



BMJ Open Combination of mesenchymal stem cells (MSCs) and platelet-rich plasma (PRP) in the treatment of knee osteoarthritis: a meta-analysis of randomised controlled trials

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ABSTRACT

Objectives The purpose of this meta-analysis was to investigate the efficacy and safety of mesenchymal stem cells (MSCs) combined with platelet-rich plasma (PRP) in the treatment of knee osteoarthritis (KOA).

Design Systematic review and meta-analysis.

Participants Patients with KOA.

Interventions Use of MSCs+PRP.

Primary and secondary outcomes Visual Analogue Scale (VAS) score, Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) score, Knee Injury and Osteoarthritis Outcome Score (KOOS) and adverse reactions.

Data sources PubMed, Cochrane Library, Embase and China National Knowledge Infrastructure were searched from inception to 15 July 2021.

Measures The OR or weighted mean difference (WMD) of relevant outcome indicators was calculated. Study quality was evaluated using the risk-of-bias assessment tool version 2.0. Heterogeneity among studies was evaluated by calculating I^2 . If $I^2 < 50\%$, a fixed-effect model was applied; conversely, if $I^2 \geq 50\%$, a random-effect model was applied.

Results Six controlled clinical trials with 493 cases were included. The meta-analysis results showed that in terms of the VAS score 3 months after treatment, MSCs+PRP had no significant effect on the reduction of the VAS score in patients with KOA compared with the control ($p=0.09$), hyaluronic acid (HA) ($p=0.15$) or PRP alone ($p=0.07$). MSCs+PRP was more effective in reducing the VAS score at 6 and 12 months after treatment than the control (WMD=-0.55, 95% CI -0.87 to -0.22, $p<0.001$), HA (WMD=-1.20, 95% CI -2.28 to -0.13, $p=0.03$) or PRP alone (WMD=-0.54, 95% CI -0.89 to -0.18, $p=0.003$). Regarding the decrease in the total WOMAC score at 3 and 6 months after treatment, MSCs+PRP showed better clinical efficacy than the control or HA alone ($p<0.01$). Compared with the control, MSCs+PRP exhibited no significant difference in reducing the total WOMAC score 12 months after treatment ($p=0.39$). There was no significant difference between MSCs+PRP and the control in terms of improvement of the KOOS 12 months after treatment ($p=0.16$). Compared with MSCs alone,

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ This is the first meta-analysis to investigate the efficacy and safety of mesenchymal stem cells (MSCs) combined with platelet-rich plasma (PRP) in the treatment of knee osteoarthritis (KOA).
- ⇒ Most of the included studies were of high quality with a low risk of bias.
- ⇒ The number of studies and sample sizes included in this meta-analysis was limited.
- ⇒ The follow-up time of the included studies was short, which is not conducive to deducing the long-term efficacy of MSCs combined with PRP for the treatment of KOA.

MSCs+PRP exhibited no significant difference in the incidence of adverse reactions ($p=0.22$) 12 months after treatment.

Conclusions Treatment with MSCs+PRP showed good clinical efficacy in improving pain and joint function in patients with KOA. Compared with MSCs alone, there was no significant difference in the incidence of adverse reactions with MSCs+PRP.

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INTRODUCTION

With the increase in the ageing population worldwide, the incidence rate of knee osteoarthritis (KOA) has increased annually.¹ Epidemiological investigations have shown that KOA is more common in middle and old age groups, especially in the population over 65 years old, in which the incidence rate is approximately 50%.² The effective prevention and treatment of KOA has become the key to improving the quality of life of middle-aged and elderly individuals.² The purpose of non-surgical treatment for KOA is to reduce pain, improve function and avoid complications of surgical treatment as much as possible. Conservative pain control methods include

drug treatment, increased daily activity and periarticular analgesia, but the optimal method is still controversial.^{3 4}

In recent years, cell therapy has shown good potential for the treatment of many diseases, including osteoarthritis, and studies have preliminarily shown that bone marrow mesenchymal stem cells (MSCs) can promote the repair of articular cartilage and be used to treat osteoarthritis.^{5 6} In addition, adipose-derived stem cells can be directionally differentiated into chondrocytes and osteoblasts, playing a role in repairing and treating osteoarthritis.^{7 8} Platelet-rich plasma (PRP) contains a large number of growth factors, such as platelet-derived growth factor and vascular endothelial growth factor, which play an important role in cartilage repair.^{9 10} Previous studies have shown that PRP has good safety and efficacy in the treatment of KOA, but it shows only short-term clinical effects.^{11 12} Interestingly, *in vitro* studies have shown that PRP can promote the proliferation and differentiation of mesenchymal stem cells (MSCs).¹³ Coculture of MSCs with PRP *in vitro* can promote the proliferation and chondrogenic differentiation of MSCs.¹⁴ In addition, *in vivo* animal experiments have confirmed that the combination of PRP and MSCs can promote the healing of rabbit radial bone defects more than MSCs alone.¹⁵ Moreover, we believe that the injection of MSCs and PRP into the knee can theoretically promote chondrocyte regeneration, eliminate nonbacterial inflammation of the synovium, and alleviate symptoms through the aetiological treatment of osteoarthritis.^{16 17}

PRP has shown good safety and efficacy in treating KOA, but the therapeutic effect decreases with increasing patient age.^{18 19} In addition, studies have shown that MSCs can promote the repair of articular cartilage to treat KOA.²⁰ Whether there is a synergistic effect of PRP combined with MSCs in the treatment of KOA has attracted the attention of researchers. However, the efficacy and safety of MSCs+PRP in the treatment of KOA are still controversial, and there is a lack of higher-level clinical evidence. This study will use meta-analysis and quantitative evaluation to explore the efficacy and safety of MSCs+PRP in the treatment of KOA to provide an evidence-based foundation for clinical application and basic research.

