our tendinopathy rats.

The findings of (2) and (3) indicated that the analgesic effect of PRP at the platelets concentration of $1.0 \times 10^6/\mu L$ was greater than that of PRP at the platelets concentration of $5.0 \times 10^5/\mu L$ in the tendinopathy rat model and that an injection of leukocyte-reduced PRP at the platelets concentration of $1.0 \times 10^6/\mu L$ completely relieved the pain in the tendinopathy rat. It was estimated that the abnormal capillary vessels with free nerve endings were rapidly degraded enough to completely relieve the pain by a reaction induced with a leukocyte-reduced PRP at the platelets concentration of $1.0 \times 10^6/\mu L$. The finding of (7) indicated that the effects for the histological restoration and the apoptosis inhibition of leukocyte-reduced PRP at the platelets concentration of $1.0 \times 10^6/\mu L$ or $5.0 \times 10^5/\mu L$ were almost identical whereas the analgesic effects between them were
late-stage tendinopathy. It was still uncertain whether the PRP therapy was effective for the late-stage tendinopathy or not. Further examinations were necessary to dissolve those problems.

Conclusion

One injection of an autologous leukocyte-reduced PRP into the space between the patellar tendon and fat pad effectively relieved pain and initiated partial restoration of the patellar tendon in the tendinopathy rat model. The analgesic effects of the leukocyte-reduced PRP at the platelets concentration of 1.0×10^6/L was greater than those at the half concentration of the platelets, although no difference was observed for the effect of the histological restoration between them. A multiple dry-needleing showed no effects for the treatment of the tendinopathy. It was suggested that an administration of an autologous leukocyte-reduced PRP was clinically effective modality in treatment of tendinopathy.

Abbreviations

PRP Platelet-rich plasma
NS Normal saline
EDTA Ethylenediamine tetra-acetic acid
H&E Hematoxylin-eosin
TUNEL TdT-mediated dUTP nick end labeling

Key points

Our goal was to assess the effects of pain-relieving and histological restoration for PRP therapy and to determine the optimal concentration of platelets in leukocyte-reduced PRP for the treatment of the tendinopathy.

We used patellar tendinopathy in a rat model prepared using a repetitive running exercise on a rodent treadmill machine with adequate controls.

We measured spontaneous locomotor activities of rats for 12h every day for the assessment of a pain-relieving and we histologically analyzed a restoration of the tendon and an occurrence of apoptosis.

We demonstrated that a PRP therapy was effective for pain-relieving and histological restoration of the tendinopathy. The injections of a PRP at the platelets concentration of 1.0×10^6/L completely relieved the pain and were more effective than those at the platelets concentration of 5.0×10^5/L whereas there was no difference for the effects of histological restoration and apoptosis inhibition between them.

We estimated that the platelets concentration between 5.0×10^5/L and 1.0×10^6/L in leukocyte-reduced PRP was optimal for the treatment of the tendinopathy rat model.

The study results suggested that an administration of an autologous leukocyte-reduced PRP was clinically effective modality in the treatment of the tendinopathy.

Conflicts of interests

The Authors declare that they have no competing interests.
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