Human umbilical cord blood-derived MSCs trans-differentiate into endometrial cells and regulate Th17/Treg balance through NF-κB signaling in rabbit intrauterine adhesions endometrium

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Abstract

Purpose: The fundamental cause of intrauterine adhesions (IUAs) is the destruction and reduction in stem cells in endometrial basal layer, resulting in endometrial reconstruction very difficult. The purpose of this study was to investigate the effects and underlying mechanism of human umbilical cord blood-derived mesenchymal stem cells (hUCB-MSCs) on the endometrial reconstruction after transplantation.

Methods: hUCB-MSCs were isolated and identified by flow cytometry, osteogenic, adipogenic and chondrogenic differentiation assays. The rabbit IUA models were established and set five groups (control, 14/28th day after surgery, estrogen and hUCB-MSCs treatment). The number of endometrial glands and the fibrosis rate were evaluated using HE and Masson staining, respectively. Endometrial proliferation, angiogenesis and inflammation were evaluated by immunohistochemical staining of ER, Ki-67 and TGF-β1, respectively. Single-cell RNA sequencing (scRNA-seq) was applied to explore the cell differentiation trajectory after hUCB-MSCs transplanted into IUA endometrium. Finally, molecular mechanism of hUCB-MSCs repairing damaged endometrium was investigated by RNA sequencing, qRT-PCR and Western blot assays.

Results: After transplantation of the hUCB-MSCs, the increase in endometrial gland number, estrogen receptor (ER) and Ki-67 expression, and the decrease in fibrosis rate and TGF-β expression (P < 0.05), suggested the endometrial repair, angiogenesis and inflammatory suppression. The therapeutic effect of hUCB-MSCs was significantly improved compared with 28th day after surgery and estrogen group. ScRNA-seq demonstrated that the transplanted hUCB-MSCs can trans-differentiate into endometrial cells: epithelial, fibroblast and macrophage. RNA sequencing of six IUA samples combined with qRT-PCR and Western blot assays further revealed that hUCB-MSCs may regulate Th17/Treg balance through NF-κB signaling, thus inhibiting the immune response of damaged endometrium.

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In women of reproductive age, severe injuries to the endometrium are often accompanied by intrauterine adhesion (IUA), which can result in menstrual disorders, dysmenorrhea, amenorrhea, recurrent spontaneous abortion and infertility [1, 2]. Currently, the gold standard treatment for IUA is hysteroscopic resection of the adhesions [3], followed by estrogen therapy to stimulate regeneration of the endometrium [4]. This treatment strategy is useful for patients with mild-or-moderate disease severity but not effective for severe IUAs [5]. The recurrence rate of severe IUA is up to 62.5% [1, 3], and the postoperative pregnancy rate is only 33.3% [6]. Current treatments can only temporarily restore the anatomy of the uterus, but cannot restore the structure and function of the endometrium. Due to impaired revascularization and ischemia in severe IUA [7], the clinical effects of these medications that promote blood perfusion such as aspirin and granulocyte colony-stimulating factor are limited [8, 9].

The cellular and molecular pathogenesis of IUA remains obscure and controversial. Some studies suggested that fibroblasts are activated by CTGF (connective tissue growth factor) through TGF-β after endometrial injury and then produce excessive fibrillar collagen, which inhibit the normal regeneration of endometrial mesenchymal stem cells [10, 11]. Studies also found that the expression of NF-κB in IUA endometrium was significantly higher than that in normal endometrium, which can promote the occurrence of endometrial inflammation [12]. Besides, damaged endometrial cells can promote Th2 immune response to promote fibrosis [13]. Thus, these cellular changes and molecular pathways may provide potential clues for new treatment of IUA to reverse inflammation and promote endometrial regeneration.

