Human Novel MicroRNA Seq-915_x4024 in Keratinocytes Contributes to Skin Regeneration by Suppressing Scar Formation

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Early in gestation, wounds in fetal skin heal by regeneration, in which microRNAs play key roles. Seq-915_x4024 is a novel microRNA candidate confirmed by deep sequencing and mirTools 2.0. It is highly expressed in fetal keratinocytes during early gestation. Using an in vitro wound-healing assay, Transwell cell migration assay, and MTS proliferation assay, we demonstrated that keratinocytes overexpressing seq-915_x4024 exhibited higher proliferative activity and the ability to promote fibroblast migration and fibroblast proliferation. These characteristics of keratinocytes are the same biological behaviors as those of fetal keratinocytes, which contribute to skin regeneration. In addition, seq-915_x4024 suppressed the expression of the pro-inflammatory markers TNF-α, IL-6, and IL-8 and the pro-inflammatory chemokines CXCL1 and CXCL5. We also demonstrated that seq-915_x4024 regulates TGF-β isoforms and the extracellular matrix. Moreover, using an in vivo wound-healing model, we demonstrated that overexpression of seq-915_x4024 in keratinocytes suppresses inflammatory cell infiltration and scar formation. Using bioinformatics analyses, luciferase reporter assays, and western blotting, we further demonstrated that Sar1A, Smad2, TNF-α, and IL-8 are direct targets of seq-915_x4024. Furthermore, the expression of phosphorylated Smad2 and Smad3 was reduced by seq-915_x4024. Seq-915_x4024 could be used as an anti-fibrotic factor for the treatment of wound healing.

INTRODUCTION

The healing of cutaneous wounds is a highly orchestrated, multistep process that is composed of a complex series of molecular events that prevent infection, repair the damaged tissue, and restore its function.1-3 Wound healing includes three phases: inflammation, re-epithelialization, and remodeling.4,5 During the remodeling stage, wounds in adult cutaneous tissues are usually filled with hyperproliferative granulation tissues, which gradually evolve into scar tissues (fibrosis).1

To restore skin tissue, regeneration that leads to scarless healing is the ideal method. Before the third trimester of gestation (early to mid-gestation), wounds in fetal mammalian skin heal rapidly, without scarring or inflammation.4-9 The fetal skin is gradually replaced by the adult skin, which is characterized by scar formation.10-12

Epidermal keratinocytes (KCs) play important roles in the process of skin tissue regeneration. Early in gestation, fetal KCs exhibit a higher proliferation rate. In addition, fetal KCs enhance fibroblast (FB) proliferation and contribute to the extracellular matrix (ECM), which is related to scarless wound healing.13,14 MicroRNAs (miRNAs), which are a large family of highly conserved, small, noncoding RNAs, play important regulatory roles over a vast number of protein-coding genes.1,15 Previously, we identified a novel miRNA candidate, seq-915_x4024, which was highly expressed in mid-gestational fetal KCs, and we predicted that seq-915_x4024 may target the transforming growth factor (TGF)-β1-Smad signaling pathway, which plays important roles in skin regeneration.10,14

In this study, we investigated the roles that seq-915_x4024 plays in wound healing, using the HaCaT cell line (an immortalized human KC cell line). After being transfected with seq-915_x4024 mimics,
HaCaT cells exhibited higher proliferative activity and the ability to promote FB migration and FB proliferation. These characteristics of KCs were the same as the biological behaviors of fetal KCs at mid-gestation. In addition, seq-915_x4024 suppressed the pro-inflammatory cytokines tumor necrosis factor-α (TNF-α), interleukin-6 (IL-6), and IL-8 and the pro-inflammatory chemokines CXCL1 and CXCL5. We also demonstrated that seq-915_x4024 regulates the expression of TGF-β isoforms and ECM, including collagen and hyaluronan. Moreover, using an in vivo wound-healing model, we demonstrated that overexpression of seq-915_x4024 in KCs suppresses inflammatory cell infiltration and scar formation. The in vivo expression of collagen type I (Col I), collagen type III (Col III), phosphorylated Smad2 (p-Smad2), and phosphorylated Smad3 (p-Smad3) was suppressed by seq-915_x4024. Furthermore, we used bioinformatics analyses, luciferase reporter assays and western blotting to demonstrate that Sar1A, Smad2, TNF-α, and IL-8 are targets of seq-915_x4024. Overexpression of seq-915_x4024 in KCs suppresses the inflammatory response and scar formation by targeting the TGF-β1-Smad signaling pathway and the inflammatory factors TNF-α and IL-8.

