Research Article

Fetal Dermal Mesenchymal Stem Cell-Derived Exosomes Accelerate Cutaneous Wound Healing by Activating Notch Signaling

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Received 5 January 2019; Revised 2 May 2019; Accepted 14 May 2019; Published 10 June 2019

Guest Editor: Ming Li

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Fetal dermal mesenchymal stem cells (FDMSCs), isolated from fetal skin, are serving as a novel MSC candidate with great potential in regenerative medicine. More recently, the paracrine actions, especially MSC-derived exosomes, are being focused on the vital role in MSC-based cellular therapy. This study was to evaluate the therapeutic potential of exosomes secreted by FDMSCs in normal wound healing. First, the in vivo study indicated that FDMSC exosomes could accelerate wound closure in a mouse full-thickness skin wound model. Then, we investigated the role of FDMSC-derived exosomes on adult dermal fibroblast (ADFs). The results demonstrated that FDMSC exosomes could induce the proliferation, migration, and secretion of ADFs. We discovered that after treatment of exosomes, the Notch signaling pathway was activated. Then, we found that in FDMSC exosomes, the ligands of the Notch pathway were undetectable except for Jagged 1, and the results of Jagged 1 mimic by peptide and knockdown by siRNA suggested that Jagged 1 may lead the activation of the Notch signal in ADFs. Collectively, our findings indicated that the FDMSC exosomes may promote wound healing by activating the ADF cell motility and secretion ability via the Notch signaling pathway, providing new aspects for the therapeutic strategy of FDMSC-derived exosomes for the treatment of skin wounds.

1. Introduction

The skin is the largest tissue of the human body and its main function is to guard the underlying tissues. Wound healing is a complex process, and successful cutaneous wound healing needs a series of steps including inflammation, new tissue formation, and remodeling. Furthermore, skin cell migration, proliferation, differentiation, and apoptosis make great contributions to this process. These steps are tightly coordinated and well regulated to restore the multilayered structure of the skin in the normal wound-healing process [1]. Dermal fibroblasts are one of the most important cell lines involved in the normal wound-healing process [2]. The main functions of the dermal fibroblast are extracellular matrix (ECM) production, collagen synthesis, wound contraction, reepithelialization, and tissue remodeling. Once hurt, hemostasis takes place immediately. Fibroblasts, along with other cells including neutrophils, macrophages, and endothelial cells, are attracted to the wound by the blood clot. Then, fibroblasts are activated by macrophages and play a vital role in the proliferative and remodeling phase. Fibroblasts start proliferating and producing ECM proteins like collagen, hyaluronan, and fibronectin to provide a foundation of wound repair [3]. There is a paucity of pharmacological therapeutics that can accelerate wound healing of a large area burn wound and chronic, nonhealing wounds. These wounds adversely affect the life quality of the patients and put great economic pressures on the family and society. Therefore, it is important
to seek an effective therapeutic method to promote wound healing [4].

Mesenchymal stem cells (MSCs) have a significant promise for regenerative medicine. Previous studies demonstrated the therapeutic potential of MSCs for tissue regeneration, including the liver, heart, bone, cartilage, neural, and skin [5–10]. Recent literature suggests that the regenerative effect of MSCs is mainly mediated through paracrine signaling to regulate host cells, instead of cell replacement [5, 11]. Fetal dermal MSCs (FDMSCs), which are derived from the dermis of accidentally aborted fetuses, exhibit advantages of high expansion potential, high differentiation properties, and low immunogenicity. As an advantageous MSC source, FDMSCs have great potential in the tissue regeneration field for their scarless wound-healing characteristic [12–14]. In our previous research, we found that FDMSCs can inhibit the biactivity of keloid fibroblasts by a paracrine manner.

In the last decades, researchers have shown increased interest in exosomes. Exosomes are 40-100 nm small membranous vesicles secreted by most cell types. There are nuclear acids, lipids, and proteins in them, and their main function is to transfer bioactive molecules in cell-cell communication [15, 16]. Moreover, recent studies have shown the role of exosomes in pathogenesis, tissue regeneration, diagnosis, and drug delivery [17–21]. Exosomes are released from MSCs due to paracrine signaling and transfer their cargo of proteins, RNAs, and lipids to recipient cells to regulate the cell state and behaviors. Exosomes derived from MSCs are involved in the acceleration of wound healing [20–22]. We used the promising MSC type, FDMSCs, to investigate the paracrine effect on wound healing process in vivo and in vitro, and to analyze the signal pathway associated with this process.