MATERIALS AND METHODS

This meta-analysis was conducted in strict accordance with the Preferred Reporting Items for Systematic reviews and Meta-Analyses statement.²¹

Data source and retrieval strategy

Two researchers conducted a comprehensive search of four databases: PubMed, Cochrane Library, Embase and China National Knowledge Infrastructure (CNKI). The retrieval time limit was from the establishment of each database to 15 July 2021. A combination of subject words and free words was used to establish a retrieval model to search the above four databases. The search terms

included “mesenchymal stem cell”, “stem cell”, “platelet-rich plasma”, “knee osteoarthritis” and “osteoarthritis”. The search strategy for each database can be found in online supplemental material 1.

Inclusion and exclusion criteria

The inclusion criteria were as follows: (1) All the included patients had KOA, and clear diagnostic criteria for KOA were described in the study. (2) The experimental group was treated with MSCs combined with PRP, and the control group was treated with MSCs, PRP or hyaluronic acid (HA) alone. It should be noted that the source of MSCs could be bone marrow, adipose tissue or umbilical cord tissue. (3) The type of study was a clinical controlled trial. (4) The relevant outcome indicators of clinical efficacy or safety were reported in the literature. To better evaluate the changes after treatment in patients with KOA, the Visual Analogue Scale (VAS) and Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) scores were selected as primary outcomes, and the Knee Injury and Osteoarthritis Outcome Score (KOOS) and adverse reactions were selected as secondary outcomes. (5) There were no restrictions regarding the language of the published studies.

The exclusion criteria were as follows: (1) incomplete original data; (2) results of repeated studies in the same population and (3) interventions involving surgical treatment.

Data extraction and literature quality evaluation

Two researchers independently conducted literature screening and data extraction according to the inclusion and exclusion criteria. If there were different opinions, they were discussed and resolved with a third researcher. The researchers standardised the relevant data included in the literature. The extracted data included basic information of the study (first author, year of publication, sample size, patient age, etc), outcome indicators and information related to the literature quality evaluation.

Study quality was evaluated using the risk-of-bias assessment tool version 2.0 (ROB 2.0) recommended by Cochrane.²² The evaluation tool evaluates the risk of bias in five areas. If the evaluation results in five areas indicate low risk, the overall risk of bias is low. If the assessment result of any one of these areas indicates high risk or the assessment result of multiple areas indicates possible risk, the overall risk is high. If neither of the above two conditions is met, the clinical control trial is judged as having a possible risk of bias.

Data analysis

Review Manager V.5.3 software (Cochrane Collaboration, UK) was used for data analysis in this meta-analysis. Continuous variables are represented by the weighted mean difference (WMD) and 95% CI, and classified variables are represented by the OR and 95% CI. The WMD and OR were calculated in our study referring to the method in the Cochrane system evaluation manual.²³ A

