Epidermal stem cell-derived exosomes promote skin regeneration by downregulating transforming growth factor-β1 in wound healing

Mengna Duan 1,2, Yan Zhang 1, Haiyang Zhang 1, Yupeng Meng 1, Ming Qian 1* and Guokun Zhang 3,4*

Abstract

Background: Scar formation, which may be caused by myofibroblast aggregations, is the greatest challenge during skin wound healing in the clinical setting. Studies have indicated that epidermal stem cells (EPSC) improve wound healing and reduce scar formation.

Methods: We investigated the therapeutic effects of EPSC-derived exosomes (EPSC-Exos) on skin wound healing in a skin-defect rat model. We also examined the roles of EPSC-Exos-specific microRNAs in inhibiting the differentiation of human dermal fibroblasts (HDF) into myofibroblasts.

Results: We found that EPSC-Exos increased the wound healing rate and reduced scar formation in rats. Also, EPSC-Exos improved the regeneration levels of skin appendages, nerves and vessels, as well as the natural distribution of collagen. Furthermore, we found these functions may be achieved by inhibiting the activity of transforming growth factor-β1 (TGF-β1) and its downstream genes. The results showed that some specific microRNAs, including miR-16, let-7a, miR-425-5p and miR-142-3p, were enriched in EPSC-Exos. EPSC-Exos-specific microRNAs, especially miR-425-5p and miR-142-3p, played vital roles in inhibiting myofibroblast differentiation via reducing the TGF-β1 expression in dermal fibroblasts.

Conclusion: We found a novel function of EPSC-Exos-specific microRNAs, suggesting that EPSC-Exos might represent a strategy to prevent scar formation during wound healing in the clinical setting.

Keywords: Exosome, Epidermal stem cells, MicroRNA, Myofibroblast, Transforming growth factor-β1

Background

The skin is a multilayer interface between the body and the environment, which can regulate temperature, prevent dehydration, keep out pathogens, provide sensation and transport water [1–3]. Once the skin is damaged, the body will immediately start a repair programme to complete the recovery of the skin’s structure and function. However, fibrosis or scarring during the healing process cannot be avoided, even under optimal conditions [3, 4]. Many approaches have attempted to achieve the best healing result, namely the double reconstruction of structure and function. Stem cell-based therapy, with unique advantages, is emerging as a promising candidate for wound healing and skin regeneration [5, 6]. Epidermal stem cells (EPSC) are a type of autologous adult stem cells that are easily obtainable from the skin. EPSC reportedly differentiated into hair follicles and sweat
glands for use in tissue repair, decreasing scar formation and providing long-term regeneration [7–11]. However, there is only a small amount of EPSC in the healing tissue after EPSC treatment. According to the mainstream view [12], the strong paracrine ability of EPSC may be the essential mechanism resulting in these therapeutic effects.

Exosomes, secreted by parental cells, are a type of membrane structure, encapsulating various mRNAs, miRNAs, LncRNAs, CircRNA and proteins to regulate the activity of recipient cells. Exosomes play important roles in tissue regeneration [13–15]. They have a similar function to their parent cells, but they are easier to store and transport, protecting the bioactive substances they carry from adverse conditions such as pH environments, high temperature and repeated freezing and thawing [16–18]. Moreover, they avoid many of the risks associated with cell transplantation. Therefore, exosome therapy may be a more safe and efficient option.

Transforming growth factor-β1 (TGF-β1) is the primary regulator in wound healing and tissue repair. It is responsible for cellular proliferation, differentiation and metabolism [3, 19]. Excessive scar formation and fibrosis are closely related to high expression levels of TGF-β1 in wound healing [1, 3, 20]. TGF-β1 induces the differentiation of myofibroblasts via the phosphorylation of the SMAD family of proteins and activating downstream-related genes such as α-smooth muscle actin (α-SMA) and collagen I [3, 19, 20]. Therefore, inhibiting the activities of TGF-β1 and its downstream genes might reduce myofibroblast formation and over-aggregation to decrease scarring.