RESULTS

**Seq-915_x4024 Regulates Adult KC Biological Behavior**

To determine the effect of seq-915_x4024 on regulating the biological behavior of KCs, we transfected an HaCaT cell line with seq-915_x4024 mimics or anti-seq-915_x4024 and their respective negative controls (NCs). Transfection efficiency is shown in Figures 1A–1C. The percentage of FAM-positive HaCaT cells was 94.96% ± 3.76%. Furthermore, we estimated the effects of seq-915_x4024 on the proliferation and migration abilities of HaCaT using the MTS proliferation assay and Transwell cell migration assay. The data demonstrated that seq-915_x4024 exhibits a significant promotional effect on the growth of HaCaT cells (Figure 1D). The results of the Transwell cell migration assay showed that overexpression of seq-915_x4024 obviously inhibits the migration of HaCaT cells.
Significantly fewer seq-915_x4024-transfected HaCaT cells (18 ± 7, p < 0.05) passed through the membrane than NC-transfected HaCaT cells (37 ± 7) or parental HaCaT cells (38 ± 7) (Figures 1E and 1F).

**KCs with Overexpressed Seq-915_x4024 Promote FB Migration and FB Proliferation**

To investigate the effect of seq-915_x4024 in KCs on FB migration, an *in vitro* wound-healing assay was used. The data demonstrated that, after being cocultured with seq-915_x4024-transfected HaCaT cells, FBs showed a significant increase in cell migration ability (Figures 2A and 2B). In addition, HaCaT cells transfected with anti-seq-915_x4024 significantly inhibited the migration ability of FBs (Figures S1A and S1B).

We used a Transwell cell migration assay to further confirm that the overexpression of seq-915_x4024 in KCs promotes FB migration. The data showed that significantly more FBs cocultured with seq-915_x4024-transfected HaCaT cells (58 ± 11, p < 0.01) passed through the membrane than FBs cocultured with NC-transfected HaCaT cells (29 ± 7) and parental HaCaT cells (32 ± 20; Figures 2C and 2D). In contrast, significantly fewer FBs cocultured with anti-seq-915_x4024-transfected HaCaT cells (16 ± 4, p < 0.01) passed through the membrane than FBs cocultured with NC-transfected HaCaT cells (28 ± 5) and parental HaCaT cells (31 ± 7; Figures S1C and S1D). An MTS proliferation assay was used to detect the effect of seq-915_x4024 in KCs on FB proliferation and demonstrated that FBs cocultured with HaCaT cells showed a positive correlation between the expression of seq-915_x4024 in KCs and FB proliferation ability (Figures 2E and S1E).

**Seq-915_x4024 Regulates ECM and TGF-β Isoforms**

We detected the mRNA and protein levels of the ECM and TGF-β isoforms by real-time qRT-PCR and western blotting. After being transfected with seq-915_x4024 mimics, HaCaT cells expressed more Has3 and Col III compared with cells transfected with NCs (Figures 3A and 3B). In addition, an increased ratio (2.45 ± 0.57 of mRNA and 1.79 ± 0.12 of protein) of type III versus type I collagen was found in seq-915_x4024-transfected HaCaT cells (Figure 3C). FBs cocultured with seq-915_x4024-transfected HaCaT cells expressed less TGF-β1 and p-Smad3 and more TGF-β3 than cells cocultured with NC-transfected or parental HaCaT cells (Figures 3D and 3E).

**Seq-915_x4024 Suppresses the Inflammatory Response in KCs**

The addition of TNF-α to HaCaT cells was used to reflect inflammatory conditions. HaCaT cells were transfected with seq-915_x4024 mimics or NCs, and then inflammation was induced using TNF-α (20 ng/mL) for 24 h. The expression levels of IL-6, IL-8, CXCL1, and CXCL5 were significantly lower in seq-915_x4024-transfected HaCaT cells compared with levels in NC-transfected HaCaT cells, as detected by real-time qRT-PCR (Figure 3F). Western blot analysis showed that the expression of IL-6, IL-8, and TNF-α was significantly suppressed in seq-915_x4024-transfected HaCaT cells compared with those of NC-transfected HaCaT cells (Figure 3G).