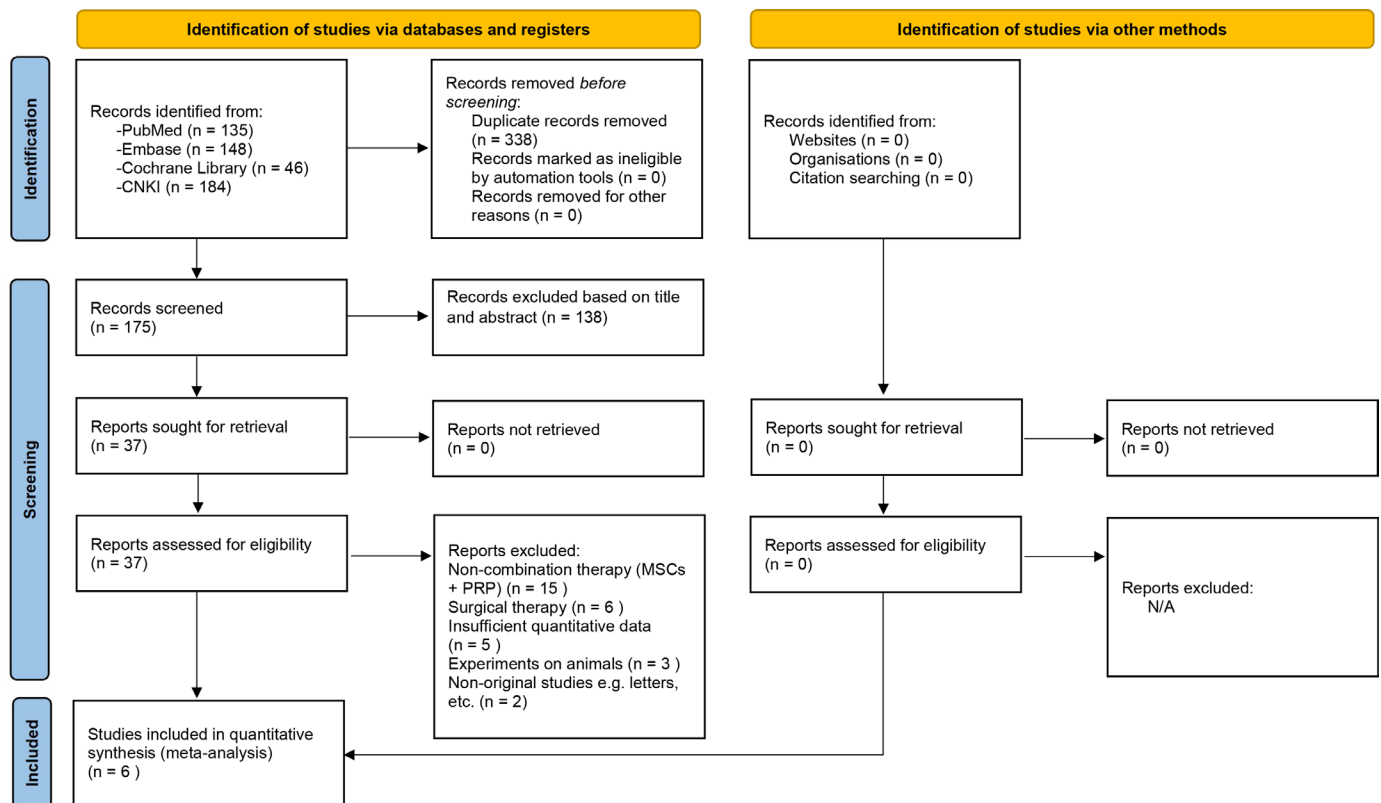


Figure 1 Flow chart of the included studies.

$p < 0.05$ was considered statistically significant. We used the Cochrane Q test to evaluate the heterogeneity among studies, which was judged according to the I^2 value. $I^2 < 50\%$ was considered to indicate low heterogeneity, and the data were analysed by a fixed-effect model; $I^2 \geq 50\%$ was considered to indicate high heterogeneity, and the data were corrected by a random-effect model and interpreted carefully. In addition, if more than 10 articles were included, we created a funnel chart to evaluate publication bias; if the number of included studies was less than 10, we did not analyse publication bias.

Patient and public involvement

Neither the patients nor the public substantially participated in any stage of this study.

RESULTS

Literature search results

A total of 513 studies were identified, including 135 in PubMed, 148 in Embase, 46 in the Cochrane Library and 184 in CNKI. After removing duplicate studies and reading the full text, six studies^{24–29} were finally included, including two from Portugal,^{24 26} three from China^{27–29} and one from Spain.²⁶ The document retrieval process and reasons for elimination are shown in figure 1. The six studies included in this study were clinical controlled trials, and the level of clinical evidence was 1–2. One study was a four-arm trial,²⁹ and the control groups were treated with MSCs, PRP or HA alone, without a blank group or

placebo among the control groups. A total of 493 cases were included, including 204 cases in the experimental group and 289 cases in the control group. Five studies were conducted with MSCs derived from bone marrow,^{24–28} and one study was conducted with stem cells derived from umbilical cord tissue.²⁹ All the included studies^{24–29} used MSCs combined with PRP in the experimental groups. The average age of the patients included in each study was higher than 54 years. One included study²⁷ did not report the OA stage of the included patients. The other five studies^{24–26 28 29} reported the OA stage of the included patients (using the Kellgren-Lawrence (K-L) staging method), and the stage ranged from 1 to 4. The follow-up time was 6–12 months. The basic characteristics of the studies included in this analysis are shown in table 1.

Study quality evaluation

In this study, the Cochrane ROB 2.0 for randomised controlled trials was used to evaluate the quality of the six included studies. Overall, five studies were evaluated as having low risk.^{24–27 29} In terms of the blinding method, five studies all described the random allocation method used in the study,^{24–27 29} and only one study²⁸ did not specify the specific method of randomization. In terms of data integrity, six studies^{24–29} were reported according to the research scheme, and the follow-up protocol and excluded cases were accurately explained. In terms of selective reporting, one article was not clear about the risk due to the lack of necessary information for selective