In the current study, we investigated whether EPSC-derived exosomes (EPSC-Exos) are effective in the healing of cutaneous wounds using full-thickness skin-defect rat models. Moreover, we identified two specific microRNAs carried by EPSC-Exos as critical components contributing to the suppression of the differentiation of fibroblasts to myofibroblasts by inhibiting TGF-β1 expression. Thus, the results suggest that EPSC-Exos treatment could be a potential strategy to prevent excessive fibrosis and scar formation during wound healing in injured skin.

Materials and methods

Cell culture
The cell lines of EPSC and human dermal fibroblasts (HDF) were obtained from the China-Japan Union Hospital of Jilin University. Cells were cultured in Dulbecco's modified Eagle medium (+10% foetal bovine serum; Biological Industries, Beit-Haemek, Israel) with 5% CO₂ at 37 °C. Cells at the 3–5 passages were used for this study.

Exosome isolation
EPSC-Exos were purified by ultracentrifugation according to previously reported methods [15, 18]. Briefly, EPSC were cultured in a serum-free medium (Gibco, Grand Island, NY) for 48 h. The culture suspension was filtered (0.1-μm filter device) and concentrated (100-kDa molecular weight cutoff), then loaded onto a 30% sucrose/deuterium oxide (D2O) cushion and ultracentrifuged at 100,000×g for 3 h. The exosomes were then washed with phosphate-buffered saline (PBS) 3 times and centrifuged at 1500×g for 30 min (100-kDa molecular weight cutoff). The protein concentration of the exosomes was determined using a bicinchoninic acid protein assay kit (Solarbio, Beijing, China). The exosomes were stored at −80 °C.

Creation of skin injury model in rats and treatment with EPSC-Exos
The Animal Experimental Ethics Committee of Jilin University approved all study protocols and procedures (Approval No. SY201902010). Thirty Sprague-Dawley rats (8-week-old, female, 200 g) were purchased from Liaoning Changsheng Biotechnology Co., Ltd. (Benxi, China) and reared under standard experimental animal feeding conditions. The rats were anaesthetised with 10% chloral hydrate (3 ml/kg), after which their hair was removed from the dorsal surface. A 12-mm diameter was removed to establish a full-thickness skin-defect model.

EPSC-Exos was dissolved into HydroMatrix (Sigma-Aldrich, St. Louis, MO) used as a scaffold. Briefly, 100 μg/ml of EPSC-Exos dissolved in PBS and 10 mg/ml (1%) hydrogel mixed at a ratio of 1:1 and injected (200 μl) around the wound once a week for 4 weeks. Skin damage was recorded photographically every 7 days. The area of each wound was calculated out using Adobe Photoshop CS6. Firstly, lasso tool was used to trace the edge of a wound on a photograph and to circle it; then, we calculate the circled area based on the pixels of that area (1 cm = 28.346 pixels). PBS alone served as a negative control, and epidermal growth factor (EGF; Beyotime, Shanghai, China) mixed with hydro-matrix served as a positive control, both PBS and EGF + hydro-matrix were also injected around the wound once a week for 4 weeks.

The rats were euthanised after being anaesthetised with 10% chloral hydrate (3 ml/kg), and the healing wound tissues collected for later use.

Histopathological analysis
Skin tissue was embedded in paraffin, then sliced into 4-μm sections. The sections were stained with haematoxylin and eosin (Solarbio, Beijing, China), Masson (Solarbio), immunohistochemistry (IHC), following the manufacturer’s instructions and conventional methods. The primary antibodies used in IHC were anti-CD31 (bs-0468R, 1:500 dilution; Bios, Beijing, China), anti-Nestin (bs-0008R, 1:500 dilution; Bios) and Ki67 (bs-23105R, 1:500 dilution; Bios). Sections were photographed using the
Digital Imaging Scanning System (Precipoint M8; Precipoint, Freising, Germany). The Masson staining and IHC results were analysed using Image-Pro Plus software. Firstly, we counted the area of the target item in the picture and then calculated its ratio to the total area of the picture.