Although many methods have been used to treat severe IUA, the high recurrence rate and endometrial thinning limit the therapeutic effect. Therefore, how to regenerate the damaged endometrium and restore its function is an important problem and great challenge for IUA treatment. Recently, cell therapy for IUA is promising in the treatment of endometrial dysfunction [14]. Mesenchymal stem cells (MSCs) are considered to be an ideal cell type for tissue regeneration due to their high potential differentiation, self-renewal and immune regulation [15]. At present, the animal and clinical trials of MSCs in IUA treatment have been abundantly carried out, showing the ability of obvious endometrial repair and regeneration for MSCs [16]. Although MSCs have demonstrated to be a promising cell therapy in IUA treatment, these studies are still in the experimental research stage, and its advantages over estrogen therapy and the exact mechanism of treatment are worthy of further investigation.

Cord blood contains both hematopoietic and non-hematopoietic cells, of which mesenchymal stem cells are a very small percentage of non-hematopoietic cells. They can be considered as “very young” and have the potential for multilineage differentiation [17]. Human umbilical cord blood-derived MSCs (hUCB-MSCs) have been regarded as a favorable source for cell-based therapies because of their easy collection, low immunogenicity, and high proliferative potential. hUCB-MSCs have been used in the treatment of type 1 diabetes [18], osteoarthritis [19], acute graft-versus-host disease [20], etc. However, few studies have reported the application of hUCB-MSCs in endometrial repair.

In the present study, we used hUCB-MSCs to treat rabbit IUA model and explored the underlying therapeutic mechanisms. We used scRNA-seq to determine whether hUCB-MSCs transplanted into endometrium differentiate into endometrial cells and used RNA sequencing to investigate the role of immunoregulation and signaling pathway during hUCB-MSCs transplantation.

Conclusions: Our study demonstrated that hUCB-MSCs can repair damaged endometrium through trans-differentiation, immunomodulatory capacities and NF-κB signaling, suggesting the treatment value of hUCB-MSCs in IUA.

Keywords: Intrauterine adhesion (IUA), Human umbilical cord blood-derived mesenchymal stem cells (hUCB-MSCs), scRNA-seq, Trans-differentiation, NF-κB signaling
Isolation of hUCB-MSCs
Umbilical cord blood samples (about 50 mL each) with anticoagulant (EDTA) were collected from umbilical cord vein. Mononuclear cells (MNCs) were isolated by density gradient centrifugation for 30 min at 400 g and washed 3 times in PBS (Beyotime). The isolated MNCs were plated in 75 cm² cell culture flask containing mesenchymal stem cell basal medium (Yocon, China) and incubated at 37 °C in a humidified atmosphere of 5% CO₂. Around the tenth day of culture, we can see adherent cells, while non-adherent cells were removed, and then the medium was changed every 3 days. After expansion to 80–90% confluence, the cells were harvested by 0.25% trypsinization and subcultured for further experiments. Inverted microscope was used to analyze the morphology of hUCB-MSCs. According to the International Society for Cellular Therapy, MSCs have three characteristics: (1) plastic adherent, (2) express CD105, CD73 and CD90 and not express CD45, CD34, CD14, CD11b, CD79α, CD19 and HLA-DR surface antigen, (3) differentiate into osteoblasts, adipocytes and chondroblasts in vitro [21]. Thus, hUCB-MSCs were analyzed by flow cytometry, osteogenic, adipogenic and chondrogenic differentiation assays.

Flow cytometric analysis
Detached hUCB-MSCs were washed twice with ice-cold PBS, centrifuged and fixed in 4% paraformaldehyde. Then, cells were incubated with mouse anti-human CD45-FITC, CD34-FITC, CD11b-FITC, CD19-FITC, CD29-FITC, CD73-PE, CD105-APC-A750, CD90-APC and HLA-DR-FITC (all from BioLegend) in dark for 20 min. Cells were analyzed by flow cytometer (Beckman Coulter GmbH, Krefeld, Germany).