Table 1 Characteristics of the included studies

First author	Year	Country	Level of evidence	No of participants (M/F)		Intervention method in CG		Age, years (mean or mean±SD)		BMI, kg/m ² (mean or mean±SD)		Follow-up period, months	Stem cell source	K-L grade
				MSCs+PRP	CG	MSCs	CG	MSCs+PRP	CG	MSCs+PRP	CG			
Bastos ²⁴	2019	Portugal	II	14 (5/9)	16 (10/6)	MSCs		60.8±9.9	55.7±7.8	28.9±4.9	30.6±4.5	12	Bone marrow	1-4
Lamo-Espinosa ²⁵	2020	Spain	I	24 (17/7)	26 (16/10)	PRP		56	54.6	27	25.3	12	Bone marrow	2-4
Bastos ²⁶	2018	Portugal	II	9 (4/5)	9 (5/4)	MSCs		60.4±11.3	54.7±7.2	NR	NR	12	Bone marrow	2-4
Hou ²⁷	2016	China	I	92 (40/52)	88 (38/50)	HA		57±8.3	55±9.2	NR	NR	6	Bone marrow	NR
Cheng ²⁸	2019	China	I	20 (8/12)	20 (9/11)	HA		54.6±6.2	52.9±5.3	22.1±1.6	21.5±1.5	6	Bone marrow	1-3
Ha ²⁹	2018	China	I	45 (15/30)	44 (14/30) 43 (12/31) 43 (11/32)	PRP MSCs HA		56.8±6.1	55.6±3.6 57±3.2 56.2±6.7	25.5±2	25.4±0.8 25.8±3.3 24.9±1.6	12	Umbilical cord	1-3

BMI, body mass index; CG, control group; F, female; HA, hyaluronic acid; KL, Kellgren-Lawrence; M, male; MSCs, mesenchymal stem cells; NR, not reported; PRP, platelet-rich plasma.

**Figure 2** Risk of bias assessment.

reporting evaluation.²⁸ The above literature quality evaluation results show that the methodological quality of the literature included in this study is generally high, which is of high value for excluding clinical heterogeneity. The risk-of-bias results of each study are shown in figure 2.

Meta-analysis results

VAS score

VAS score three months after treatment

Three studies^{25 28 29} reported a comparison of VAS scores 3 months after treatment. There was no heterogeneity among the studies ($p=0.67$, $I^2=0\%$); thus, a fixed-effect model was used. The results of the meta-analysis showed that there was no significant difference in the VAS scores of patients with KOA 3 months after treatment with MSCs+PRP compared with the VAS scores of patients in the control group (WMD -0.30 , 95% CI -0.64 to 0.04). Similarly, compared with HA alone (WMD -0.57 , 95% CI -1.34 to 0.20 , $p=0.15$) or PRP alone (WMD -0.34 , 95% CI -0.72 to 0.03 , $p=0.07$), MSCs+PRP did not have an advantage in reducing the degree of pain (online supplemental figure 1).

VAS score 6 months after treatment

Three studies^{25 28 29} reported a comparison of VAS scores 6 months after treatment. The results of the meta-analysis showed that compared with the control, MSCs+PRP could reduce the VAS score of patients with KOA after 6 months (WMD -0.40 , 95% CI -0.72 to 0.07), and the difference was statistically significant ($p=0.02$). Compared with HA alone (WMD -0.99 , 95% CI -1.75 to -0.23 , $p=0.01$) or PRP alone (WMD -0.38 , 95% CI -0.73 to -0.03 , $p=0.03$), MSCs+PRP showed more advantages in reducing the VAS score of patients with KOA (online supplemental figure 2).

VAS score 12 months after treatment

The results of the meta-analysis showed that compared with the control (WMD -0.55 , 95% CI -0.87 to -0.22 , $p=0.0009$), HA alone (WMD -1.20 , 95% CI -2.28 to -0.13 , $p=0.03$) and PRP alone (WMD -0.54 , 95% CI -0.89 to -0.18 , $p=0.003$), MSCs+PRP could significantly reduce

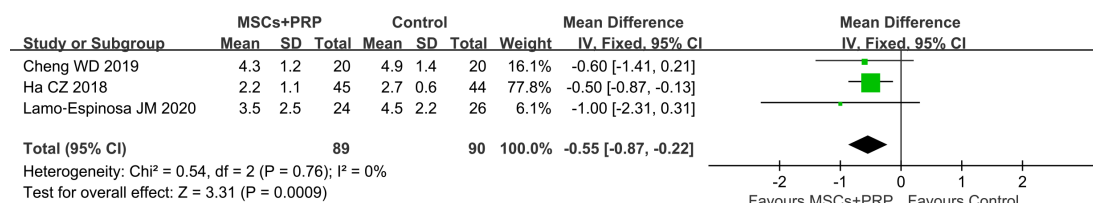


Figure 3 Forest plot of VAS score 12 months after treatment: MSCs+PRP versus control group (PRP and HA in control group). HA, hyaluronic acid; MSCs, mesenchymal stem cells; PRP, platelet-rich plasma; VAS, Visual Analogue Scale.

the VAS score of patients with KOA 12 months after treatment (figure 3 and online supplemental figure 3).

WOMAC score

WOMAC score 3 months after treatment

Meta-analysis showed that compared with the control (WMD -5.20, 95% CI -8.55 to -1.86, $p=0.002$) and HA alone (WMD -6.65, 95% CI -8.06 to -5.05, $p<0.00001$), MSCs+PRP could significantly reduce the total WOMAC score 3 months after treatment (online supplemental figure 4).