Co-culture system of EPSC and HDF
The 5 × 10⁴ HDF and ESCs were seeded with Dulbecco’s modified Eagle medium and 10% foetal bovine serum on the upper (0.4 μm) and lower chambers of 24-well plates (Corning Life Sciences, NY). After incubating plates for 48 h at 37 °C, TGF-β1 expression levels of HDF were performed using quantitative immunofluorescence (IF) staining and quantitative real-time polymerase chain reaction (qRT-PCR).

qRT-PCR
Trizol reagent (Tiangen, Beijing, China) was used for total RNA isolation of the skin tissue. cDNA was prepared via reverse transcription of RNA. cDNA, primers, and SYBR premix (Roche, Basel, Switzerland) were combined on qRT-PCR using the ABI 9700 Detection System (Thermo Fisher Scientific, Waltham, MA), and GAPDH was used as control. Supplemental Table S1 shows the primers. All experiments were repeated three times. qRT-PCR for miRNA was done using miScript SYBR Green PCR kit (Qiagen, Hilden, Germany) according to the manufacturer’s instruction. All results were normalised to U6 small RNA levels measured using the Hs_RNU6B_2 miScript Primer Assay kit (Qiagen). Supplemental Table S1 shows the primers. All experiments were repeated three times.

IF staining
HDF with a density of 70–80% were incubated for 10 min with 10% formaldehyde and for 30 min with 1% bovine serum albumin (Beyond, Shanghai, China). Next, the cells were incubated with anti-TGF-β1 (BSM-33345M, 1:500 dilution; Bioss) at 4 °C overnight and incubated with anti-IgG (A0408, 1:500 dilution) at 25 °C for 30 min. The nucleus was labelled with 4′,6-diamidino-2-phenylindole (Beyotime, Shanghai, China).

Statistical analysis
All quantitative data were expressed as the mean ± SD. One-way analysis of variance with Tukey’s multiple comparisons test was used for statistically significant
differences. Results were considered significant at a $P$ value of less than 0.05.

**Results**

**EPSC-Exos improve wound healing rate and suppress scar formation**

We first identified the purified EPSC-Exos. Transmission electron microscopy results showed that the particle size of EPSC-Exos was between 30 and 100 nm (Fig. 1a). The RNA in EPSC-Exos was analysed via agarose gel electrophoresis. There was only a clear band at 60 nt (Fig. 1b). Analysis using the Agilent 2100 Bioanalyzer confirmed these results (Fig. 1b). Western blot analysis was used to detect the expression of exosomal markers CD9 and CD63, and serum-free medium was used as a control. The results showed that CD9 and CD63 were abundant in EPSC-Exos (Fig. 1c). In summary, the isolated EPSC-Exos has high purity and can be used in subsequent experiments.

To clarify the effects of EPSC-Exos in the wound healing rate and scar formation, we used full-thickness skin-defect rats and injected equal quantities of hydrogel-coated EPSC-Exos, PBS or EGF around the wounds. The results are shown in Fig. 2. After 1 or 2 weeks of treatment, the wound areas and scars of the EPSC-Exos group were smaller than those of the EGF and control groups. After 3 weeks, the defects in the EPSC-Exos group were closed.
but those in the EGF and control groups were not. After 4 weeks, the defects in the EGF and control groups were also closed. However, they exhibited more scar formation than that observed in the EPSC-Exos group. These results suggest that EPSC-Exos can improve the wound healing rate and suppress scar formation.