Osteogenic, adipogenic and chondrogenic differentiation
To investigate the osteogenic and adipogenic differentiation potential of hUCB-MSCs, third-passage cells were plated at a concentration of 3 x 10^5 cells/cm² and cultured with osteogenic and adipogenic medium (Sigma-Aldrich) for 3 weeks with medium changes twice weekly, respectively. At the end of differentiation, cells were stained with Alizarin Red S and Oil Red O, respectively. To induce chondrogenic differentiation, we cultured 3 x 10^5 hUCB-MSCs/well in chondrogenic medium (Procell) for 4 weeks. Medium changes were carried out twice weekly, and chondrogenesis was assessed at 2–3-day intervals. Cells were fixed in 4% formaldehyde, dehydrated in an ethanol series and embedded in paraffin blocks. Blocks were cut, and sections were stained with Alcian Blue to evaluate chondrogenic differentiation. Osteogenic medium consists of IMDM supplemented with 0.1 μM dexamethasone, 10 mM β-glycerol phosphate and 0.2 mM ascorbic acid. Adipogenic medium consists of IMDM supplemented with 0.5 mM 3-isobutyl-1-methylxanthine, 1μM hydrocortisone, 0.1 mM indomethacin and 10% rabbit serum. Chondrogenic medium consists of DMEM supplemented with 6 μg/mL insulin–transferrin–selenium premix, 0.1 mM ascorbic acid 2-phosphate, 10 mM sodium pyruvate, 10 ng/mL transforming growth factor-β1 and 100 nM dexamethasone.

Treatment for IUA model
The rabbits (n = 6) were randomly assigned to two groups, including estrogen treatment group (n = 3, rabbits underwent surgery of intrauterine adhesions, estradiol benzoate (0.5 mg/kg) was administered intramuscularly every 4 days) and hUCB-MSCs treatment group (n = 3, rabbits underwent surgery of intrauterine adhesions, 1 weeks after surgery, a relaparotomy was performed and the rabbits were injected with 1 x 10^6 hUCB-MSCs in each of the uterine wall, Fig. 1D). After 28 days of treatments, all rabbits were killed for the collection of uterine tissue.

Histological analysis
The endometrial gland number and fibrosis rate were examined via hematoxylin–eosin (HE) and Masson staining, respectively. The excised uteri were fixed in 4% paraformaldehyde, then embedded in paraffin, sliced
into 5-mm-thick sections, and routinely stained with HE and Masson stains according to standard protocols. Sections were examined under an inverted microscope (Leica, German, DMIL-PH1). Four high-power fields (HPF) were selected on each HE-stained slice to count the number of glands. Four high-power fields (HPF) were selected on each Masson-stained slice, and the fibrosis rate was calculated using ImageJ software (National Institutes of Health) as follows: the area of endometrial stromal fibrosis/the area of endometrial stroma and glands (excluding the uterine cavity).

**Immunohistochemistry**

In order to evaluate expressions of endometrial receptivity-related estrogen receptor (ER), angiogenesis related Ki-67, and inflammation-related transforming growth factor-β1 (TGF-β1), immunohistochemistry was used. The transverse paraffined uterine sections were deparaffinized, rehydrated, and then incubated in 5% bovine serum albumin (Beyotime) for 30 min at 37 °C to block the nonspecific antibody. Sections of the uterus were deparaffinized and gradually dehydrated. Slides were incubated with rabbit anti-Ki-67 (1:200 dilution), anti-ER

Fig. 1 Establishment of IUA model and characterization of hUCB-MSCs. A Normal uterus after laparotomy without treatment. B Uterus after IUA surgery. An LPS surgical suture placed in the uterine cavity after curettage. C The abdomen after suture. D Schematic diagram of hUCB-MSCs injected into rabbit uterus wall. E Representative image of hUCB-MSCs. Scale bar: 100 μm. F, G Representative image of hUCB-MSCs adipogenic (F) and osteogenic (G) differentiation. Scale bar: 200 μm. H Representative image of hUCB-MSCs chondrogenic differentiation. Scale bar: 100 μm. I-L Surface antigens of hUCB-MSCs detected by flow cytometry assay. Cells were positive for CD73, CD90 and CD105, but negative for CD34, CD45, CD11b, CD19 and HLA-DR.
ordered into a trajectory with branch points. As called in space. Single cells are projected onto this space and dimensional expression profiles to a low-dimensional embedding, a machine learning technique to learn a par-

between clusters. Then, monocle uses reversed graph ordering genes which showed differential expression.