**Seq-915_x4024 Suppresses Inflammatory Cell Infiltration and Scar Formation In Vivo**

After transplanting the KC cell sheets onto the surface of wounds, we found that overexpression of seq-915_x4024 in KC cell sheets had no significant effect on accelerating wound closure (Figures 4A and 4B). However, on days 15, 35, 45, and 60 after healing, the areas of scar tissue in the seq-915_x4024 group were significantly smaller than those of the NC group. On day 60 after full healing, the scars in the seq-915_x4024 group were 37% ± 12% smaller than those of the control group (Figures 4C and 4D). On day 28 after transplantation, there were fewer α-SMA (α-smooth muscle actin; marker of myofibroblasts)-positive cells, more-organized tissues (skin appendages and new vessels), and uniformly arranged collagen bundles in the deep dermis of the seq-915_x4024 group, than in those of the NC group (Figures 4E–4G and 5A). The wounds on day 12 detected by Masson’s Trichrome staining demonstrated that the closure of open wounds in seq-915_x4024 group rats occurred with deposition of more uniformly organized collagen fiber bundles (Figure 5A). In addition, on days 1, 3, and 6 after transplantation, the inflammatory cell infiltration at the wound sites of the seq-915_x4024 group was significantly lower than that of the NC group (Figures 5B and 5C). At the wound sites on day 28 after transplantation, there were significantly fewer macrophages in the seq-915_x4024 group than in the NC group (Figure 5D).

**Seq-915_x4024 Suppresses the Expression of Col I, Col III, p-Smad2, and p-Smad3 In Vivo**

On days 3, 6, 9, and 12 after transplantation, the wound tissues were harvested and analyzed using real-time qRT-PCR and western blot. Compared with the NC group, the protein expression of Col I and Col III and the mRNA expression of Col I were decreased significantly in the wounds of the seq-915_x4024 group (Figures 6A and 6B). On days 3 and 6 after transplantation, the mRNA expression of Col III in the wounds of the seq-915_x4024 group was higher than that in the NC group. However, on day 12, the mRNA expression of Col III in the NC group was significantly increased and higher than that in the seq-915_x4024 group.
the seq-915_x4024 group (Figure 6C). Smad3 protein expression and mRNA expression of Smad2 and Smad3 were not significantly different between the seq-915_x4024 group and NC group (Figures 6D–6F). However, Smad2, p-Smad2, and p-Smad3 protein expression was significantly decreased in wounds of the seq-915_x4024 group compared with those of the NC group (Figures 6F and 6G).

Figure 3. Seq-915_x4024 Regulates the ECM, TGF-β Isoforms, and Inflammatory Factors
(A) Real-time qRT-PCR demonstrated that the RNA levels of Col III and Has3 are significantly higher in seq-915_x4024-transfected HaCaT cells than in NC-transfected HaCaT cells. The expression of Col I showed no significant difference. (B) Western blotting confirmed that the expression levels of Col III and Has3 proteins were significantly higher in seq-915_x4024-transfected HaCaT cells compared with those in NC-transfected HaCaT cells. The expression of Col I showed no significant difference. (C) Realtime qRT-PCR and western blotting demonstrated that the ratio of type III versus type I collagen (2.45 ± 0.57 mRNA and 1.79 ± 0.12 protein) was significantly higher in seq-915_x4024-transfected HaCaT cells compared with that in NC-transfected HaCaT cells. (D) Western blotting demonstrated that FBs cocultured with seq-915_x4024-transfected KCs expressed less TGF-β1 and p-Smad3 and more TGF-β3 compared with FBs cocultured with NC-transfected KCs. (E) Real-time qRT-PCR demonstrated that FBs cocultured with seq-915_x4024-transfected KCs expressed less TGF-β1 and more TGF-β3 compared with FBs cocultured with NC-transfected KCs. (F) Real-time qRT-PCR was used to detect the RNA levels of IL-1α, IL-1β, IL-6, TNF-α, CXCL1, and CXCL5. The RNA levels of IL-6, IL-8, CXCL1, and CXCL5 were significantly lower in seq-915_x4024-transfected HaCaT cells than in NC-transfected HaCaT cells under inflammatory stimulation. (G) Western blotting showed that under inflammatory stimulation, the expression levels of IL-6, IL-8, and TNF-α in seq-915_x4024-transfected HaCaT cells were significantly lower than those in NC-transfected HaCaT cells. The results represent the means of the values. Bars indicate SD. *p < 0.05, statistical significance between groups.