Notch signaling is an evolutionarily conserved pathway with numerous functions ascribed. Studies over the past decades have proved that Notch plays key roles in stem cell maintenance, development, homeostasis regulation, and cell fate decisions, and its dysfunction can contribute to a variety of diseases in humans [23]. There are 5 ligands (delta-like-1 (Dll-1), Dll-3, Dll-4, Jagged 1, and Jagged 2) in mammals, of which 3 (Dll-1, Dll-3, and Dll-4) are expressed in humans. Delta-like ligands are processed by γ-secretase, which results in the release of their extracellular domains of the Notch signal-binding protein to regulate target gene expression [24, 25]. Cleaved NICD can translocate into the nucleus and combine with a DNA-binding protein to activate the transcriptional expression of the downstream target gene [26]. Furthermore, the nuclear import of NICD is mediated by a nuclear import signal that is recognized by the importin α/β heterodimeric complex [27].

2. Material and Method

2.1. Cell Culture. FDMSCs were extracted from the dorsal skin of fetal samples while adult dermal fibroblasts (ADFs) were extracted from adult skin samples of patient surgical waste. The extraction and identification steps were described in our previous study [30]. These cells were cultured in DMEM/low glucose (HyClone, USA) containing 10% fetal bovine serum (FBS, Gibco, USA) and 1% 100 U/ml Penicillin-Streptomycin (Gibco, USA).

2.2. Isolation and Identification of FDMSC Exosomes. The exosomes were isolated using an ExoQuick-TC kit (SBI, USA) following the instruction. In brief, approximately 80% confluent FDMSCs were washed with PBS twice and cultured for an additional 48 hours in serum-free medium (SFM) containing 1% 100 U/ml Penicillin-Streptomycin. The CM (conditioned medium) was collected and centrifuged at 3,000 × g for 15 minutes to remove cells and cell debris. The supernatant was filtered using a 0.22 μm filter sterilized Sterilop™ (Millipore, USA), and then the supernatant was transferred to an Amicon® Ultra-15 10K Centrifugal Filter Unit (Millipore, USA) to concentrate to 1/5 volume. Appropriate volume of ExoQuick-TC was added in the supernatant in a ratio of 1 : 5 and mixed with the supernatant. After storing at 4°C overnight, the mixture was centrifuged at 1500 × g for 30 minutes to collect the exosomes. The exosomes were quantitated using the BCA Protein Assay Kit (Beyotime, China) following the manufacturer’s protocol. The morphology of the exosomes was observed using a FEI Tecnai G2 Spirit transmission electron microscope (TEM, FEI, USA) after being fixed with 2% glutaraldehyde and counterstained with 4% uranyl acetate. The exosome markers, CD63, Alix, and Tsg101, were detected by Western blot using the specific antibodies. The diameter of exosomes was measured by a ZetaView Nanoparticle Tracking Microscope (Particle Metrix Inc., USA).

2.3. Animal Assay. Animal experiments were approved by the Ethics Committee of the Second Hospital of Shandong University. Studies were performed in 8–10-week-old BALB/c mice weighing 25 ± 5 g. Mice were anesthetized using tribromoethanol and the dorsal hair was shaved. 1 cm × 1 cm full-thickness dermal wounds were created in the skin on the back of the mouse. 200 μl FDMSC-exosomes in 200 μl PBS or 200 μl PBS were injected subcutaneously at four sites around the wound. On days 0, 7, and 14, digital photographs of the injury site were taken. Some mice in each group were euthanized to obtain the skin tissue samples from the wound site by dissection. These samples were collected for histopathological examination by hematoxylin and eosin (H&E) and immunohistochemistry (IHC). In IHC, primary antibodies PCNA (Servicebio, China) and CK19 (Servicebio, China) were used.