Monocle, cells in the same segment of the trajectory have the same ‘state.’

RNA expression profiling by RNA sequencing

The rabbit IUA samples with and without treated with hUCB-MSCs were used for transcriptome sequencing (3 vs 3). Firstly, total RNAs were extracted by Trizol method, RNA purity was detected by spectrophotometer, and RNA integrity was analyzed by agarose gel electrophoresis and Agilent 2100 BioAnalyzer. The Library was constructed using Illumina’s NEBNext® UltraTM RNA Library Prep Kit. Then, Illumina platform was used for library sequencing and 150 bp paired terminal reading was generated to obtain the sequence information of the fragment to be measured. After quality control and sequence alignment based on reference genome, DESeq2 software [24] was used to analyze the differentially expressed genes (DEGs) between the two groups. Finally, the DEGs were used for gene enrichment analysis based on gene ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG).

Western blot

Proteins were extracted from endometrium samples with RIPA buffer containing proteinase inhibitors. The protein concentrations were quantified using BCA Protein Assay Kit (Beyotime, Shanghai, China) and separated in SDS-PAGE gel (Solarbio, Beijing), and then they were transferred onto the PVDF membrane. The membranes were incubated with anti-NF-κB-p65 and anti-GADPH for 1 h at room temperature. The blots were visualized using ECL chemiluminescence kit (enhanced), and the band intensity was quantified with ImageJ software.

qRT-PCR

Total RNA was extracted from excised endometrium using trizol reagent (Invitrogen, USA). cDNA was generated by reverse transcription of per RNA sample using PrimeScript RT reagent kits (Takara, Japan) following the manufacturer’s guidelines. The PCR reaction system was performed with specific primers and SYBR Premix Ex Taq II (Takara, Japan), with a final volume of 20 μL. The PCR cycling conditions were as follows: pre-denaturation at 95 °C for 30 s, denaturation at 95 °C for 5 s, annealing at 60 °C for 30 s, 40 cycles, and elongation at 60 °C for 30 s. Analyses of relative gene expressions were performed using 2^−ΔΔCT methods. GAPDH was used as an internal control. Primer sequences are summarized in Table 1.
Statistical analysis

Statistical analysis was performed with SPSS 20.0 software. Numerical data were indicated as means standard deviation. For nonparametric statistics, data were analyzed using the Mann–Whitney U test and presented as populations with median values indicated by bars. For parametric statistics, data were analyzed using unpaired Student’s t test. Data were presented as a mean value with 95% confidence interval (CI). P values < 0.05 were considered to be statistically significant.

Results

Identification and characterization of hUCB-MSCs

We observed that hUCB-MSCs were adherent, with a fibroblast-like shape under optical microscope (Fig. 1E). Lipid droplets can be observed by Oil Red O staining (Fig. 1F); calcified extracellular matrix was detected by Alizarin Red S staining, (Fig. 1G); blue-stained acid proteoglycan was observed by Alcian Blue staining (Fig. 1H), demonstrating that hUCB-MSCs could differentiate into adipocytes, osteoblasts and chondroblasts. The results of flow cytometry showed that the cells were positive for CD73, CD90 and CD105, but negative for CD34, CD45, CD11b, CD19 and HLA-DR (Fig. 1I–L), indicating that these cells were MSCs, not hematopoietic stem cells or macrophages. These findings are concordant with the characteristics of mesenchymal stem cells.

The gland number and fibrosis rate of endometrium after hUCB-MSCs transplantation

We found that normal endometrial surface was covered with simple high columnar epithelial cells, and endometrial glands were primarily located in submucosa and basal layer. On 14/28th day after surgery, the endometrium showed flat low columnar epithelial cells (Fig. 2A). Compared with control group, the gland number markedly reduced, and the fibrosis rate was obviously elevated in 14/28th day after surgery ($P<0.05$, Fig. 2A–D). The situation did not improve on 28th day after surgery compared with 14th day, which indicated that the rabbit IUA models were successfully established.