**EPSC-Exos improve the regeneration level of skin appendages and collagen distribution on wound healing of rats**

After evaluating the effect of EPSC-Exos on increasing the wound healing rate and decreasing scar formation in shape and appearance, we made further evaluations at the histological level. As shown in Fig. 3a and b, the results showed that after 4 weeks of treatment, the skin in the EPSC-Exos group exhibited more appendages than that in the EGF and control groups (significant; \( P < 0.001 \)). There were significantly fewer myofibers (red) in the EPSC-Exos group than the EGF and control groups (\( P < 0.001 \)) (Fig. 3a, c). In addition, there were significantly fewer collagen fibres (blue) in the skin of the EPSC-Exos group than observed in the EGF group (\( P < 0.01 \)) (Fig. 3a, d). Moreover, among the regenerated collagen fibres, the rates of collagen I and III in the EPSC-Exos group were higher than those in the EGF and control groups.

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**Fig. 3** EPSC-Exos improved the regeneration level of skin appendages and the collagen distribution on wound healing of the rats. **a** Haematoxylin and eosin and Masson staining of the wound healing tissues. Scale bar = 100 \( \mu \)m. **b** Number of skin appendages (hair follicles and sebaceous glands) in the healing tissue according to haematoxylin and eosin staining. **c, d** The proportion of myofibers (red) and collagen fibres (blue) in the healing tissue, according to Masson staining. **e-g** Collagen I mRNA, collagen III mRNA and collagen I/collagen III levels in the healing tissue. \( \ast P < 0.05; \ast\ast P < 0.01; \ast\ast\ast P < 0.001; \) mean ± SD; \( n = 5 \). Ctrl, control; EGF, epidermal growth factor; EPSC-Exos, epidermal stem cell-derived exosomes
Exos group were lower than in the EGF and control groups (Fig. 3e–g). These results suggest that EPSC-Exos can promote skin appendage regeneration, decrease myofiber formation and increase collagen III in wound healing.

**EPSC-Exos promote nerve and vessel regeneration on wound healing in rats**

Next, we clarified the effects of EPSC-Exos on nerve and vessel regeneration of damaged skin using IHC. The results showed that after 4 weeks of treatment (as shown in Fig. 4a–c), the skin of the EPSC-Exos group exhibited significantly more vessels (CD31+) and nerves (nestin+) than in the EGF and control groups ($P<0.001$). Also, the proliferation (Ki67+) of skin cells in the EPSC-Exos group was also higher than in the EGF and control groups (Fig. 4d). These results suggest that EPSC-Exos can promote nerve and vessel regeneration and cell proliferation in wound healing.

**EPSC-Exos inhibit TGF-β1 expression from suppressing myofibroblast formation in vivo and in vitro**

TGF-β1 is the main factor in inducing dermal fibroblasts and the most important cell in skin wound healing and for differentiation into myofibroblasts [1, 3, 4]. Inhibiting TGF-β1 may be an essential means to reduce scar formation in wound healing. To explore the mechanism of EPSC-Exos inhibiting scar formation in wound healing, we measured the TGF-β1 expression levels in the healing tissue of rat models and HDF co-cultured with ESCs. The results showed that the mRNA levels of TGF-β1 and its downstream genes, including Smad2, α-SMA and collagen I, in the healing tissue of rats were all significantly lower than in the EGF and control groups ($P<\)
0.001) (Fig. 5a–d). Also, expression levels of TGF-β1 were evaluated on HDF treated with EPSC-Exos, co-cultured with ESC in Transwell plates for 48 h (Fig. 5e).

As shown in Fig. 5f and g, expression levels of TGF-β1 treated with EPSC-Exos were significantly reduced in mRNA and protein compared with the control group (P<0.001). These results suggest that EPSC-Exos may inhibit the expression of TGF-β1 to suppress myofibroblast formation in wound healing.