2.4. Exosome Internalization. Exosomes were labeled with PKH26 (Sigma-Aldrich, USA) according to the manufacturer’s protocol. Briefly, 5 mg of exosomes was resuspended in 0.5 ml 2 × Diluent C. PKH26 was diluted in 0.5 ml 2 × Diluent C (4 × 10⁻⁶ M). Immediately mix the exosomes and dye solutions to make the final concentrations of PKH26 2 × 10⁻⁶ M. Then, the exosome dye suspension was incubated for 3 min with periodic mixing. 1 ml 1% BSA was then
Table 1: Primer sequences for quantitative real-time PCR.

| Gene name     | Forward                  | Reverse                  |
|---------------|--------------------------|--------------------------|
| Type I collagen | 5′-CGGCAGGAGCATGACCGATGG-3′ | 5′-TCAATGTAGGCGACGCTGTC-3′ |
| Type III collagen | 5′-AAATAAGGGAGAAGCTGGACC-3′ | 5′-GGAGACCCGGGCTCCCATC-3′ |
| Fibronectin-1 | 5′-AAATAGCCCTTGTCAGGAGTCA-3′ | 5′-GAATTTATGGAATTTGGCTTC-3′ |
| Elastin        | 5′-GAGGCAACCTCTTGAAGCC-3′  | 5′-AGCCCCGCGGCACCTAGCCT-3′ |
| α-SMA         | 5′-CATACACCACTGGGAGCGA-3′  | 5′-TCGGTTAGCAAGGTTGAATG-3′ |
| GAPDH         | 5′-GCACGTCAGGCGTGAAC-3′   | 5′-TGGTGAAGAGCCAGCTGGA-3′ |

2.5. Western Blot. Western blotting was performed following standard protocols. Western blotting was used to identify exosome markers CD63, Alix, and Tsg101. Briefly, exosomes were resuspended by PBS and loading buffer and then heated at 95°C for 5 minutes. Cell samples were lysed in RIPA lysis buffer (Beyotime, China) on ice. Then, the samples were loaded and separated in SDS-PAGE gels and transferred onto nitrocellulose membranes (Pall Life Sciences, USA). After incubating with specific antibodies, protein expression and phosphorylation were imaged with FluorChem Q (ProteinSimple, USA). The images were quantified using ImageJ.

2.6. Cell Proliferation. 2 × 10^3 fibroblasts were seeded in 96-well plates in SFM. After overnight plating, a Cell Counting Kit (CCK-) 8 (Beyotime, China) assay was performed to evaluate the cell proliferation according to the manufacturer’s protocol. In brief, cells were treated with MSC exosomes (1 μg/ml, 10 μg/ml, and 100 μg/ml) or SFM for 24 h, and each group contained three parallel wells. 20 μl CCK-8 solution was added to each well and incubated for 2 hours at 37°C. The optical density of each well was measured at 450 nm using the Victor spectrophotometer (Thermo Fisher Scientific, USA).

2.7. Cell Migration. The migration assay was used to analyze the migration effect of FD-MSC exosomes to ADFs. 1 × 10^4 ADFs were seeded in the upper chamber in FDMSC exosomes (1 μg/ml, 10 μg/ml, and 100 μg/ml) or vehicle, and the bottom chambers contained culture media containing 10% FBS and 1% P/S. 24 hours later, cells were fixed with 4% paraformaldehyde and stained with 0.1% crystal violet, and then the upper surface cells were removed with a cotton swab. The cells of the lower membrane surface were counted under a microscope (Nikon, Japan) at ×100 magnification, and 5 random fields were selected.

2.8. Quantitative Real-Time PCR. The primers were synthesized by BGI (China), and the sequences are listed in Table 1. Total RNA was isolated from ADFs and mouse-excised skin wounds using a TRizol reagent (Invitrogen, USA) according to the manufacturer’s instructions. cDNA was synthesized using a HiScript II Q RT SuperMix for qPCR (Vazyme Biotech, China), and real-time PCR was performed using a SYBR Green Master Mix (TAKARA, China) reagent. GAPDH was used as the reference gene for calculations. The ΔΔCt method was used to analyze the real-time PCR data.