Seq-915_x4024 Targets Smad2, Sar1A, TNF-α, and IL-8
Using bioinformatics analyses, we found that TGF-β2, TGF-β3, Smad2, Smad3, Smad4, Sar1A, TNF-α, and IL-8 may be potential targets of seq-915_x4024 (Figure 7A). To confirm whether these genes are direct targets of seq-915_x4024, we performed a luciferase reporter assay. The results showed that the luciferase activity of...
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pGL3-Sar1A-3′-UTR, pGL3-Smad2-3′-UTR, pGL3-TNF-α-3′-UTR, and pGL3-IL-8-3′-UTR reporters was significantly suppressed in seq-915_x4024-transfected HaCaT cells compared with the activity in NC-transfected HaCaT cells (Figures 7B and 7C). In contrast, there were no significant differences in the relative luciferase activity of pGL3-Smad2-3-MUT, pGL3-Sar1A-MUT, pGL3-TNF-α-MUT, or pGL3-IL-8-MUT reporters in seq-915_x4024-transfected HaCaT cells compared with activity in NC-transfected HaCaT cells (Figure 7C).

To further verify these results, we examined the Sar1A, Smad2, TNF-α, and IL-8 protein levels in seq-915_x4024-transfected, NC-transfected, and parental HaCaT cells by western blot. Normalized to the endogenous reference GAPDH, the levels of the endogenous Sar1A, Smad2, TNF-α, and IL-8 showed a clear reduction in seq-915_x4024-transfected HaCaT cells (Figures 7D and 7E). These results demonstrated that seq-915_x4024 may target Sar1A, Smad2, TNF-α, and IL-8 in KCs.

**DISCUSSION**

It is beginning to be revealed that miRNAs play important roles in skin regeneration. In early to mid-gestation, several miRNAs, especially skin-specific expressed miRNAs, contribute to skin regeneration without scar formation by targeting key genes and pathways. Using next-generation sequencing and mirTools 2.0, we identified a novel miRNA candidate, seq-915_x4024, which was highly expressed in mid-gestational fetal KCs. In this study, we further investigated the role of seq-915_x4024 in the process of wound healing.

Early in gestation, wounds heal rapidly. In addition, fetal KCs exhibit a significantly increased proliferation rate and promote the migration and proliferation ability of FBs compared with late-gestational fetal KCs. Are these fetal KC cell activities associated with overexpression of seq-915_x4024? We demonstrated that KCs with overexpression of seq-915_x4024 exhibit the same biological behaviors as fetal KCs at mid-gestation. After being transfected with seq-915_x4024 mimics, HaCaT cells exhibited higher proliferative activity and the ability to promote FB migration and FB proliferation.

To investigate whether overexpression of seq-915_x4024 in KCs plays a role in the process of wound repair, an *in vivo* wound-healing model was used. We demonstrated that wounds in the seq-915_x4024 group heal with fewer myofibroblasts, less scar tissue, uniformly arranged collagen bundles in the deep dermis, more-organized tissue, and less expression of Col I and Col III. Collagens are the most important components of dermal ECM. The key features of fibrosis are increased collagen content, and the arrangement of the collagen fibers becomes disordered. The accumulation of myofibroblasts is also involved in the process of fibrosis because myofibroblasts produce a collagen-rich matrix that forms scar tissues. This evidence demonstrates that seq-915_x4024 significantly inhibits FB conversion into myofibroblasts and scar formation.