WOMAC score 6 months after treatment

The results of this study showed that MSCs+PRP could reduce the total WOMAC score 6 months after treatment compared with the control (WMD -7.65, 95% CI -12.38 to -2.92, $p=0.002$) or HA alone (WMD -9.11, 95% CI -13.43 to -4.80, $p<0.0001$) (online supplemental figure 5).

WOMAC score 12 months after treatment

The results of the meta-analysis showed that there was no significant difference between MSCs+PRP and the control in terms of the reduction of the total WOMAC score 12 months after treatment (WMD -4.63, 95% CI -15.22 to 5.95, $p=0.39$) (figure 4).

KOOS 12 months after treatment

The heterogeneity among the studies was low ($p=0.42$, $I^2=0\%$); thus, a fixed-effect model was adopted. Meta-analysis showed that there was no significant difference between MSCs+PRP and the control in terms of improvement of the KOOS 12 months after treatment (WMD 10.08, 95% CI -4.13 to 24.30, $p=0.16$) (online supplemental figure 6).

Adverse reactions

Only two studies^{26 29} reported the incidence of adverse reactions. The specific manifestations of adverse reactions were aggravation of knee joint pain, low back pain, low fever and swelling, and the above adverse reactions

were relatively mild. There was no heterogeneity among the studies ($p=0.38$, $I^2=0\%$); thus, a fixed-effect model was used. Meta-analysis showed that there was no significant difference in the incidence of adverse reactions between MSCs+PRP and the control (OR 0.59, 95% CI 0.25 to 1.36, $p=0.22$) (online supplemental figure 7).

DISCUSSION

Early KOA is generally treated with drugs, but drug treatment has limitations, including failure to repair damaged cartilage and serious gastrointestinal and renal adverse reactions.^{30 31} Therefore, it is of great significance to explore safer and more effective treatments or combination therapies for KOA. The results of this meta-analysis showed that compared with the control, HA and PRP alone, MSCs+PRP had more advantages in reducing the VAS score of patients with KOA at 6 and 12 months after treatment; MSCs+PRP also showed better clinical efficacy than the control and HA in improving the total WOMAC score of patients with KOA 3 months and 6 months after treatment. There was no significant difference in the incidence of adverse reactions between patients treated with MSCs alone and MSCs+PRP. It is worth noting that some statistically significant results were found in this meta-analysis, but the statistically significant differences do not necessarily represent significant differences in clinical efficacy; in addition, the number of randomised controlled trials included in this systematic review was small, which may lead to false-positive statistical results. Therefore, we suggest that users should consider the KOA stage and the tolerance of patients to treatment with PRP or MSCs in combination with our research conclusions when considering the best treatment strategy.

According to this meta-analysis, we believe that MSCs+PRP has potential advantages in improving pain and joint activity in patients with KOA. Although many studies have indicated that MSCs can repair joint surface

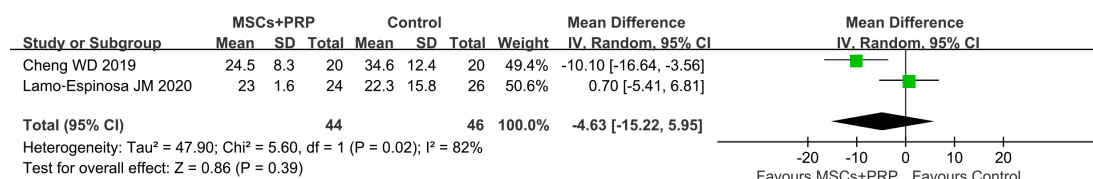


Figure 4 Forest plot of WOMAC score 12 months after treatment: MSCs+PRP versus control group (PRP and HA in control group). HA, hyaluronic acid; MSCs, mesenchymal stem cells; PRP, platelet-rich plasma; WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index.

damage, the function of MSCs depends more on the local microenvironment of the joint.³² The inflammatory and apoptotic environment formed by the loss of intra-articular dynamic balance will affect the proliferation and differentiation of MSCs.³³ The addition of PRP can improve the intra-articular microenvironment and promote the proliferation and differentiation of MSCs, which is also consistent with the conclusion that PRP can promote the proliferation and differentiation of MSCs confirmed by in vitro experiments.¹⁴ Combined with the conclusions of this study and previous basic studies,^{14 32 33} we believe that MSCs combined with PRP can improve the intra-articular microenvironment and promote the proliferation and differentiation of MSCs to repair damaged chondrocytes, inhibit the synovial inflammatory response, improve the intra-articular microenvironment and treat KOA. This may be the physiological mechanism through which MSCs+PRP improves VAS scores and total WOMAC scores in patients with KOA after treatment. In addition, due to the small number of included studies, we did not compare the efficacy of MSCs+PRP among different age, body mass index and sex subgroups. Previous studies^{34 35} of PRP for KOA have shown a shorter time of pain reappearance among patients with K-L stage 3 KOA than among patients with K-L stage I KOA and considered that age was negatively correlated with efficacy. Due to differences in age, sex and K-L stage, among other factors, different treatment schemes may show differences in efficacy in KOA patients. The study population should also be limited in future studies to clarify the therapeutic effect of MSCs+PRP.