2.9. DAPT Treatment. γ-Secretase inhibitor DAPT was purchased from Sigma-Aldrich. In an inhibiting assay, ADFs were incubated with SFM, 100 μg/ml FDMSC exosomes, or 10 μM DAPT + 100 μg/ml FDMSC exosomes for 24 hours. Then, the cell proliferation and migration were evaluated by CCK-8 and Transwell assays.

2.10. Jagged 1 Peptide Treatment. Jagged 1 peptide (CDDYYGFGCNKFCPRR) with Notch agonist activity and scrambled control (SC) peptide (RCGPDCFDNYGRKYCF) was synthesized at the Qiangyao Biological Technology Company (China) [31]. Peptide stock solutions (10 mM) were prepared in sterile distilled water and diluted to 15 μM in culture medium before use.

2.11. siRNA Knockdown. FDMSCs were transfected with Jagged 1 siRNA and control RNA (RiboBio, China). A Lipofectamine™ RNAiMAX Transfection Reagent (Thermo Fisher Scientific, USA) was used according to the manufacturer’s instructions. 10 hours after transfection, ADFs were washed with PBS twice and cultured with SFM. 48 hours later, CM was collected from Jagged 1 siRNA or control siRNA-transfected FDMSCs to isolate Jagged 1 siRNA exosomes and control siRNA exosomes separately.

2.12. Statistical Analysis. Statistical analyses were performed using GraphPad Prism 5. Three or more independent experiments were performed for each result, and the mean and SD were calculated. One-way ANOVA or Student’s t-test...
was used to detect statistically significant differences. A P value < 0.05 was considered as statistically significant.

3. Results

3.1. Characterization of FDMSC Exosomes. FDMSCs were successfully isolated from fetus dorsal skin and identified by flow cytometry analysis and differentiation potential analysis in our previous study [30]. FDMSC exosomes were isolated and then analyzed by TEM and Western blotting. We used TEM to analyze the size, shape, and morphology of exosomes, and the result clearly revealed that FDMSC exosomes have a size range of about 100 nm with an appearance of cup-shaped or round-shaped morphology (Figure 1(a), black arrow). The result showed the presence of exosome marker proteins, CD63, Tsg101, and Alix, in exosome lysates (Figure 1(b)). Then, we measured the size of the exosomes, and the results showed that the diameter of exosomes was about 100 nm (Figure 1(c)).

3.2. FDMSC Exosomes Promote Cutaneous Wound Healing In Vivo. We established a mouse full-thickness dermal wound injury model to investigate the roles of FDMSC exosomes in wound healing. In the exosome-treated group, the wounds healed more rapidly than those in the control group (Figures 2(a) and 2(b)). H&E results indicated that in the exosome-treated group, there are more cells in ECM and the ECM proteins are more regular and denser, with a thicker layer of collagen than that in the control group on 7 days and 14 days posttreatment (Figure 2(c)). Furthermore, in FDMSC exosome-treated wounds, there were more cells with a higher proliferative rate in the wound area evaluated by PCNA IHC results (Figure 2(d)). The IHC result of CK19 illustrated that the reepithelialization in the exosome-treated group was accelerated and the regenerative epidermis is thicker than that in the control group (Figure 2(e)). In summary, our in vivo results indicated that FDMSC exosomes can accelerate cutaneous wound healing by promoting cell proliferation, ECM deposition, and reepithelialization in the wound area.