Fetal skin synthesizes less Col I and more Col III than adult skin. Compared with type I collagen fibers, type III collagen fibers are smaller and may be deposited in a well-structured reticular manner that is similar to the structure of the fibers present in normal skin. In this study, an increased ratio of type III versus type I collagen was found in seq-915_x4024-transfected HaCaT cells, indicating that seq-915_x4024 contributes to the collagen composition, which leads to skin regeneration. In addition, we found that Has3 was also upregulated by seq-915_x4024. Has3, one of the three types of Has, synthesizes hyaluronic acid, which plays major roles in scarless wound healing. In dermal wounds receiving hyaluronic acid treatment, scar formation is significantly reduced. These results suggest that seq-915_x4024 regulates ECM components, including Col I, Col III, and Has3, which may contribute to skin regeneration.

Inflammation is a key determinant of fibrosis. The inflammatory phase is the beginning of wound healing. During this phase, chemokines and cytokines, such as platelet-derived growth factor (PDGF), interleukins, TGF-β, and TNF-α, are released into the bloodstream and activate inflammatory cells. Early in gestation, wounds in fetal mammalian skin do not trigger inflammation in response to damage and heal without scarring. In contrast, fetal wounds generated during late gestation heal with a strong inflammatory response and scar formation.

Chemokines and cytokines play important roles in scarless wound healing in fetal skin. Inflammatory cells exist and respond to inflammatory factors, even at the stage when fetal cutaneous wounds heal without an inflammatory response. TNF-α is an important pro-inflammatory cytokine. After wounding, in response to TNF-α, CXC family members, particularly CXCLs 1, 5, and 8, are released.
Figure 5. KCs with Overexpressed Seq-915_x4024 Suppress Inflammatory Cell Infiltration and Scar Formation In Vivo

(A) On days 12 and 28 after transplantation, Masson’s Trichrome staining showing uniformly arranged collagen bundles in the deep dermis of the seq-915_x4024 group.

(B) On days 1, 3, and 6 after transplantation, the tissues at the wound site were stained with H&E. The inflammatory cells were stained with blue nuclei, as indicated by black lines.
to recruit neutrophils into inflamed tissues. To investigate whether overexpression of seq-915_x4024 in KCs plays an anti-inflammatory role in wound healing, we used TNF-α as an inflammatory stimulus to demonstrate that the expression levels of IL-6, IL-8, CXCL1, CXCL5, and TNF-α are significantly suppressed in KCs transfected with seq-915_x4024. TNF-α and IL-8 are direct targets of seq-915_x4024 in KCs. IL-6 and IL-8 (pro-inflammatory cytokines) are expressed at low levels in fetal wounds. Attenuating IL-6 and IL-8 expression leads to reduced recruitment of inflammatory cells and decreased scar formation. Using in vivo experiments, we demonstrated that KC cell sheets with overexpressed seq-915_x4024 suppress inflammatory cell infiltration. We believe that seq-915_x4024 also functions as an anti-inflammatory factor and contributes to skin regeneration by suppressing the pro-inflammatory cytokines TNF-α, IL-6, and IL-8 and the pro-inflammatory chemokines CXCL1 and CXCL5.

Among all the growth factors involved in wound healing, TGF-β family members are the most important factors that play roles in all three phases, especially in the remodeling phase. The mammalian genome contains three TGF-β isoforms known as TGF-β1, TGF-β2, and TGF-β3. TGF-β1 is well known as a key mediator in tissue fibrosis via the Smad signaling pathway. The activated TGF-β1-Smad signaling pathway induces myofibroblastic differentiation, induces FB production of type I collagen, and promotes ECM deposition. In contrast, TGF-β3 appears to play different roles in wound healing, as it inhibits myofibroblast proliferation and type I collagen deposition, which leads to scar tissue formation. Overexpression
of seq-915_x4024 leading to higher TGF-β3 and lower TGF-β1 may contribute to suppressing fibrosis.

