SDF-1-edited human amniotic mesenchymal stem cells stimulate angiogenesis in treating hindlimb ischaemia

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Abstract
Although stem cells have extensively been studied as a novel vehicle for tissue repair, their sustained efficacy remains controversial. In this study, we aimed to investigate the angiogenic potency over time of stromal cell-derived factor-1 (SDF-1) gene-edited amniotic mesenchymal stem cells (AMM/S) in a hindlimb ischaemia model. An SDF-1 transgene was inserted into the AMM cell genome via transcription activator-like effector nuclease (TALEN) mediated knock-in, and cell migration, Matrigel tube formation, and in vivo Matrigel plug assays were performed. AMM/S were also transplanted into hindlimb ischaemia model mice. Blood perfusion, therapeutic potential, histology, capillary density and in vivo angiogenic assays were performed. AMM/S exhibited high expression of the SDF-1 gene, and robustly promoted migration, proliferation and microvascular formation. AMM/S transplantation significantly increased blood perfusion and limb loss prevention compared with AMM. AMM/S also significantly inhibited increased capillary density and expression of angiogenic factors in the ischaemic hindlimb. Our study demonstrated that AMM/S provides a significant therapeutic effect in ischaemic hindlimbs by enhancing angiogenesis.

KEYWORDS
angiogenesis, cell therapy, ischaemia, SDF-1, stem cells

1 | INTRODUCTION
Peripheral artery disease (PAD) is a vascular disease caused by abnormal narrowing of peripheral arteries, restricting blood flow to peripheral tissues, causing pain, ulceration and gangrene. The PAD patient population is estimated to include more than 200 million worldwide. These patients’ symptoms usually progress to critical limb ischaemia (CLI) eventually requiring limb amputation.1,2 Treatment options for CLI patients are limited primarily to surgical and endovascular procedures. Novel, more effective treatment strategies, including revascularization or therapeutic angiogenesis, are being actively pursued.

Recently, stem and/or progenitor cells have been shown to contribute to therapeutic angiogenesis through paracrine effects (transdifferentiation).3,4 Several studies have demonstrated that mesenchymal stem cells (MSCs) can robustly stimulate angiogenesis by secreting angiogenic factors, facilitating favourable therapeutic outcomes.5,6 However, the remaining uncertainty in therapeutic outcomes of procedures using stem cells is a major obstacle to their use for damaged tissue regeneration. One interesting angiogenic
factor, stromal cell-derived factor-1 (SDF-1), also known as C-X-C motif chemokine 12, has been reported to exert potent angiogenic effects. SDF-1 signals to fibroblasts, osteoblasts and endothelial cells, and plays a key role in cell trafficking and homing of hematopoietic progenitor cells. SDF-1 overexpressing rat MSCs promote angiogenesis in a rat model of myocardial infarction. Thus, SDF-1 introduction by tissue engineering or transplantation of SDF-1-overexpressing cells has been successfully applied to regenerate damaged tissues and organs.

Recently, genome-editing technologies have been applied to generate functionally improved stem cells. Such technologies can target specific genomic sites, and artificial restriction enzymes allow genome modification via homology-directed repair for knock-in construction.

In this study, we generated long-term SDF-1 secreting human amniotic mesenchymal stem cells (AMM/S) via TALEN-mediated editing and investigated to determine whether gene-edited AMM/S are novel angio-vasculogenic cell population in experimental ischaemia.

2 | MATERIALS AND METHODS
2.1 | Cell culture

Human AMMs were obtained from Thermo Fisher Scientific, Inc. and cultured in low-glucose Dulbecco’s modified Eagle medium (DMEM), supplemented with 10% foetal bovine serum (FBS), 100 U/ml penicillin and 100 μg/ml streptomycin (Gibco). The characteristics of AMMs were described in Figure S1.

2.2 | Donor construction

SDF-1 was synthesized and inserted into the AAVS1 genome site targeting donor vector (System Biosciences).

2.3 | Transfection and selection

Transfection and selection were conducted as described in a previous study. AMMs (1 × 10⁵) were suspended with 0.6 μg of AAVS1 left, right TALE-Nuclease vector (System Biosciences) and AAVS1 HR Donor (System Biosciences) in 10 μl electroporation buffer, and electroporated using a Neon Transfection System (Thermo Fisher Scientific). Four days after, SDF-1 knock-in cells were selected by incubating them in 5 μg/ml puromycin for 7 days.

