Original Article

Exendin-4-assisted adipose derived mesenchymal stem cell therapy protects renal function against co-existing acute kidney ischemia-reperfusion injury and severe sepsis syndrome in rat

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Abstract: This study tested the hypothesis that combined therapy with exendin-4 (Ex4) and autologous adipose-derived mesenchymal stem cells (ADMSCs) was superior to either alone for protecting renal function against acute kidney ischemia-reperfusion (IR; 40-min ischemia/27-h reperfusion) injury when complicated by sepsis syndrome (SS; by cecal-ligation-puncture). Adult-male Sprague-Dawley rats (n=40) were equally divided into group 1 (sham-control), group 2 (IR-SS), group 3 (IR-SS + Ex4, 10 μg/kg subcutaneously 30 min after reperfusion and daily for 3 days), group 4 (IR-SS + ADMSC (1.2 × 10⁶)), and group 5 (IR-SS + Ex4 + ADMSC). The circulating levels of BUN and creatinine and the ratio of urine protein to creatinine were highest in group 2, lowest in group 1, significantly higher in groups 3 and 4 than group 5, and significantly higher in group 3 than in group 4 (all P<0.0001). Microscopic findings of kidney injury score, inflammatory cells (CD14+, F4/80+), and expressions of glomerular-damage indicators (FSP-1+/WT-1+) and renal tubular-damage indicators (KiM-1+/snail+) showed an identical pattern, whereas expressions of indices of glomerular-integrity (ZO-1+/p-cadherin+/podocin+/synaptopodin+) and angiogenesis (CD31+/vWF+/number of small vessels) biomarkers demonstrated an opposite pattern, to that of creatinine level (all P<0.001). Protein expressions of inflammatory (MMP-9/IL-1β/TNF-α/TLR-2/TLR-4), apoptotic (cleaved caspase-3/PARP/mitochondrial Bax), and oxidative-stress (NOX-1/NOX-2/oxidized protein) biomarkers exhibited an identical pattern, whereas anti-inflammatory (IL-10/IL-4) biomarkers displayed an opposite pattern, to that of creatinine level (all P<0.001). In conclusion, combined Ex4 and ADMSC therapy significantly protected kidney from acute IR-SS injury.

Keywords: Ischemia-reperfusion injury, sepsis syndrome, exendin-4, adipose derived mesenchymal stem cells, oxidative stress

Introduction

Acute kidney injury (AKI) is a common, heterogeneous clinical syndrome encompassing a spectrum of risk factors, etiologies and precipitatory acute insults, which occurs in multiple settings, including acute ischemia-reperfusion (IR) injury [1-4]. AKI can be classified according
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to the RIFLE (Risk, Injury, Failure, Loss and End-stage) and/or the Acute Kidney Injury Network (AKIN) criteria [5, 6]. Acute IR injury is one of the most common etiologies referred to within the Injury domain of the RIFLE classification and AKI is commonly caused by severe septic shock [6-9]. Previous studies have revealed that acute kidney IR injury complicated by severe sepsis is the leading cause of hospitalization for AKI and is associated with high morbidity and mortality [6-9], especially in patients who are immunocompromised, elderly or have diabetes mellitus (DM). There is currently no accepted effective treatment for acute kidney IR injury/AKI complicated by severe sepsis with septic shock.

Consistent with other causes of sepsis syndrome, acute kidney IR injury/AKI complicated by severe sepsis triggers a variety of inflammatory changes, culminating an overwhelming host immune response. This enhances the generation of oxidative stress and reactive oxygen species (ROS), increases the metabolic and humoral pathways and leads to remote organ dysfunction with detrimental effects on both short and long-term outcomes [9-13].