What is the molecular mechanism by which seq-915_x4024 acts as an anti-fibrotic factor? TGF-β1 plays a central role in tissue fibrosis via the Smad signaling pathway. TGF-β1 interacting with type I and type II transmembrane receptors (TGF-βRI and TGF-βRII, respectively) activates canonical Smad pathway members, including Smad2 and Smad3, forming hetero-oligomeric complexes with Smad4 (common Smads, co-Smads), and finally translocating into the nucleus and activating transcription. We investigated whether TGF-β1-Smad signaling pathway members are targets of seq-915_x4024 and demonstrated that Smad2 and Smad3 are direct targets. In addition, the expression of TGF-β1 significantly decreased by seq-915_x4024 in vivo. These results demonstrated that seq-915_x4024 suppresses the TGF-β1-Smad signaling pathway by targeting Smad2 and Smad3. Seq-915_x4024 acts as an anti-fibrotic factor by suppressing the TGF-β1-Smad signaling pathway members Smad2 and Smad3.

In conclusion, our study shows that the overexpression of seq-915_x4024 in KCs suppresses inflammatory cell infiltration and scar formation by targeting the TGF-β1-Smad signaling pathway and inflammatory factors TNF-α and IL-8. Seq-915_x4024 could be a new anti-fibrotic factor for the treatment of wound healing.

MATERIALS AND METHODS

Skin Samples and Cell Culture

The immortalized human KC cell line HaCaT was kindly provided by Chundi He and propagated in high-glucose Dulbecco’s modified Eagle’s medium (DMEM [HG]; Invitrogen Life Technologies, Carlsbad, CA, USA). Skin samples were obtained from First Hospital of China Medical University. This study was performed in accordance with the ARRIVE guidelines and under a research protocol approved by the Ethics Committee of China Medical University. Written informed consent was obtained.

Primary culture of FBs was obtained from skin samples. Cells were prepared as previously described. Briefly, full-thickness skin samples were de-epithelialized after incubating at 4°C overnight in Dispase II (Roche Applied Science, Indianapolis, IN, USA). After mincing the dermal components, the isolated FBs were cultured in DMEM (HG) supplemented with 10% FBS (Invitrogen Life Technologies), 100 U/mL penicillin and 100 μg/mL streptomycin. The cultures were incubated at 37°C in a humidified incubator with 5% CO2. When the FBs reached 80–90% confluence, the cells were passaged for expansion. For the experiments, human dermal FBs were used at passages 3–6.

RNA Isolation and Real-Time qRT-PCR

Total RNA was extracted using the mirVana miRNA Isolation Kit (Thermo Fisher, Uppsala, Sweden) according to the manufacturer’s instructions. The concentration and purity of RNA were determined by ultraviolet spectrophotometry (A260/A280 > 1.9), using a nanophotometer UV-visible spectrophotometer (Implen, Schatzbogen, Germany). The 3’-termini of the RNA sequences were polyadenylated using the Poly(A) Tailing Kit (Thermo Fisher). Then, RNA was extracted with phenol-chloroform and precipitated with ethanol.

Real-time qRT-PCR was used to confirm the expression levels of seq-915_x4024, IL-1α, IL-1β, IL-6, IL-8, TNF-α, CXCL1, CXCL5, TGF-β1, TGF-β3, Has3, Col I, and Col III. Reverse transcription was performed using a PrimeScript RT reagent Kit with gDNA Eraser (Takara Bio, Shiga, Japan). Real-time qRT-PCR was performed on a 7500 Real-Time PCR system (Applied Biosystems, Foster City, CA, USA) supplied with analytical software, using an SYBR Premix Ex Taq II Kit (Takara Bio) according to the manufacturer’s instructions. The U6 or GAPDH mRNA level, as an endogenous reference, was used for normalization. The relative expression levels of the genes were calculated using the 2^{-ΔΔCT} equation in which ΔΔCT = CT_T mRNA – C_T U6 or ΔΔCT = CT_T mRNA – CT_GAPDH. The primers used for real-time qRT-PCR are given in Table S1.

Transfection

All RNA oligoribonucleotides and their respective NCs were purchased from GenePharma (Shanghai, China). The pyrimidine nucleotides in the seq-915_x4024 mimics, anti-seq-915_x4024, and their respective NCs were substituted with their 2-O-methyl analogs to improve RNA stability. HaCaT cells were plated 1 day before transfection. RNA oligoribonucleotides were transfected using Lipofectamine 2000 (Thermo Fisher) according to the manufacturer’s instructions. After 48 h, the cells were harvested for further experiments.