2.4 | Fluorescence-activated cell sorting

After puromycin selection, the cells were washed once with phosphate-buffered saline (PBS) following the 0.05% trypsin/EDTA treatment for cell detachment. The cells were resuspended in PBS for sorting. AMM/S were sorted on FACSaria™ III Cell Sorter (BD Biosciences).

2.5 | Genomic DNA extraction and junction polymerase chain reaction

Genomic DNA from cells was extracted using a G-spin™ Total DNA Extraction Mini Kit (Intron Biotechnology), and 120ng of genomic DNA was amplified by touch-down polymerase chain reaction (PCR) (36 cycles) as described in a previous study. PCR primer was used as following: Forward (J212): aactctgcccttaaacgctg, (J195): cgggctggctcaggttctgtgta.

2.6 | Quantitative reverse transcriptase-PCR (qRT-PCR) and reverse transcriptase-PCR (RT-PCR) analyses

qRT-PCR or RT-PCR analysis was conducted as described in a previous study. Briefly, the total RNA was isolated from cells using RNA-stat (Iso-Tex Diagnostics), and RNA was reverse-transcribed using Taqman Reverse Transcription Reagents (Applied Biosystems). The synthesized cDNA was subjected to qRT-PCR using human-specific primers and probes. RNA levels were quantitatively assessed using the ABI PRISM 7000 Sequence Detection System (Applied Biosystems). The relative mRNA expression normalized to GAPDH expression was calculated as described previously.

2.7 | qPCR primer

The primers used in qRT-PCR were for human SDF-1 (Hs00171022_m1) and GAPDH (Hs99999905_m1) and for mouse fibroblast growth factor-2 (FGF-2) (Mm01285715_m1), hepatocyte growth factor (HGF) (Mm01135184_m1), insulin growth factor-1 (IGF-1) (Mm00439561_m1), vascular endothelial growth factor A (VEGF-A) (Mm01281448_g1) and GAPDH (Mm99999915_g1). All of the primer/probe sets were purchased from Applied Biosystems.

2.8 | Conditioned media preparation

Conditioned media (CM) was prepared as described in previous literature. Briefly, AMMs, AMM/sh-S or AMM/S (each 1 × 10⁶) were seeded into T-75 cell culture flasks and grown in normal medium or low-glucose DMEM (Gibco) containing 10% FBS, 100 μg/ml penicillin and 100mg/ml streptomycin (Gibco) for 48h until the cells reached approximately 90% confluence. Each of CM was then centrifuged, and the supernatants were collected and used for this study.
2.9 | Scratch migration assay

Scratch wound assays were performed by a method established in previous literature with modifications. Human dermal fibroblast (1 x 10^5) were seeded in 24-well culture plates coated with type I collagen (0.2 mg/ml) and incubated at 37°C in 5% CO₂ to produce confluent monolayers. Monolayers were scratched using a sterile pipette tip and incubated with each CM groups (AMMs, AMM/sh-S or AMM/S), which had been cultured for 5 days. To measure cell mobility, images were obtained at 5 random fields after scratching. Wound areas were examined, using the NIH Image program (http://rsb.info.nih.gov/nih-image/).19

2.10 | Matrigel tube formation assay

Each type of AMMs, AMM/sh-S or AMM/S was seeded with endothelial growth medium (EGM)-2 (Lonza Inc.) at a concentration of 1.5 x 10^6 cells/well in basement membrane matrix gel (Matrigel, BD)-coated 12-well culture plate. After 12h of incubation, culture plates were randomly photographed using microscopy. Tube length and the number of branching point from each sample were measured.

2.11 | In vivo Matrigel plug assay

AMMs or AMM/S with 500μl of Matrigel were subcutaneously transplanted in Male athymic nude mice (Joongang Laboratory Animal Inc.), aged between 7 and 9 weeks. After 2 weeks, Matrigel plugs were harvested, and the haemoglobin content was measured as previously reported.20

2.12 | Transplantation of cells in the ischaemic hind limb mouse model

All procedures were performed in accordance with the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (NIH Publication No. 85–23, revised 1996). Experimental protocols of animal were approved by the Catholic Kwandong University Institutional Animal Care and Use Committee. Female athymic nude mice (Joongang Laboratory Animal Inc., Seoul, South Korea) aged between 6 and 9 weeks and weighing between 19 and 22 g were used. To induce hind limb ischaemia, the mice were anaesthetised with 2% isoflurane, and their right femoral artery was surgically ligated. 1,1'-dioctadecyl-3,3',3''-tetramethylindocarbocyanine (Dil)-labelled 1 x 10^6 AMM, AMM/S or PBS were intramuscularly injected into the hind limb area after surgery (n = 7 for each group).21 Immediately before euthanasia, an overdose of pentobarbital (200 mg/kg) was injected into each mouse. We used a laser Doppler perfusion image (LDPI) analyser (Moor Instrument) to measure the serial blood flow in the hind limb after cell injection.