After the hUCB-MSCs transplantation, the gland number statistically increased and the fibrosis rate decreased compared with 28th day after surgery ($P<0.05$, Fig. 2A–D). Though the tendency of increased gland number and decreased fibrosis rate can be observed, there was still no significance on the two indexes between the estrogen treatment group and 28th day after surgery ($P>0.05$, Fig. 2A–D). Compared with estrogen treatment group, the two indexes in hUCB-MSCs transplantation group were closer to control group, which suggested that hUCB-MSCs could be superior to estrogen in IUA treatment.

Expression of ER, Ki-67 and TGF-β1 of endometrium after hUCB-MSCs transplantation

To further explore the hUCB-MSCs contributor on IUA endometrium, we detected the ER, Ki-67 and TGF-β1 expression in endometrial issues (Fig. 3A, C, E), which reflect the hormone level, the ability of angiogenesis and the degree of fibrosis of uterine tissue, respectively. After the hUCB-MSCs transplantation, there is an obvious increase in ER and Ki-67 expression and obvious reduction in TGF-β1 expression compared to 28th day after surgery ($P<0.05$, Fig. 3B, D, F). Notably, compared with 28th day after surgery, the increase in ER expression and reduction in TGF-β1 expression were observed in estrogen treatment group ($P<0.05$, Fig. 3B, F), but no significance in Ki-67 expression between the two groups ($P>0.05$, Fig. 3D). Furthermore, compared with estrogen treatment group, the expression of the three protein in hUCB-MSCs transplantation group was more similar to control group. These pieces of evidence proved that hUCB-MSCs have advantages over estrogen for IUA treatment.

ScRNA-seq revealed hUCB-MSCs trans-differentiate into endometrial cells

To investigate the cellular architecture and cell differentiation trajectory of IUA from single cell level, a rabbit IUA sample with hUCB-MSCs transplantation was utilized to scRNA-seq and bioinformatics analysis (Fig. 4A). After quality control, 4097 human-derived cells were used to perform t-SNE analysis and 8 cell clusters were obtained (Fig. 4B). By analyzing the differential expressed genes across the clusters and combining with classical markers of human cell types from a previous study [25], the relationship of the clusters and cell types was displayed (Fig. 4C, D). After purification and re-annotation manually, four main cell types including MSCs (68 cells, 1.66%), fibroblasts (664 cells, 16.21%), epithelial cells (2770 cells, 67.61%) and macrophages (595 cells, 14.52%)

| Table 1. Primers of specific genes used in qRT-PCR analyses |
|------------------------|------------------------|
| Gene        | Sequence (5’−3’)             |
| NFKB1 Forward | CCTGAGTCTTTTGACCGCT    |
| Reverse     | GCAGGCTATGGCTCAACCG    |
| GAPDH Forward | CTTGATCGTGGAGGAAGGA |
| Reverse     | AGGGATGATGTTCGAGAG   |
were revealed (Fig. 4E). The MSCs annotated were the residual transplanted hUCB-MSCs, which did not take long enough to transform into any other cell types, while the remaining three cell types (fibroblasts, epithelial cells and macrophages) could be the result of endometrial reconstruction after hUCB-MSCs transplantation for IUA treatment.

Since MSCs displayed stem cell-like signatures and have potential to differentiate into other cell types, we speculated that pseudotime series analysis based on the four cell types may capture the main differentiation...
Fig. 3  Immunohistochemistry of ER, Ki-67 and TGF-β1 (A, C, E). Micrographs of ER (A), Ki-67 (C) and TGF-β1 (E) expression from 5 groups, i.e., rabbits without surgery (control), 14/28th day after surgery, and treated with estrogen/hUCB-MSCs. (B, D, F). The expression of ER (B), Ki-67 (D) and TGF-β1 (F) in endometrium in 5 groups. All original magnification at ×400. Each group has three independent experiments.
processes of MSCs during treatment of IUA using hUCB-MSCs. A cell trajectory was reconstructed by Monocle, which mainly contained three branches (Fig. 5A). Notably, the trajectory’s root was MSCs, which trans-differentiate into epithelial cells throughout the whole process. And some MSCs trans-differentiate into fibroblasts firstly, while some followed to trans-differentiate into macrophages (Fig. 5B). The results suggested that hUCB-MSCs can trans-differentiate into endometrial tissues, which will benefit for the mechanism understanding and clinical treatment of MSCs transplantation in uterine inflammatory injury.