**EPSC-Exos-specific microRNAs target TGF-β1 to suppress myofibroblast differentiation**

We found that miR-16, let-7a, miR-425-5p and miR-142-3p were highly expressed in EPSC-Exos detected using qRT-PCR (Fig. 6a). To predict the potential target genes of these microRNAs, we used TargetScan (http://www.targetscan.org/) and Microrna (http://www.microrna.org) for analysis. The results showed that miR-425-5p and miR-142-3p were all directly targeted to TGF-β1 (Fig. 6b) To verify the role of miR-425-5p and miR-142-3p in EPSC-Exos, we first added agomirs in HDF cultural system to test whether they can affect the expression of TGF-β1. The qRT-PCR and IF analysis showed that these 2 microRNAs significantly suppressed the expression of TGF-β1 (Fig. 6c, d). The results suggest that EPSC-Exos may inhibit the differentiation of fibroblasts to myofibroblasts via suppressing TGF-β1 expression via miR-425-5p and miR-142-3p.

**Fig. 5 EPSC-Exos inhibited the expression of transforming growth factor-β1 (TGF-β1) and its downstream genes to suppress myofibroblast formation.**

**a–d** mRNA levels of TGF-β1, Smad2, α-SMA and collagen I in the healing tissue. **e** ESCs were co-cultured with HDF in Transwell plates (membrane well, 0.4 μm). IF and qRT-PCR were performed after culture for 48 h. **f, g** Expression levels of TGF-β1 of HDF in a co-cultured system using qRT-PCR and IF. Scale bar = 200 μm. ***P < 0.001; mean ± SD; n = 5. Ctrl, control; EGF, epidermal growth factor; EPSC-Exos, epidermal stem cell-derived exosomes; ESC, epidermal stem cell; HDF, human dermal fibroblast; TGF-β1, transforming growth factor β-1; α-SMA, α-smooth muscle actin; IF, immunofluorescence staining.
**Discussion**

Scar formation is an undesirable and significant result of both wound healing and fibrosing disorders. Excessive accumulation of myofibroblasts is a crucial factor for scar development, which can even lead to tissue or organ contraction [1, 4, 21]. Regulating myofibroblast formation may be an effective approach to preventing abnormal scarring. EPSC-based therapies have been proven to improve wound healing and reduce scar formation [22, 23]. Researchers have evaluated the effects of EPSC on tissue repair and wound healing. However, no previous studies have focused on the effects of EPSC-Exos on wound healing and scar formation. This study is the first to report that EPSC-Exos suppress myofibroblast differentiation in wound healing, which may be achieved by EPSC-Exos-specific microRNAs. Our research provides new insights for the application of EPSC-Exos to prevent scar formation.

EPSC, similar to adipose stem cells, are a cell type widely used in clinical research and regenerative medicine because they are numerous, are easy to obtain and have fewer ethical controversies than embryonic stem cells [22, 24, 25]. EPSC promote re-epithelisation, angiogenesis and hair growth in wound healing, as well as stimulate the recruitment and proliferation of endogenous stem cells [22, 23, 25]. We speculated that EPSC might play a major role in cell therapy through paracrine. Therefore, the paracrine functions of EPSC were the focus of this study. We observed that the healing skin in the EPSC-Exos group was smooth with less scarring, which was significantly lower than that in the control and EGF groups.

Furthermore, we confirmed that the expression levels of TGF-β1 were lower than in the control and EGF groups. TGF-β1 is reportedly a pivotal factor in simulating fibroblast differentiation into myofibroblasts [26, 27]. Therefore, intervention is required during wound healing...

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**Fig. 6 EPSC-Exos-specific microRNAs inhibited TGF-β1 expression of HDF.**

- **a** Main miRNAs in EPSC-Exos using qRT-PCR.
- **b** A list of predicted binding sites of EPSC-Exos-specific miRNAs and their targets.
- **c, d** TGF-β1 mRNA levels of HDF treated with miR-425-5p agomir and miR-142-3p agomir of HDF, respectively. Scale bar = 200 μm. *P < 0.05; ***P < 0.001; mean ± SD; n = 5. Ctrl, control; EGF, epidermal growth factor; EPSC-Exos, epidermal stem cell-derived exosomes; HDF, human dermal fibroblast; TGF-β1, transforming growth factor β1; IF, immunofluorescence staining.
healing to prevent excessive formation of myofibroblasts, rather than taking remedial measures after scar formation.