3.3. FDMSC Exosomes Enhance Proliferation, Migration, and Secretion of ADFs. The main functions of ADFs are to synthesize, secrete, and deposit collagen and elastic fibers of the ECM. Therefore, the proliferation, migration, and protein synthesis abilities of ADFs are vital factors in wound healing. To explore the mechanisms for FDMSC exosome-induced repair, we treated ADFs with FDMSC exosomes. FDMSC exosomes (red) were found to be internalized by the ADFs (Figure 3(a)). To determine the effect of FDMSC exosomes on ADF growth and mobility, CCK-8 and Transwell assays were performed. The results showed that cell proliferation ability of ADFs was significantly improved after being treated with exosomes in a dose-dependent manner (Figure 3(b)). Compared to the control group, the migratory capabilities of ADFs were also significantly improved in the presence of exosomes (Figures 3(c) and 3(d)). These results demonstrated that exosomes significantly enhanced the proliferation
and migration of ADFs in a concentration-dependent manner. Fibroblasts, due to their abilities of synthesis and secretion of ECM proteins, play a significant role during repair of skin wounds. These proteins, to a certain extent, determine the speed and quality of wound healing. Here, we analyzed the mRNA expressions of ECM proteins and wound-healing-related proteins (Type I and III collagen, fibronectin, elastin, and α-SMA) of ADFs by real-time PCR after being treated with FDMSC exosomes. We found that in ADFs incubated with exosomes (1 μg/ml, 10 μg/ml, and 100 μg/ml) for 48 hours, Type I and III collagen, elastin, and fibronectin mRNA production was increased in a dose-dependent manner (Figure 3(e)). The results suggested that the FDMSC exosomes can promote the ECM secretion of ADFs.

Figure 2: FDMSC exosomes accelerate cutaneous wound healing in vivo. (a) Representative photos of mouse dorsal full-thickness wound healing. (b) Quantitative analysis of wound size (n = 5–7 per group). (c) Representative H&E stain images of the wound at 1 week and 2 weeks after treatment (×100 magnification). (d) Representative images of IHC of PCNA in each group (×200 magnification). (e) Representative images of IHC of CK19 in each group (×200 magnification). *P < 0.05.
3.4. FDMSC Exosomes Activate the Notch Signaling Pathway. Recently, researchers have shown the importance of the Notch signal in skin development and tissue regeneration. Therefore, we hypothesized that Notch signaling might be involved in the exosome-mediated wound-healing process. To investigate the underlying mechanism of the effect of FDMSC exosomes on ADFs, the expression level of Notch1, Jagged 1, components of the Notch signaling pathway, and hairy and enhancer of split-1 (Hes 1), a Notch target gene, were analyzed by Western blot. The results showed the increased expression of active Notch1, Jagged 1, and Hes 1, which illustrated the activation of Notch signaling in the presence of FDMSC exosomes (Figures 4(a) and 4(b)). To find how the Notch signaling was activated, we detected the
Notch ligands in exosomes and found that Jagged 1 was packaged into MSC exosomes while the others were undetectable by Western blot (Figure 4(c)).

3.5. DAPT Can Partly Block the Promoting Effect of FDMSC Exosomes of ADF Proliferation and Migration.

To determine whether exosomes can promote ADF proliferation and migration in a Notch-dependent manner, we treated ADFs with DAPT, the \( \gamma \)-secretase inhibitor, to block Notch receptor cleavage at the cell surface. ADFs were treated with SFM, 100 \( \mu \)g/ml exosomes or DAPT 10 \( \mu \)M + exosomes 100 \( \mu \)g/ml. We found that DAPT partly abolished the positive regulating effect in cell proliferation (Figure 5(a)) and migration (Figures 5(b) and 5(c)) of FDMSC exosomes on ADFs. These results indicate that FDMSC exosomes can activate the wound-healing capacity of ADFs via the Notch signaling pathway, and these effects can also be inhibited when DAPT was used, illustrating the role of the Notch pathway in wound healing.

3.6. Jagged 1 in FDMSC Exosomes Promote the Wound-Healing Capacity of ADFs.

To further investigate the functional role of Jagged 1 expressed in exosomes in wounding, we used the Jagged 1 peptide to mimic Jagged 1 in activating the Notch signal and knockdown Jagged 1 expression in FDMSCs by siRNA. The expression of Jagged 1 in FDMSC exosomes was reduced after siRNA knockdown (Figure 6(a)). ADFs were incubated with SFM, 100 \( \mu \)g/ml FDMSC exosomes, 15 \( \mu \)M Jagged 1 peptide, or 100 \( \mu \)g/ml Jagged 1 knockdown exosomes for 24 hours. We found that in the FDMSC exosome and Jagged 1 peptide treatment groups, the Notch pathway was activated, and the proliferation and migration ability of ADFs was increased, while depletion of Jagged 1 in FDMSC exosomes by siRNA blocked the activation of Notch signaling and blocked the promoting ability of FDMSC exosomes on the proliferation and migration of ADFs (Figures 6(b)–6(e)). These results indicated that Jagged 1 in FDMSC exosomes can activate the wound-healing capacity of ADFs via the Notch signaling pathway.