2.13 | Histological analysis

Mice hind limbs were harvested and fixed for 4 h in 4% paraformaldehyde and incubated overnight in 15% sucrose solution. The tissues were embedded in OCT compound (Sakura Finetek, Torrance, CA, USA), snap-frozen in liquid nitrogen and sectioned (thickness, 10–20 μm).22 Nuclei were counterstained with 4′,6-diamidino-2-phenylindole (1:5000; Sigma Aldrich). For capillary density measurement, five frozen sections of tissue from each group were stained with primary biotinylated ILB4 (1:250; Vector Laboratory Inc.) and secondary strepta-avidin Alexafluor 488 (1:400; Invitrogen).21 Five fields from four tissue sections were randomly selected, and the number of capillaries was counted in each field.21

Sections were also labelled with primary antibody to SDF-1 (Santa Cruz Biotechnology, Inc.) and detected by Cy3 conjugated antibody (Jackson Immunoresearch Laboratories). Photographs were taken using confocal microscopy (Zeiss LSM 510; Carl Zeiss Inc.).

2.14 | Statistical analysis

All data were presented as mean ± standard deviation. Statistical analyses were performed with Student’s t test for comparisons between two groups, and analysis of variance followed by Bonferroni’s correction was performed for more than two groups using SPSS version 11.0 (IBM Corp.).16,17 A p value <0.05 was considered statistically significant. Graphs were drawn using MedCalc software® (Mariakerke, Belgium).

3 | RESULTS

3.1 | Targeted SDF-1 knock-in into the AMMs cell genome

To generate a stable stem cell line expressing SDF-1 using the TALEN integration method, we chose a safe locus on chromosome 19 (AAVS1) as the target site. The donor plasmid was designed to carry the PGK promoter, SDF-1 and EF1α promotors, and GFP-T2A-puromycin (Figure 1A). The donor plasmid was transfected into human AMMs using TALEN. Only 4%–5% of cells were GFP-positive after transfection. Purity of SDF-1 knock-in cells was enriched to approximately 99.1% of GFP-positive cells by puromycin treatment followed by fluorescence-activated cell sorting (Figure 1B). To confirm integration of the donor plasmid into the AMM genome, we conducted genomic DNA PCR. Correct insertion of the donor plasmid was confirmed. (Figure 1C). Next, we confirmed SDF-1 expression in transfected AMMs by RT-qPCR. We found that transfected AMMs...
expressed high levels of SDF-1 compared to control AMMs, suggesting successful establishment of an SDF-1 expressing AMM/S cell line (Figure 1D).

3.2 | Culture Media (CM) from AMM/S promotes endothelial cell proliferation and migration

To examine whether protein factors secreted from AMM/S promote endothelial cell proliferation and migration, we measured rates of cell proliferation, and performed a scratch migration assay. In addition, to further confirm the function of SDF-1 in AMMs, we also generated SDF-1 silencing AMMs using shRNA lentiviral construct (AMM/sh-S) and AMD3100 (CXCR4 inhibitor) treated AMM/sh-S used as an another control group. The cell proliferation assay showed that AMM/S CM (333 ± 30%, p = 0.001, 0.002) significantly increased rates of HUVEC proliferation compared with control AMMs CM (206 ± 21%) or AMM/sh-S CM (188 ± 19%) (Figure 2A and B). Scratch closure assay also showed that AMM/S CM (66.3 ± 11%, p = 0.001, 0.002) significantly increased rates of wound closure compared with those under exposure to control AMMs (24.3 ± 4%) or AMM/sh-S CM (24.6 ± 5%) (Figure 2C and D).

3.3 | AMM/S display angiogenic and vasculogenic properties

To investigate vasculogenic potential of AMM/S in vitro, we performed a Matrigel tube formation assay. AMM/S displayed significantly higher tube lengths (8.9 ± 3 mm, p = 0.007, 0.006) and branching points (16.6 ± 4, p = 0.002, 0.002) than in AMMs (1.8 ± 0.3, 3.2 ± 0.2), or AMM/sh-S (1.8 ± 0.2, 2.7 ± 0.2) (Figure 3A and B).