Exosomes have content that includes mRNAs, regulatory miRNAs, functional protein and signalling lipids. Thus, they involve cell signalling and cell-to-cell communication and they influence tissue responses to injury, infection and disease [14, 28]. In this study, we found that the primary function of promoting skin healing and reducing scar formation may be caused by EPSC-Exos-specific miRNAs. The EPSC-Exos-specific miRNAs inhibited myofibroblast formation, which might be related to the targeting of TGF-β1-targeting microRNAs. In our previous study, several highly expressed specific microRNAs derived from EPSC-Exos were identified, including miR-16, let-7a, miR-425-5p and miR-142-3p. miR-425-5p [29] and miR-142-3p [30] have been previously reported to suppress fibrotic diseases, and most studies have suggested that these two microRNAs directly target TGF-β1, consistent with our present findings. We believe the functions of microRNAs in different organs and tissues might be the same. Therefore, our study on the effects of EPSC-Exos on these two microRNAs might help us better understand microRNAs’ functions. We believe these EPSC-Exos miRNAs could be essential inhibitors of TGF-β1, reducing the myofibroblast formation during skin wound healing.

Exosome-derived microRNAs are more stable than their parental cells. They can resist degradation in the circulation process in the body through the vesicle structure released by the parental cells [18, 31]. In the subsequent production and application, we can use EPSC as a factory to produce exosomes, making EPSC produce more functional microRNAs via transfection. Compared with EPSC,
EPSC-Exos might be safer and more efficient in clinical applications. They may also have other advantages, such as easier storage and production, a lower risk of side effects and easier quality control. Thus, we suggest that EPSC-Exos could be a candidate strategy for promoting healing and reducing scar formation in the future.

Conclusions
This study has shed light on the specific microRNAs of EPSC-Exos and clarified a new approach for using stem cell therapy to promote wound healing and reduce scarring. miR-425-5p and miR-142-3p from EPSC-Exos might suppress myofiber and collagen I deposition via downregulating TGF-β1 expression (Fig. 7). As an alternative to cell therapy, EPSC-Exos might have a clinically beneficial anti-scarring effect.

Supplementary information
Supplementary information accompanies this paper at https://doi.org/10.1186/s13287-020-01971-6.

Abbreviations
α-SMA: α-Smooth muscle actin; CD31: Platelet endothelial cell adhesion molecule-1; EGF: Epidermal growth factor; EPSC: Epidermal stem cells; EPSC-Exos: EPSC-derived exosomes; HDF: Human dermal fibroblasts; IF: Immunofluorescence staining; IHC: Immunohistochemical staining; PBS: Phosphate-buffered saline; TGF-β1: Transforming growth factor-β1

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Authors’ contributions
MD and GZ designed and supervised the research; YM contributed to experiments; MD and YM contributed to histopathological experiments; and MD, MQ and GZ wrote and edited the paper. All authors read and approved the final manuscript.

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Availability of data and materials
The datasets used and/or analysed during the present study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
This study was approved by the Administration Committee of Experimental Animals Jilin University (Approval No. SY201902010).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

Author details
1Department of Prosthodontics, Hospital of Stomatology, Jilin University, 1500 Qinghua Rd., Changchun 130021, Jilin, China. 2Jilin Provincial Laboratory of Biomedical Engineering, Jilin University, 1500 Qinghua Rd., Changchun 130021, Jilin, China. 3Department of Prosthodontics, Hospital of Stomatology, Jilin University, 1500 Qinghua Rd., Changchun 130021, Jilin, China. 4Institute of Antler Science and Product Technology, Changchun Sci-Tech University, 1345 Pudong Rd, Changchun 130600, Jilin, China. 5Institute of Special Animal and Plant Sciences, Chinese Academy of Agricultural Sciences (CAAS), 4899 Juye St., Changchun 130112, Jilin, China.

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