Similarly, 8792 rabbit-derived cells were annotated to four main cell types: endothelial cells (2431 cells, 27.65%), fibroblasts (5721 cells, 65.07%), epithelial cells (208 cells, 2.37%) and macrophages (312 cells, 3.55%) and three rare cell types (Fig. 5C). Integrated human with rabbit-derived cells, we observed that the proportion of endothelial cells (2431 cells, 18.7%) and fibroblasts (6385 cells, 49.54%) was elevated, while the proportion of epithelial cells (2978, 23.1%) and macrophages (907 cells, 7.04%) was decreased (Fig. 5D). By analyzing the percentage of normal uterine tissue cells in a previous study [25], we found that the distribution of merged cell types was more similar to that of normal endometrium compared with that of rabbit-derived cell types (Fig. 5D). This indicates that MSCs can differentiate endometrial tissue cells and restore the damaged endometrial environment to normal, which reflects the value of mesenchymal stem cell transplantation in the treatment of IUA. With the prolongation of action time after MSC transplantation and the initiation of body self-repair, the proportion of cells in the damaged endometrial tissue may tend to be more and more normal.

RNA Sequencing revealed hUCB-MSCs regulate Th17/Treg balance and NF-κB signaling in treating IUA

To clarify the biological role of treatment with hUCB-MSCs in IUA, RNA sequencing of 6 samples (3 for 28th day after surgery and 3 for hUCB-MSCs treatment) was performed to observe the molecular expression changes in IUA samples before and after hUCB-MSCs transplantation. We found that hUCB-MSCs treatment caused 2188 genes to be up-regulated and 1193 genes to be down-regulated (Fig. 6A, B). GO and KEGG pathway enrichment analysis was performed based on these differential expressed genes. GO result showed that treatment-associated genes were mainly enriched in cell proliferation and cell migration (Fig. 6C). KEGG analysis depicted that hUCB-MSCs treatment may participate in the pathways related to immune cells (Th1, Th2 and Th17) differentiation, T cell receptor signaling and NF-κB signaling pathway (Fig. 6D).

NF-κB signaling has been reported to be possible involved in MSCs transplantation to treat IUA [11], thus becoming to our focus. NFKB1 mRNA expression was significantly decreased after hUCB-MSCs treatment in RNA-Seq experiment (Fig. 6E), which was validated by qRT-PCR assay (Fig. 6F). WB assay also confirmed that NF-κB-p65 protein level indeed reduced compared with pre-treatment (28th day after surgery) (Fig. 6G, H), indicating that NF-κB signaling may play a crucial role in hUCB-MSCs regulating inflammatory microenvironment of endometrium. According to previous studies [25, 26], MSCs can regulate the polarization of naive T cells into anti-inflammatory regulatory T cells (Treg) and reduce the ratio of Th17/Treg cells to exert immunomodulatory function, which can mediate through the activation of NF-κB signaling pathway. Thus, we believed that hUCB-MSCs can regulate Th17/Treg balance and NF-κB signaling in treating IUA, which promoting the mechanism understanding and potential clinical application for treating IUA with hUCB-MSCs transplantation.

Discussion

The components of endometrial tissue include epithelial cells, stromal cells, vascular smooth cells and vascular endothelial cells. The healthy endometrium with regenerative capacity can divide into two zones: the upper functional layer and lower basal layer. During a menstrual cycle, the functional layer shed, while the permanent basal layer can regenerate into new functional layer every menses as well as postpartum [27–29]. It has become evident that endometrial stem cells are responsible for endometrial regeneration [30], while their number of endometrial stem cells decreases and their function was weaken in IUAs due to the damaged basal layers [1]. Exogenous MSCs transplantation may compensate for this endogenous reduction to exert the role of regeneration.