Exendin-4 (Ex4), a glucagon-like peptide-1 (GLP-1) analogue, was originally used to treat type 2 DM patients. Copious data have displayed that Ex4 therapy has protective effects against ischemia-induced tissue and organ damage mainly via its anti-oxidative and anti-inflammatory properties [14-17]; these are in addition to its hypoglycemic function [14-19]. Our recent studies have also shown that Ex4 possesses anti-apoptotic, anti-oxidative and anti-fibrotic capacity in the setting of ischemia-reperfusion organ injury [15] and ischemia-related organ dysfunction [14, 16, 17]. Plentiful investigations have shown that cell-based therapy provides a new therapeutic modality for ischemia-related organ dysfunction refractory to conventional therapy. Mesenchymal stem cells (MSC), particularly adipose-derived MSCs (ADMSCs), have immunomodulatory capacity and can suppress inflammation and oxidative stress [20-24]. We recently reported that ADMSC therapy effectively reduced acute urogenital organ damage and mortality rate in rat sepsis syndrome [13]. Accordingly, this study, tested the hypothesis that Ex4-assisted ADMSC therapy might be superior to either one alone for preserving kidney function in the setting of acute kidney IR injury complicated by sepsis syndrome (SS).

**Materials and methods**

**Ethics**

Both animal experimental protocols and procedures were approved by our Institute of Animal Care and Use Committee at Kaohsiung Chang Gung Memorial Hospital (Affidavit of Approval of Animal Use Protocol No. 2015032502) and executed in accordance with the Guide for the Care and Use of Laboratory Animals [The Eighth Edition of the Guide for the Care and Use of Laboratory Animals (NRC 2011)].

Animals were housed in an Association for Assessment and Accreditation of Laboratory Animal Care International (AAALAC)-approved animal facility in our hospital with controlled temperature and light cycles (24°C and 12/12 light cycles).

**Procedure and protocol of acute kidney ischemia-reperfusion (IR)**

Pathogen-free, adult male Sprague-Dawley (SD) rats (n=40) weighing 320-350 g (Charles River Technology, BioLASCO Taiwan Co. Ltd., Taiwan) were utilized in the present study. The protocol for the acute kidney IR procedure has been detailed in our previous reports [15, 17, 23]. Briefly, animals were anesthetized by inhalational 2.0% isoflurane and placed supine on a warming pad at 37°C for midline laparotomies. The sham control (SC) animals underwent laparotomy only. Acute IR injury of both kidneys was induced in all animals in acute kidney IR groups by clamping the renal pedicles for 40 minutes using non-traumatic vascular clips.

**Procedure for SS induction by cecal ligation and puncture (CLP)**

The procedure and protocol for SS induction, which was performed immediately after acute kidney IR procedure, was based on our previous report [24]. In the experimental CLP groups, the cecum of each animal was prolene suture-ligated at its distal portion (i.e. distal ligation), the cecum distal to the ligature was punctured twice with an 18G needle, and the distal cecal contents were manually expressed intraperitoneally. The abdominal muscle and skin were sutured and the animal was allowed to recover from anesthesia.
The animals in each group were euthanized and kidney specimens were collected for individual study by day 3 after the IR procedure.

**Animal grouping and rationale for the therapeutic regimen**

The animals were equally categorized into five groups (i.e., n=8 each group): Sham control (SC; laparotomy plus intra-peritoneal administration of 3.0 mL normal saline at 30 minutes and days 1 to 3 after IR procedure), IR-SS (receiving the same treatment as SC except for IR of both kidneys and SS induction), IR-SS + exendin-4 [(IR-SS + Ex4), subcutaneous administration of exendin-4 10 µg/kg at 30 minutes and at days 1 to 3 after IR procedure], IR-SS + ADMSC (1.2 × 10^6 cells intravenous administration at 3 h after IR procedure), and IR-SS + Ex4 + ADMSC.

The dosages and timings for exendin-4 administration were based on our recent reports [15-17]. The dosages and timings for ADMSC administration were based on our recent reports [13, 20, 21, 23-25].