The limitations of this study are as follows: (1) Although the quality of the included studies was generally high, the total number of studies and sample size were small; thus, the results still need to be confirmed by more studies in the future. (2) The follow-up time reported in the studies included in this meta-analysis was short, no more than 12 months, which is not conducive to deducing the long-term efficacy of MSCs combined with PRP for the treatment of KOA. (3) The outcome indicators of this study lack an objective test index and imaging indicators, which is not conducive to the multidimensional evaluation of the efficacy of MSCs combined with PRP for the treatment of KOA.

CONCLUSIONS

MSCs+PRP has a good clinical effect in improving pain and joint function in patients with KOA. Compared with MSCs alone, there was no significant difference in the incidence of adverse reactions with the clinical application of MSCs+PRP. Future research should focus on changes in the efficacy of MSCs+PRP over short-term, medium-term and long-term follow-up, which would be very valuable for clinicians in selecting treatment prescriptions. Due to the limited number of studies included, the conclusions of the above meta-analysis still need to be confirmed by larger multicentre, clinical, randomised controlled trials

with longer follow-up periods and evaluations of more objective physical, chemical and imaging indexes.

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Contributors JL and L-FZ conceived of the project, acting as guarantor, wrote the article and critically revised it for methodological and intellectual substance. JZ, GL and YH wrote the first draft of the manuscript. The search strategy, data extraction and literature search were all done by NX, WY, ML and JP who also critically revised the work for methodological and intellectual content. The final paper was approved by all authors.

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article or uploaded as online supplemental information. No data are available. The authors confirm that the data supporting the findings of this study are available within the article and its online supplemental materials.

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Supplementary material

Supplementary material 1 Retrieval strategy.

PubMed

#1 (((((Mesenchymal Stromal Cell*[Title/Abstract]) OR (Mesenchymal Stem Cell*[Title/Abstract])) OR (Mesenchymal Progenitor Cell*[Title/Abstract])) OR (Bone Marrow Stromal Cell*[Title/Abstract])) OR (Stem cell*[Title/Abstract])

#2 ((((((Osteoarthritis[MeSH Terms]) OR (Osteoarthritis[Title/Abstract])) OR (Osteoarthritis[Title/Abstract])) OR (Osteoarthritis[Title/Abstract])) OR (knee osteoarthritis[Title/Abstract])) OR (knee osteoarthritis[Title/Abstract])) OR (KOA[Title/Abstract])

#3 (((((((platelet-rich plasma[MeSH Terms]) OR (PRP[Title/Abstract])) OR (plasma rich in growth factors[Title/Abstract])) OR (platelet derived growth factor[Title/Abstract])) OR (platelet derived[Title/Abstract])) OR (platelet gel[Title/Abstract])) OR (platelet lysate[Title/Abstract])) OR (platelet rich plasma[Title/Abstract])

#4 #1 AND #2 AND #3

EMBASE

#1 'mesenchymal stromal cell*':ab,ti OR 'mesenchymal stem cell*':ab,ti OR 'mesenchymal progenitor cell*':ab,ti OR 'bone marrow stromal cell*':ab,ti

#2 osteoarthritis:ab,ti OR osteoarthritis:ab,ti OR osteoarthritis:ab,ti OR osteoarthritis:ab,ti OR 'knee osteoarthritis':ab,ti OR 'knee osteoarthritis':ab,ti OR koa:ab,ti

#3 'platelet-rich plasma':ab,ti OR prp:ab,ti OR 'plasma rich in growth factors':ab,ti OR 'platelet derived growth factor':ab,ti OR 'platelet derived':ab,ti OR 'platelet gel':ab,ti OR 'platelet lysate':ab,ti

#4 #1 AND #2 AND #3

Cochrane Library

#1 (knee osteoarthritis):ti,ab,kw OR (knee osteoarthritis):ti,ab,kw OR (osteoarthritis of Knee):ti,ab,kw OR (Osteoarthritis):ti,ab,kw OR (KOA):ti,ab,kw

#2 (platelet-rich plasma):ti,ab,kw OR (plasma rich in growth factors):ti,ab,kw OR (PRP):ti,ab,kw OR (platelet gel):ti,ab,kw OR (platelet lysate):ti,ab,kw