MSCs have effects on tissue repair by homing to the injured site, secreting chemokines, modulating the immune function, differentiating into other types of cells and potentially having antimicrobial ability. They
Fig. 4 (See legend on previous page.)
can suppress inflammation by secreting immunomodulatory factors, such as, IL-6, IL-8, MCP-1, CCL5 and TLR-4, thereby decreasing proliferation and activation of CD4+ T cells [31, 32]. Clinical trials have demonstrated the initial safety and efficacy of mesenchymal stem cells derived from bone marrow, umbilical cord, menstrual blood and adipose tissue in restoring menstruation, fertility outcomes and endometrial regeneration. Studies have showed transplantation with the MSCs promoted endometrial regeneration and collagen resurgery and enhanced the expression of estrogen receptor of IUA model [32]. Animal experiments have corroborated that MSCs transplanted into uterine cavity of mice can exert immunomodulatory effects by secreting anti-inflammatory cytokines IL-6 [33]. Current knowledge on IUA pathogenesis was mostly derived from tissue studies without considering the multicellular structures and their orchestration; thus

Fig. 5 Cell differentiation trajectory of hUCB-MSCs and the cell type proportion of rabbit-derived cells. A The trajectory with 3 branches identified by pseudotime analysis. B The specific cell distribution of four cell types in differentiation trajectory. C Four mainly cell types and three rare cell types identified in rabbit-derived cells. D The proportion of four mainly cell types in IUA rabbit-derived cells, merged (human and rabbit) endometrial cells and normal human endometrial cells from a reference literature.
it is quite essential for in-depth mechanistic investigations, such as trans-differentiation, immunoregulation and signaling pathway.

Previous studies showed MSC-induced functional improvement in injured tissues by paracrine effect rather than direct differentiation [34, 35]. However, our study revealed the potential of differentiation for MSCs in tissue repair. The annotation analysis of human-derived cell types from scRNA-seq showed that hUCB-MSCs may differentiate fibroblasts, epithelial cells and macrophages (Fig. 4E), suggesting the potential of endometrial repair and regeneration after hUCB-MSCs transplantation in IUA treatment. (1) Previous studies demonstrated that MSCs transplantation can affect the composition and morphology of endometrial epithelial cells, and promote the regeneration of endometrial epithelial cells [36, 37]; (2) MSCs were considered as one of the sources for fibroblasts and even immature fibroblasts, which could regenerate during the repair of inflammatory damage [38]; (3) MSCs were reported to promote macrophage polarization inhibit inflammatory progression, and increased anti-inflammatory responses in vivo and in vitro [39]. These pieces of evidence also provide clues of the differentiation possibility into the three cell types during IUA.
treatment after MSCs transplantation. In our study, the pseudotime series analysis also showed MSCs can transdifferentiate into epithelial cells, fibroblasts and macrophages throughout the differentiation process or in different periods (Fig. 5A, B). The results suggested that scRNA-seq could reveal the cellular architecture and evolution of IUA from single cell level.

Excitingly, compared with rabbit-derived cell types, we found that the distribution of merged cell types was more similar to that of normal endometrium (Fig. 5D). The merged cell types represent IUA endometrium after hUCB-MSCs transplantation, while the rabbit-derived cell types roughly represent IUA endometrium without treatment. This indicates that MSCs can differentiate endometrial tissue cells and restore the damaged endometrial environment to normal. Of course, the proportion of merged cell types was still slightly different with normal endometrium. This could be due to the influence of distinct sample sources and processing methods, as well as the diversity of cell type annotations. On the other hand, we speculated that the proportion of cells in the damaged endometrial tissue may tend to be completely normal as MSCs continue to function as well as body self-repair.