**Flow cytometric analysis of circulatory inflammatory biomarkers day 3 after acute kidney IR-SS procedure**

**Whole blood-derived Ly6G (lymphocyte antigen 6 complex locus G6D):** Population of Ly6G positive cells from whole blood was determined by CytomicsTM FC500 flow cytometer (BD Biosciences, California, USA). Briefly, whole blood was diluted by PBS and further incubated with anti-rat Ly6G-FITC antibody for 30 minutes at room temperature. Incubated cells were washed with RBC lysis buffer (155 mM NH₄Cl, 12 mM NaHCO₃, 0.1 mM EDTA, in 1 L di-water, pH7.3) for 5 minutes and re-suspend in PBS for analysis. The results were recorded and evaluated by the CXP Analysis software (BD Biosciences, California, USA).

**Peripheral blood mononuclear cell (PBMCs)-derived CD12/TLR2 CD12/TLR4:** Population of toll-like receptor (TLR)-2, TLR-4, CD12/TLR2 and CD12/TLR4 positive cells were determined by (BD Biosciences, California, USA). Briefly, whole blood was diluted by PBS and further incubated with anti-rat Ly6G-FITC antibody for 30 minutes at room temperature. Incubated cells were washed with RBC lysis buffer (155 mM NH₄Cl, 12 mM NaHCO₃, 0.1 mM EDTA, in 1 L di-water, pH7.3) for 5 minutes and re-suspend in PBS for analysis. The results were recorded and evaluated by the CXP Analysis software (BD Biosciences, California, USA).

**Histological analysis of kidney injury scores at day 3 after acute kidney IR-SS procedure**

Histopathology scoring was assessed in a blinded fashion as we have previously described [15, 23, 25]. Briefly, the kidney specimens from all animals were fixed in 10% buffered formalin, embedded in paraffin, sectioned at 5 μm and stained with hematoxylin and eosin (H&E) for light microscopy. The scoring system reflects the grading of tubular necrosis, loss of brush border, cast formation, and tubular dilatation in 10 randomly chosen, non-overlapping fields (200 x) as follows: 0 (none), 1 (≤10%), 2 (11-25%), 3 (26-45%), 4 (46-75%), and 5 (≥76%) [23].

**Immunohistochemical (IHC) and immunofluorescent (IF) staining**

The procedure and protocol for IHC and IF staining have previously been reported in detail [13-17, 20, 21, 23-25]. For IHC and IF staining, rehydrated paraffin sections were first treated with 3% H₂O₂ for 30 minutes and incubated with Immuno-Block reagent (BioSB, Santa Barbara, CA, USA) for 30 minutes at room temperature. Sections were then incubated with primary antibodies specifically against zonula occludens-1 (ZO-1) (1:300, Abcam, Cambridge, MA, ClaySciences, USA), anti-rat TLR2-PE (abcam), and anti-rat TLR4-PE (abcam). Incubated PBMCs were then washed by PBS for 5 minutes and were subjected to analyze. The results were recorded and evaluated by the CXP Analysis software (BD Biosciences, California, USA).
USA), Wilm’s tumor suppressor gene 1 (WT-1) (1:1000, Abcam, Cambridge, MA, USA), kidney injury molecule (KIM)-1 (1:200, R&D system, Minneapolis, MN, USA), fibroblast specific protein (FSP)-1 (1:200, Abcam, Cambridge, MA, USA), P-cadherin (1:100, Novus, Littleton, CO, USA), snail (1:300, Abcam, Cambridge, MA, USA), podocin (1:100, Sigma, St. Louis, Mo, USA), dystrophin (1:100, Abcam, Cambridge, MA, USA), fibronectin (1:200, Abcam, Cambridge, MA, USA), and synaptopodin (1:100, Santa Cruz, Santa Cruz, CA, USA) while sections incubated with irrelevant antibodies served as controls. Three kidney sections from each rat were analyzed. For quantification, three randomly selected HPFs (200 x for IHC; 400 x for IF) were analyzed in each section. The mean number of positively-stained cells per HPF for each animal was then determined by summation of all numbers divided by 9.