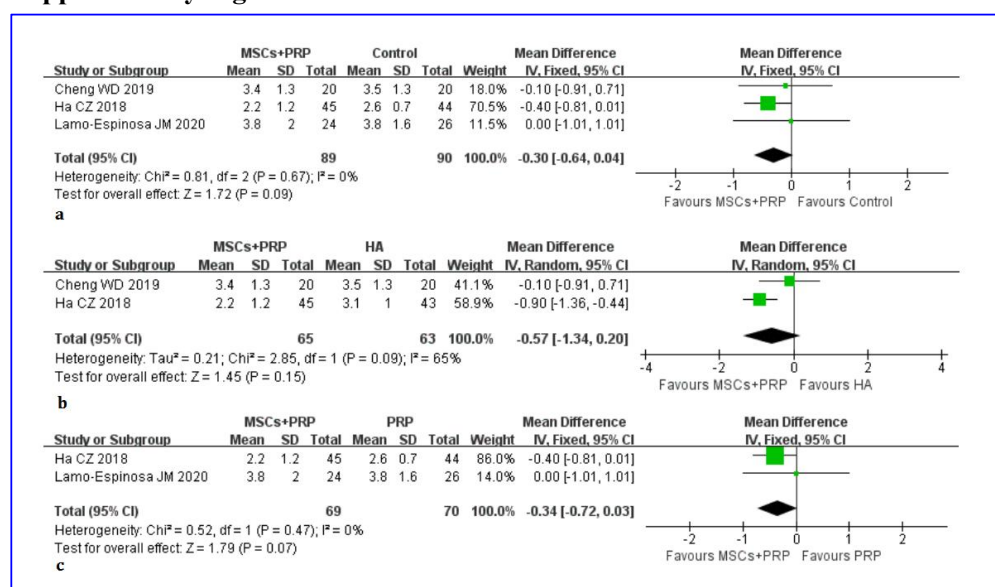
#3 (platelet-rich plasma):ti,ab,kw OR (plasma rich in growth factors):ti,ab,kw OR (PRP):ti,ab,kw OR (platelet gel):ti,ab,kw OR (platelet lysate):ti,ab,kw

#4 #1 AND #2 AND #3

CNKI

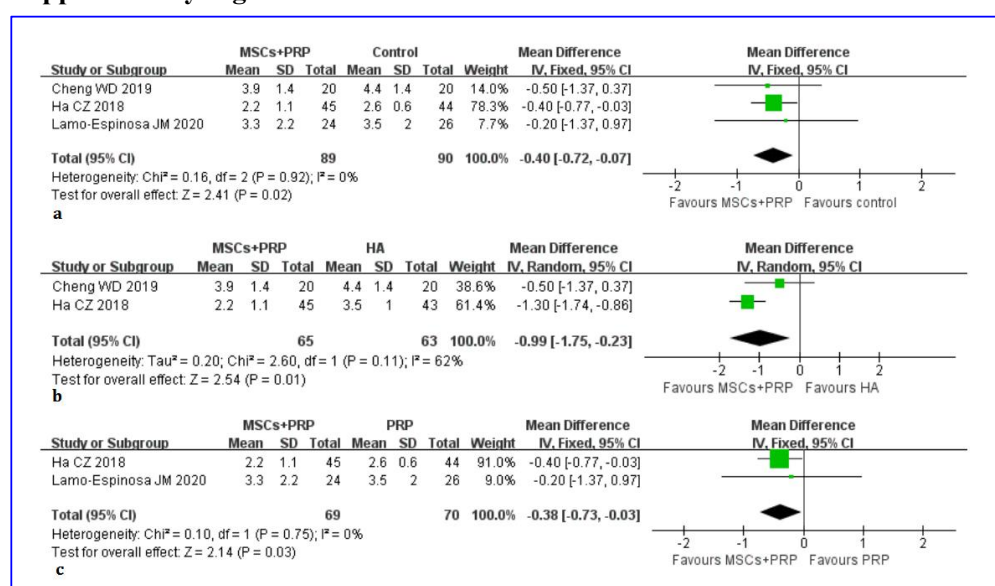
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Supplementary Figure 1



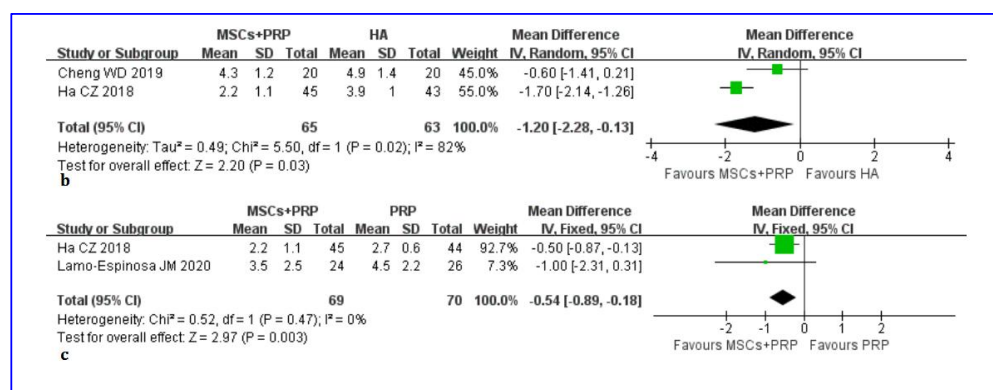
Supplementary Figure 1 Forest plot of VAS score 3 months after treatment : (a) MSCs+PRP vs control group (PRP and HA in control group); (b) MSCs+PRP vs HA; (c) MSCs+PRP vs PRP.