4. Discussion

Wound healing is an integrated and coordinated process of different cells functionally relevant to skin tissue repair, alone with the microenvironment around them. There are a large number of published studies that describe the treatment methods for the management of cutaneous wound healing; however, the questions and difficulties are still remaining in this field. Especially for nonhealing and chronic wounds, effective therapeutic approaches need to be further explored to deal with this prevalent and costly public health issue. Thus, it is urgent to find an effective approach to prompt wound healing [32].

In the last decades, researches and clinical trials of MSC applications in tissue regeneration have made great progress. Studies focused on MSC transplantation suggested that instead of direct cell differentiation and replacement, MSCs play regulation and stimulation roles via paracrine signaling by releasing factors that promote angiogenesis, immunomodulation, and recruitment of different cells [5, 9, 33]. Literature has proved the positive effect of MSC CM on tissue regeneration [5, 34, 35]. Growth factors, cytokines, immunomodulatory proteins, and other biologically active proteins are the major components of CM. Besides, the discovery of exosomes helps us gain a better understanding of the underlying mechanism of the multiple effects of MSCs throughout.
the body [36–38]. Exosomes can mediate the cell-cell communication by transferring RNAs, proteins, and lipids to recipient cells and modifying their bioactivity state [15]. Nowadays, exosomes are considered as novel therapeutic tools and diagnostic markers [17, 39, 40].

MSC exosomes can exhibit repair effect, consistently with the MSCs, on the injured tissues through modifying recipient cell gene expression, protein production, and status, as well as activating regeneration-associated pathways including Wnt/β-catenin, AKT, ERK, and STAT3 [41–43]. Recently, investigators have examined the regenerative effects of exosomes derived from MSCs on tissues of the lung, heart, kidney, liver, brain, and so on. Therefore, exosomes derived from MSC exosomes may become potential therapeutic agents in cell-free tissue regeneration therapy. According to advanced research in wound healing, MSC exosomes can increase the proliferation and migration of skin cells and inhibit their apoptosis.

Fetal MSCs are a new potential source of MSCs. The dorsal skin of aborted fetuses, which is considered as clinic discards, is an alternative abundant source of MSCs, and the clinic significance needs to be further explored. Compared with adult MSCs, fetal MSCs exhibit low immunogenicity, higher proliferation, and differentiation potential. FDMSCs are derived from accidental aborted fetuses, and they are thought as the main functional cells involved in scarless wound healing [12]. Furthermore, owing to the histological origin of FDMSCs, they may deserve unique properties on skin regeneration. In summary, FDMSCs are better candidates than adult MSCs in wound healing.

Fibroblasts, as the important target of exosomes in wound healing, are the major cell type to synthesize, secrete, and deposit collagen and elastic fibers of the ECM [2]. Recently, there has been renewed interest in the different fibroblast lineages [24, 44]. Researchers found that fibroblasts isolated from different dermal sources exhibit diverse functions, and the underlying mechanism needs to be explored further. Fibroblasts from diabetic patients showed impaired function in wound healing with reduced migration response and growth factor expression [45, 46]. In summary, the proliferation, migration, and protein synthesis abilities of dermal fibroblasts are vital for wound repair. Activation of fibroblasts in the early phase of wound healing can accelerate the wound closure and matrix protein production, providing a foundation for wound repair.