Coculture of Dermal FBs with HaCaT

Dermal FBs were cocultured with parental and transfected HaCaT cells in Transwell chambers (3-μm pore size; Corning, Corning, NY, USA). After 96 h, FBs were harvested for further experiments.
Transwell Cell Migration Assay
We used a Transwell cell migration assay to estimate the migration ability of the cells in vitro. Before use, the bottom of the culture inserts (8-μm pores) in 24-well tissue culture plates (Transwell; Corning) was coated with serum-free medium at 37°C for 1 h. A total of 5 × 10^4 cells were harvested by trypsinization, washed with serum-free medium, resuspended at a concentration of 5 × 10^5/mL, and placed in the upper chamber. The lower chamber contained 10% FBS for use as a chemoattractant. After 24 h of incubation at 37°C with 5% CO2, the number of cells that had migrated to the basal side of the membrane was quantified by counting 16 independent symmetrical visual fields under the microscope, and cell morphology was observed by staining with H&E (hematoxylin-eosin).

Quantification of Cell Proliferation
Cell proliferation was determined using a colorimetric microculture assay with MTS (3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulphophenyl)-2H-tetrazolium) dye. A total of 5 × 10^3 HaCaT cells or 2 × 10^3 FBs were seeded separately into 96-well culture plates for 24, 48, 72, and 96 h; after incubation with 20 μL MTS for 2 h at 37°C, the absorbance was measured using an iMARK microplate reader (Bio-Rad, Hercules, CA, USA) at a wavelength of 495 nm.

In Vitro Wound-Healing Assay
For the migration assay, culture inserts (Ibidi, Munich, Germany) were placed into the wells of a six-well plate. After being cocultured, 5 × 10^3 FBs with 100 μL medium were seeded on each side of a culture insert and incubated under normal culture conditions to allow the cells to grow to confluence. Afterward, inserts were removed, leaving a cell-free area. The medium was switched to 2 mL DMEM (HG) supplemented with 1% FBS. The status of the scratch wounds was monitored using an inverted microscope at 0, 12, 24, and 48 h, and representative images were collected. The results are presented as the percentage of wound healing, calculated as follows: (wound area (initial) − wound area (final))/wound area (initial) × 100.

KC-Sheet Preparation and Wound-Healing Model In Vivo
HaCaT cells (5 × 10^5) were seeded into Nunc UpCell (Thermo Fisher) dishes and incubated under normal culture conditions to allow cells to grow to confluence. After 24 h, the dishes were transferred to another incubator, set at 20°C with 5% CO2, for 4 h to release the cultured cells as a KC cell sheet. In vivo assessment of the effect of seq-915_x4024 on cutaneous regeneration was performed using a wound-healing model. Female Wistar rats (10–12 weeks old, approximately 250 g) were obtained from Vital River Laboratory Animal Technology (Beijing, China) and housed in the animal care facilities of China Medical University under specific-pathogen-free conditions. After hair removal from the dorsal surface, the rats were anesthetized with pentobarbital sodium (40 mg/kg; Solarbio, China). Full-thickness skin, 10 mm in diameter, was removed from each side of the dorsal skin of each rat. The seq-915_x4024-transfected KC cell sheet was transplanted onto the surface of the left-hand wound bed, and the NC-transfected KC cell sheet was transplanted on the other side as an NC, followed by coating with Hydrosorb (Hartmann, Germany), Tegaderm film dressing (3M Health Care, Neuss, Germany), and medical gauze. After rats were sacrificed, the wound tissues, including a border of normal tissue, were harvested and immediately fixed in 4% paraformaldehyde in phosphate-buffered saline and were dehydrated in a 30% sucrose solution. Frozen sections were prepared in a freezing microtome (Leica, Germany) and subjected to the following procedures: (1) H&E staining for histological evaluation; (2) Masson’s trichrome staining (Solarbio, Beijing, China) for collagen fibers; (3) immunohistochemical assay for the expression of α-SMA (α-smooth muscle actin); and (4) immunofluorescence assay for expression of Ly6G and F4/80.