Supplementary Figure 2



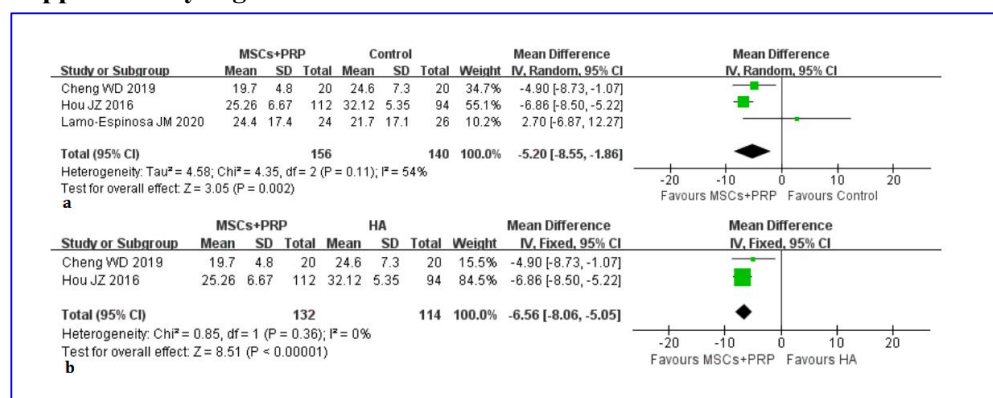
Supplementary Figure 2 Forest plot of VAS score 6 months after treatment : (a) MSCs+PRP vs control group (PRP and HA in control group); (b) MSCs+PRP vs HA; (c) MSCs+PRP vs PRP.

Supplementary Figure 3



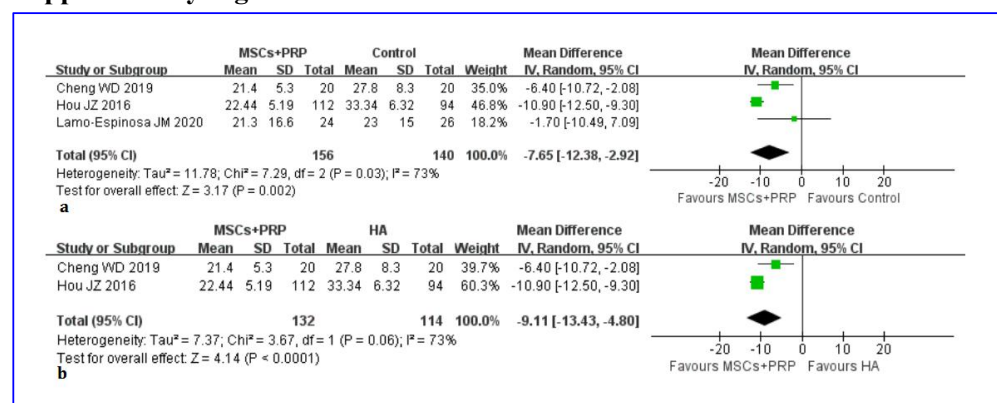
Supplementary Figure 3 Forest plot of VAS score 12 months after treatment : (b) MSCs+PRP vs HA; (c) MSCs+PRP vs PRP.

Supplementary Figure 4



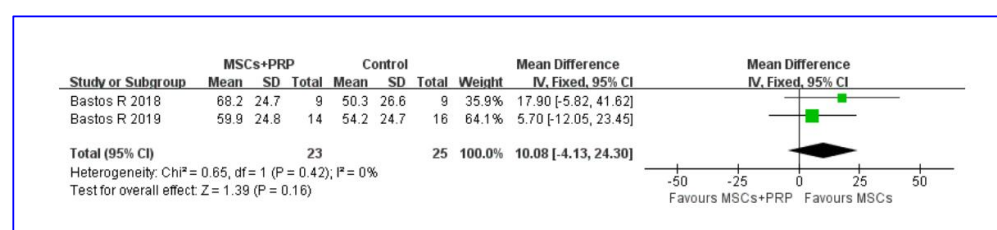
Supplementary Figure 4 Forest plot of WOMAC score 3 months after treatment : (a) MSCs+PRP vs control group (PRP and HA in control group); (b) MSCs+PRP vs HA.

Supplementary Figure 5



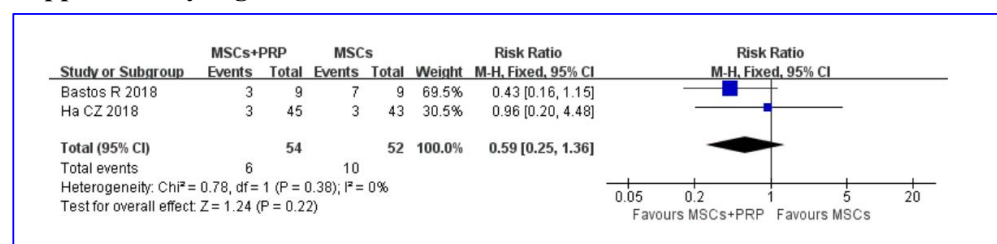
Supplementary Figure 5 Forest plot of WOMAC score 6 months after treatment : (a) MSCs+PRP vs control group (PRP and HA in control group); (b) MSCs+PRP vs HA.

Supplementary Figure 6



Supplementary Figure 6 Forest plot of KOOS 12 months after treatment : MSCs+PRP vs MSCs.

Supplementary Figure 7



Supplementary Figure 7 Forest plot of adverse reaction : MSCs+PRP vs MSCs.