An IHC/IF-based scoring system was adopted for semi-quantitative analyses of KIM-1, ZO-1, podocin, FSP-1, synaptopodin, snail and WT-1 as percentage of positive cells in a blind fashion [Score of positively-stained cell for podocin and WT-1: 0= no stain %; 1≤15%; 2=15~25%; 3=25~50%; 4=50~75%; 5≥75%-100%/per high-power filed (400 x)].

Western blot analysis

The procedure and protocol for Western blot analysis have been described in our previous reports [13-17, 20, 21, 23-25]. Briefly, equal amounts (50 μg) of protein extract were loaded and separated by SDS-PAGE using acrylamide gradients. After electrophoresis, the separated proteins were transferred electrophoretically to a polyvinylidene difluoride (PVDF) membrane (Amersham Biosciences, Amersham, UK). Non-specific sites were blocked by incubation of the membrane in blocking buffer [5% nonfat dry milk in T-TBS (TBS containing 0.05% Tween 20)] overnight. The membranes were incubated with the indicated primary antibodies [matrix metalloproteinase (MMP)-9 (1:3000, Abcam, Cambridge, MA, USA), tumor necrosis factor (TNF)-α (1:1000, Cell Signaling, Danvers, MA, USA), nuclear factor (NF)-κB (1:600, Abcam, Cambridge, MA, USA), NADPH oxidase (NOX)-1 (1:1500, Sigma, St. Louis, Mo, USA), NOX-2 (1:750, Sigma, St. Louis, Mo, USA), and actin (1:10000, Chemicon, Billerica, MA, USA)] for 1 hour at room temperature. Horseradish peroxidase-conjugated anti-rabbit immunoglobulin IgG (1:2000, Cell Signaling, Danvers, MA, USA) was used as a secondary antibody for one-hour incubation at room temperature. The washing procedure was repeated eight times within one hour. Immunoreactive bands were visualized by enhanced chemiluminescence (ECL; Amersham Biosciences, Amersham, UK) and exposed to Biomax L film (Kodak, Rochester, NY, USA). For quantification, ECL signals were digitized using Labwork software (UVP, Waltham, MA, USA).

Assessment of oxidative stress

The procedure and protocol for assessing the protein expression of oxidative stress have been detailed in our previous reports [13-17, 20, 21, 23-25]. The Oxyblot Oxidized Protein Detection Kit was purchased from Chemicon, Billerica, MA, USA (S7150). DNPH derivatization was carried out on 6 μg of protein for 15 minutes according to the manufacturer’s instructions. One-dimensional electrophoresis was carried out on 12% SDS/polyacrylamide gel after DNPH derivatization. Proteins were transferred to nitrocellulose membranes which were then incubated in the primary antibody solution (anti-DNP 1:150) for 2 hours, followed by incubation in secondary antibody solution (1:300) for 1 hour at room temperature. The washing procedure was repeated eight times within 40 minutes. Immunoreactive bands were visualized by enhanced chemiluminescence (ECL; Amersham Biosciences, Amersham, UK) which was then exposed to Biomax L film (Kodak, Rochester, NY, USA). For quantification, ECL signals were digitized using Labwork software (UVP, Waltham, MA, USA). For Oxyblot protein analysis, a standard control was loaded on each gel.

Statistical analysis

Quantitative data are expressed as means ± SD. Statistical analysis was adequately performed by ANOVA followed by Bonferroni multiple-comparison post hoc test. Statistical analysis was performed using SPSS statistical software for Windows version 13 (SPSS for Windows, version 13; SPSS, IL, U.S.A.). A P value of less than 0.05 was considered statistically significant.