To measure the vasculogenic potential of AMM/S in vivo, we performed a Matrigel plug assay. After 2 weeks, Matrigel plugs were harvested, and their haemoglobin content was examined. Interestingly, Matrigel plugs containing AMM/S (4.1 ± 1.5, p = 0.02)
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contained significantly higher levels of red blood cells compared with plugs containing AMMs (1.5 ± 0.6), indicating high potential for inducing formation of vasculature (Figure 3C and D).

3.4 | AMM/S transplantation exerts strong therapeutic effects in response to hindlimb ischaemia

We induced hind limb ischaemia in nude mice to examine the therapeutic effects of AMM/S. AMM/S, AMM or PBS were injected intramuscularly into the hind limb. Laser Doppler perfusion image analysis revealed that AMM/S (47 ± 7.1, p = 0.03, 0.0006) injection remarkably increased blood perfusion on Day 5 compared with limbs injected with AMM (32 ± 8.6) or PBS (12 ± 3.0) (Figure 4A and B). In addition, the AMM/S-injected group demonstrated a higher limb function salvage ratio than the AMM and PBS control groups (Figure 4C and D).

3.5 | AMM/S enhance capillary density in ischaemic hind limb muscle

To elucidate the mechanism(s) underlying the therapeutic effects of AMM/S treatment, we harvested hind limb muscle and immunostained tissue sections using SDF-1 and isoelectin B4 (ILB4) as a marker for endothelial cells. The number of ILB4 positive capillaries was significantly higher on Day 7 in the AMM/S-injected hind limb (67 ± 1.5, p = 0.04, 0.0003) than in PBS (17.3 ± 5.5) or AMM (44.3 ± 9.7) treated control hind limbs (Figure 5A and B). In addition,
immunostaining and Western blot analysis revealed that the expression of SDF-1 ($2.8 \pm 0.5, p = 0.001, 0.001$) was higher in AMM/S-injected hind limb than AMM or AMM/sh-S (1.0 ± 0.3) or AMM (0.9 ± 0.2) treated control hind limbs (Figure S2A,B).

To further elucidate the mechanisms underlying the enhanced recovery of blood perfusion, we examined the expression of angiogenic factors in hind limb tissues via RT-qPCR analysis. Interestingly, AMM/S injection significantly upregulated FGF-2 ($p < 0.01, n = 5$ per group), HGF ($p = 0.02, p = 0.0001$), VEGF-A ($p = 0.005, p = 0.007$) compared to AMM or PBS injection five days after cell transplantation, suggesting strong angiogenic stimulation by AMM/S, promoting vascular regeneration (Figure 5C).

### 3.6 Endothelial differentiation potential of AMM/S in vivo

The endothelial differentiation potential of AMM/S was investigated in hind limb ischaemia. Dil-labelled AMM/S were directly injected into the ischaemic area of the hind limb adductor muscle.
Hind limb muscle tissues were harvested 4 weeks after cell injection. Immunohistochemistry was performed, demonstrating that AMM/S colocalized with ILB4 in vascular structure, supporting their endothelial differentiation in vivo (Figure 6). However, we could not find colocalization with ILB4 of AMM in vascular structure.

4 | DISCUSSION

In this study, we first generated a gene-edited AMM/S cell line. We also investigated angio-vascuogenic characteristics of AMM/S and the possible therapeutic mechanism. These results showed that AMM/S injection induced the enhanced recovery of blood flow, neovascularization and prevented limb dysfunction in a mice limb ischaemia model.

Although stem cell-based cell therapies have emerged as an attractive therapeutic avenue for repair of damaged tissues, the lasting efficacy of its therapeutic effects remains in question.\textsuperscript{23} We initially evaluated allogeneic stem cell source candidates, including adult stem cells, for some important points, including low immunological response, young/unlimited cell proliferation, high transdifferentiation potential and easy accessibility for collection. We identified human amnion as an attractive stem cell resource. We evaluated isolated tissues from the amnion and found the amnion membrane to be an attractive candidate stem cell source. AMM contains abundant cells compared with fluid, cord or chorion, and displays high/unlimited cell proliferation capacity, as previously reported.\textsuperscript{24} Additionally, we tested a strategy to enhance the therapeutic potential of AMMs by genome modification using TALEN gene-editing methods. Gene-editing technology can precisely manipulate genes, preventing unintended mutations.