MSCs can regulate the polarization of naïve T cells into anti-inflammatory regulatory T cells (Treg) and reduce the ratio of Th17/Treg cells to exert immunomodulatory function [25, 40, 41]. The immunomodulatory effect of MSCs is proved to mainly communicate with T cell through IL-6-mediated paracrine way [42]. The interaction between lymphocytes and MSCs also depends on the induction of IL-6 expression by MSCs [43]. IL-6 has been demonstrated to regulate Th17/Treg balance [44], and inhibiting its expression could significantly impair the immunomodulatory function of MSCs [45]. On the other hand, MSC transplantation can mediate T cell response through the activation of NF-κB signaling pathway [26], which may rely on LAT to exhibit T cell regulatory function [46, 47]. The inhibition of LAT reduced CTLA-4 and CD25 expression in Treg cells and impairs their immunosuppression capacity [48]. These pieces of evidence indicate the key role of cellular molecules and signaling pathways in inflammation and immune regulation after MSCs transplantation [49]. In our study, functional enrichment analysis based on differential expressed genes showed that MSCs transplantation may treat IUA through T cell differentiation and NF-kappa B signaling pathway. In order to determine the immunomodulation function, the supernatants of third-generation hUCB-MSCs were collected and Legend plex TM human Th cytokine assay (BioLegend) was used to analyze the following 13 factors: IL-2, IL-4, IL-5, IL-6, IL-9, IL-10, IL-13, IL-17A, IL-17F, IL-21, IL-22, IFN-α and TNF-γ. Experiments were repeated to ensure the accuracy of the data. Finally, CytoFLEX flow cytometer (Beckman Coulter GmbH, Krefeld, Germany) was used for analysis, and the data were analyzed by LEGENDplex V8.0 and Prism7.0 software (BioLegend), the concentration was pg/ml. The results showed that the concentrations of IL-5, IL-13, TNF-α and IFN-γ were lower, the concentrations of IL-2, IL-9, IL-10, IL-17A, IL-17F, IL-4, IL-21 and IL-22 were undetectable, but the secretion level of IL-6 (891.58 pg/ml) was very high (Additional file 1: Figure S1-2). The expression of IL-6-related genes (IL6R and IL6ST) and LAT detected by RNA sequencing were significantly elevated compared with pre-treatment (Additional file 1: Figure S3). Therefore, we speculate that except for NFKB1, IL-6 and LAT may also be the key factors in the action mechanism of MSCs transplantation in the treatment of IUA, which needs to be further investigation in future studies.

Conclusion
Transplantation of hUCB-MSCs promoted structural reconstruction and regulated inflammatory environment in IUA endometrium through NF-kB signaling. We firstly demonstrated that hUCB-MSCs can differentiate into normal endometrial tissue in rabbit. It was proved that hUCB-MSCs may be a promising therapy for humans with IUA. Of course, more research is needed on the long-term safety profile and effectiveness for patients with IUA.

Abbreviations
hUCB-MSCs: Human umbilical cord blood-derived mesenchymal stem cells; ER: Estrogen receptor; H&E: Hematoxylin and eosin; WB: Western blot; IUA: Intrauterine adhesion; TGF: Transforming growth factor; IL: Interleukin; MCP: Monocyte chemoattractant protein; CCL: Chemokine ligand; TLR: Toll-like receptor; LAT: Linker for activation of T cells.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s13287-022-02990-1.

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Author contributions
YLW and LS were responsible for experimental design and project conception. QH and YZ carried out the main experiments, performed statistical analysis and wrote the manuscript. HRL, HJL, LL, RRJ, YXQ, MT and JWW participated in sample collection and manuscript revision. All the authors read and approved the final version.

Additional file 1: The concentration and expression analysis of cytokines in Discussion.
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Availability of data and materials
The RNA-seq and scRNA-seq datasets can be available in GEO with accession number GSE205997. The original images and statistic tables were attached in Additional file 1.

Declarations

Ethics approval and consent to participate
The animal study was reviewed and approved by Ethics Committee of Zhengzhou Central Hospital Affiliated to Zhengzhou University, Zhengzhou, Henan, People’s Republic of China. The study was conducted in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments. The study protocol was approved by the Ethics Committee of Zhengzhou Central Hospital Affiliated to Zhengzhou University, Zhengzhou, Henan, People’s Republic of China.

Competing interests
All authors gave consent for publication.

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