Bioinformatics Analyses and Luciferase Reporter Assays
For potential target gene prediction of seq-915_x4024, the 3' UTRs of the genes were obtained from both the Ensembl and miRanda databases. We considered the seed sequence (nucleotides 2–8) of seq-915_x4024 and at least one part of the 3' UTR of target mRNAs was reverse complemented perfectly.10 For luciferase reporter experiments, the wild-type 3' UTR segments of Sar1A, Smad2, Smad3, Smad4, TGF-β2, TGF-β3, TNF-α, and IL-8, containing the seq-915_x4024 binding sites, were amplified by PCR and inserted into the pGLO-control vector (Promega, Madison, WI, USA) using the XbaI site, which is immediately downstream of the luciferase stop codon. DNA segments with scrambled target sites (Smad2-3-MUT, Sar1A-MUT, TNF-α-MUT, and IL-8-MUT) designed to interfere with seed sequence recognition were also cloned to serve as controls. HaCaT cells were plated in 24-well plates. For each well, 20 pM seq-915_x4024 mimics or NC, 0.8 μg of the firefly luciferase report vector and 0.08 μg of the control vector containing Renilla luciferase, pRL-TK (Promega), were transfected using Lipofectamine 2000. Twenty-four hours after transfection, Firefly and Renilla luciferase activities were measured consecutively using a dual-luciferase reporter assay (Promega) on a Centro LB 960 (Berthold, Bad Wildbad, Germany).

Western Blot Analysis
Western blotting was performed as previously described.13 Briefly, total protein was extracted using a Total Protein Extraction Kit (KeyGen, Nanjing, China), followed by measurement using a BCA Protein Assay Kit (KeyGen). A total of 30 μg of each sample was separated by 12% SDS polyacrylamide gels and electrophoretically transferred to polyvinylidene difluoride membranes (Millipore, Billerica, MA, USA). After incubation with 5% bovine serum albumin in Tris-buffered saline-0.5% Tween-20 at room temperature for 1 h, the membranes were incubated at 4°C overnight with primary antibodies against IL-6 (ImmunoWay, Plano, TX, USA), IL-8 (ImmunoWay), TNF-α (ImmunoWay), Sar1A (ImmunoWay), Smad2 (ImmunoWay), TGF-β1 (Abcam, Cambridge, MA, USA), TGF-β3 (Santa Cruz Biotechnology, Santa Cruz, CA, USA), HAS3 (Abcam), Col III (ImmunoWay), and GAPDH (Santa Cruz) and then incubated with horseradish peroxidase-conjugated secondary antibody at room temperature for 2 h. The antigen-antibody
complexes were visualized, using an ECL Kit (Pierce, Rockford, IL, USA). Quantification of protein was carried out using FluorChem 2.01 (Alpha Innotech, San Leandro, CA, USA). Protein levels in seq-915_x4024-transfected HaCaT cells were presented as fold change relative to NC-transfected HaCaT cells.

**Statistical Analysis**

The results are presented as the mean ± SD from at least three separate experiments. Statistical differences between groups were analyzed using a one-way ANOVA test or Student’s t test. The statistical analyses were performed using SPSS 16.0 software (SPSS, Chicago, IL, USA). Values of p less than 0.05 were considered statistically significant.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes one figure and one table and can be found with this article online at https://doi.org/10.1016/j.omtn.2018.12.016.

**AUTHOR CONTRIBUTIONS**

F.Z. and H.L. performed the experiments; F.Z. wrote the paper; Z.W., X.L., and D.Z. provided study materials; T.Z., R.W., and X.L. contributed to data analysis and interpretation; P.S. contributed to collection and assembly of data; and X.P. designed the study and provided final approval of the manuscript.

**CONFLICTS OF INTEREST**

The authors declare no competing interests.

**ACKNOWLEDGMENTS**

This work was supported by the Program for National Basic Research Program of China (2012CB518103), the National Science Foundation of China (81370883), the National Key Research and Development Project (2017YFC1103300), the Shenyang Key Laboratory Project of China (81370883), the National Key Research and Development Program of China (2012CB518103), the National Science Foundation of China (F15-157-1-00), the Key R & D and Technology Transfer Program (2017TFC1103300), and the Program for National Basic Research (2018.12.016). The authors thank the laboratory staff for their dedicated assistance.

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