We recently reported angiogenic properties in human AMMs, and ASC overexpressing a single chemokine gene (GCP-1) displayed high angio-vasculogenic capacity.\textsuperscript{25} In addition, AMM offers great advantages as an allogeneic stem cell source due to its low immunogenicity, and unlimited cell proliferation capacity.\textsuperscript{24,26} Thus, we hypothesize that one of the representative chemokine genes (SDF-1) overexpressed in AMM might enhance their angiogenic potential, aiding development of universal stem cell resources. The marginal therapeutic effects of stem cells to date are a major limitation facing cell therapy. In addition, AMM specifically exhibits low SDF-1 expression compared to other angiogenic factors such as VEGF-A. In fact, SDF-1 overexpressing MSC promote angiogenesis and improve heart function.\textsuperscript{8} In line with this report, gene-edited AMM/S also exhibited strong proliferative and migratory properties. In addition, AMM/S also showed vasculogenic potential in in vitro and in vivo Matrigel assays.

CXC chemokines exert pleiotropic effects on immunity, angiogenesis and cancer metastasis.\textsuperscript{27} In addition, CXC chemokines are a unique family of cytokines that contain the 'ELR' motif and potently stimulate angiogenesis via binding and activating CXCR2 on endothelial cells.\textsuperscript{27} SDF-1, also known as C-X-C motif chemokine 12 (CXCL12), induces migration of human umbilical vein endothelial cells in vitro\textsuperscript{28} and
enhances formation of tube-like structures. Consistent with these studies, SDF-1 overexpressing AMM/S exhibited strong migration and tube formation properties, suggesting enhanced angiogenic potential.

SDF-1 exerts a chemoattractive effect on haematopoietic stem cells (HSCs) and induces bone marrow-derived c-kit progenitor cell differentiation towards an endothelial phenotype. SDF-1 pretreatment during endothelial progenitor cell (EPC) expansion stimulates EPC adhesion to endothelial cells, augmenting cell therapy efficacy for ischaemic vascular diseases. Based on these numerous reports, SDF-1 affect chemotaxis and differentiation in HSCs and EPCs, enhancing therapeutic angiogenesis in hind limb ischaemia.

In addition, MSCs could differentiate into endothelial cells in the presence of SDF-1. Consistent with these studies, we found that AMM/S transdifferentiated into endothelial-like cells with vascular structure, suggesting that autocrine SDF-1 action might enhance in vivo endothelial cell transdifferentiation in hind limb tissue.

Stem cell therapeutic mechanisms in ischaemic cardiovascular disease have mostly been attributed to paracrine effects. AMM/S cell injection significantly increased vascular density and resulted in high levels of the proangiogenic factors FGF-2, HGF, IGF-1 and VEGF-A in hind limb tissues. AMM/S injection may stimulate circulating endothelial stem or progenitor cells to home to ischaemic

**FIGURE 5** Therapeutic mechanism of hind limb ischaemia after cell injection. (A) Representative images of capillary density in ischaemic hind limbs 2 weeks after cell injection. (B) Quantification of capillary density in ischaemic hind limb tissues after cell injection. **p < 0.01; n = 7 per group. (C) Analysis of levels of angiogenic gene expression in ischaemic hind limb tissues 3 days after cell transplantation. Various angiogenic gene expression levels were measured by RT-qPCR. **p < 0.01; *p < 0.05; n = 7 per group.
limbs due to high chemokine levels. In addition, in vivo endothelial-transdifferentiated AMM/S contributed to neovascularization or maintenance of vascular structure in ischaemic limbs.

In conclusion, we demonstrated that SDF-1 overexpressing AMMs generated by gene-editing display potent angiogenic and vasculogenic properties. Thus, we speculate that AMM/S may represent a great therapeutic option, as an enhanced stem cell modality of allogeneic therapy in ischaemic vascular diseases.

**AUTHOR CONTRIBUTIONS**

**Hong Zhe Zhang:** Data curation (equal); funding acquisition (equal); investigation (equal). **Seongho Han:** Data curation (equal); investigation (equal); project administration (equal); resources (equal); supervision (equal). **Sung-Whan Kim:** Conceptualization (equal); data curation (equal); funding acquisition (equal); investigation (equal); methodology (equal); project administration (equal); resources (equal); supervision (equal); writing – original draft (equal); writing – review and editing (equal).

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**CONFLICT OF INTEREST**

The authors declare no conflicts of interest.

**DATA AVAILABILITY STATEMENT**

Research data are not shared.

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