In our study, results suggested that FDMSC exosomes have an enhancing effect on ADF cell growth and migration. Further analysis by real-time PCR showed significantly elevated ECM protein levels compared to those of the control group, indicating that FDMSCs can promote ECM protein synthesis. The upregulation of Notch1, Jagged 1, and Hes 1 exhibited the activating effect of FDMSC exosomes on Notch signaling. Furthermore, Western blot analysis of exosome components showed Jagged 1 was the only ligand that can be detected, and the inhibition of Notch signaling by DAPT significantly decreased the proliferation and migration of

![Figure 5: Activated Notch signaling by FDMSC exosomes can be partly blocked by DAPT. (a) DAPT can partly block the proliferation of ADFs. (b, c) DAPT can partly block the migration of ADFs. (b) Cell migration is expressed as a percentage of control. (c) Representative images of the migration (×100 magnification). **P < 0.01 or ***P < 0.001.](image)
ADFs. In contrast, ADFs treated with FDMSC exosomes and Jagged peptide showed significantly enhanced proliferation and migration, and knockdown of Jagged 1 in exosomes abolished the promoting effect. These results emphasizing that the Notch pathway is a mediator of exosome communication in regulating wound repair and Jagged 1 in exosomes play a vital role.

As one of the important Notch ligands, Jagged 1 can regulate maturation of the human epidermis by activating Notch signaling [31]. In addition, Jagged 1 is present in exosomes from different kinds of cells and is biologically active, but the role of Jagged 1 in exosomes in wound healing is largely unknown [47–49]. In this study, we found that Jagged 1 is sorted in FDMSC exosomes to regulate the Notch signal pathway activity in ADFs. However, the quantity of Jagged 1 in FDMSC exosomes is variable and unstable because the biogenesis of exosomes is largely depending on cell types, cell functions, and physiological statuses. Due to the complexity of FDMSC exosomes, the important components of the exosomal cargo and other factors which can activate Notch signaling and the mechanism are still studying. Further research is needed to elucidate a detailed molecular mechanism of the sorting process and biological functions of Jagged 1 and the exact mechanism of FDMSC exosomes in wound healing and to develop new therapeutic strategies for non-healing and chronic wounds.

In conclusion, we successfully obtained FDMSC exosomes and investigated their role on cutaneous wound healing. Our results demonstrated that FDMSC exosomes could accelerate cutaneous wound healing in vivo and promote the wound-healing capacities of ADFs by activating the Notch signal pathway in vitro. Our findings provided new aspects for the therapeutic strategy of FDMSC-derived exosomes for the treatment of skin wounds.

5. Conclusion

The results demonstrated that FDMSC exosomes could accelerate cutaneous wound healing in vivo and promote the wound-healing capacities of ADFs by activating the Notch signal pathway in vitro. Our findings provided new aspects for the therapeutic strategy of FDMSC-derived exosomes for the treatment of skin wounds.

Abbreviations

α-SMA: α-Smooth muscle actin
ADFs: Adult dermal fibroblasts
CM: Conditioned media
ECM: Extracellular matrix
FDMSCs: Fetal dermal mesenchymal stem cells
H&E: Hematoxylin and eosin
Hes 1: Hairy and enhancer of split-1
IHC: Immunohistochemistry
NICD: Notch intracellular domain
SFM: Serum-free medium
siRNA: Small interfering RNA
TACE: Tumor necrosis factor alpha converting enzyme

Data Availability
The data used to support the findings of this study are available from the corresponding author upon request.

Ethical Approval
We received the ethical approval of the Ethics Committee of the Second Hospital of Shandong University on fetal skin isolation. The ethics certificate was issued on 1st Jan 2018 and the certificate number is KYLL-2018(LW)-021.

Consent
We received the informed consent of the patients.

Conflicts of Interest
The authors declare that they have no conflicts of interest.

Authors’ Contributions
Xiao Wang designed the experiments, performed the experiments, analyzed the data, prepared the figures, and wrote the manuscript. Yi Pan and Ya Jiao performed the experiments, analyzed the data, and proofread the manuscript. Longxiao Zhang performed the histological experiments and analyzed the results. Yongjun Qi, Hongmin Gong, and Maoying Wang performed experiments. Duyin Jiang designed the experiments and supervised the research. All authors read and approved the final manuscript. Xinglei Wang and Duyin Jiang contributed equally.

Acknowledgments
We thank the members of the central laboratory of the Second Hospital of Shandong University. This work was supported by the National Natural Science Foundation of China (81873934), Science and Technology Development Projects of Shandong Province (2015GSF118041), and Youth Fund of the Second Hospital of Shandong University (2018YT14).

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