Results

Creatinine and BUN levels, urine amount and the ratio of urine protein to creatinine at baseline and day 3 after IR-SS procedure (Figure 1)

Prior to the IR-SS procedure, the circulating levels of creatinine and BUN, urine volume, and
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Figure 1. Circulating levels of creatinine, blood urine nitrogen (BUN), urine amount, and the ratio of urine protein to creatinine at baseline and day 3 after IR-SS procedure. By day 0 prior to IR-SS procedure: A. Creatinine level, SC vs. other groups, P>0.5. B. BUN level, SC vs. other groups, P>0.5. C. Urine amount, SC vs. other groups, P>0.5. D. Ratio of urine protein to creatinine, SC vs. other groups, P>0.5. By day 3 after IR-SS procedure: E. Creatinine level, * vs. other groups with different symbols (†, ‡, §), P<0.0001. F. BUN level, * vs. other groups with different symbols (†, ‡, §), P<0.0001. G. Urine amount, * vs. other groups with different symbols (†, ‡, §), P<0.0001. H. Ratio of urine protein to creatinine, * vs. other groups with different symbols (†, ‡, §), P<0.0001. All statistical analyses were performed by one-way ANOVA, followed by Bonferroni multiple comparison post hoc test (at least n=7 for each group). Symbols (*, †, ‡, §) indicate significance at the 0.05 level. SC = sham-operative control; IR-SS = ischemia-reperfusion and sepsis syndrome; Ex4 = exendin-4; ADMSC = adipose-derived mesenchymal stem cell.

Figure 2. Flow cytometric assessment of circulating inflammatory biomarkers and mortality rate by day 3 after IR-SS procedure. A. The survival rate of five groups (n=16) by rate Pairwise comparisons (without Bonferroni's correction), * vs. †, P<0.0005. B. Kaplan Meier survival cure. By log Rank test, P=0.004. C. Circulating number of Ly6G+ cells, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. D. Circulating number of tall-like receptor (TLR)-2+ cells, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. E. Circulating number of TLR-4+ cells, * vs. other groups with different symbols (†, ‡), P<0.0001. F. Circulating level of CD14/TLR-2+ cells (i.e., double stain), * vs. other groups with different symbols (†, ¶), P<0.001. G. Circulating number of CD14/TLR-4+ cells (double stain) did not differ among the five groups, P>0.2. All statistical analyses were performed by one-way ANOVA, followed by Bonferroni multiple comparison post hoc test (at least n=7 for each group). Symbols (*, †, ‡, §) indicate significance at the 0.05 level. SC = sham-operative control; IR-SS = ischemia-reperfusion and sepsis syndrome; Ex4 = exendin-4; ADMSC = adipose-derived mesenchymal stem cell.
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the ratio of urine protein to urine creatinine did not differ among the five groups (Figure 1A-D). However, by 72 h after IR-SS procedure, the creatinine and BUN levels, two indicators of renal dysfunction (Figure 1E, 1F), and the ratio of urine protein to urine creatinine, an indicator...
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Figure 4. Protein expressions of inflammatory and anti-inflammatory biomarkers in kidney parenchyma by day 3 after IR-SS procedure. A. Protein expression of matrix metalloproteinase (MMP)-9, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. B. Protein expression of interleukin (IL)-1β, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. C. Protein expression of tumor necrosis factor (TNF)-α, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. D. Protein expression of toll-like receptor (TLR)-2, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. E. Protein expression of TLR-4, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. F. Protein expression of IL-4, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. G. Protein expression of IL-10, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. All statistical analyses were performed by one-way ANOVA, followed by Bonferroni multiple comparison post hoc test (at least n=7 for each group). Symbols (*, †, ‡, §, ¶) indicate significance (at 0.05 level). SC = sham-operative control; IR-SS = ischemia-reperfusion and sepsis syndrome; Ex4 = exendin-4; ADMSC = adipose-derived mesenchymal stem cell.

Flow cytometric analysis of circulating level of inflammatory cells and mortality rate by day 3 after IR-SS procedure (Figure 2)

The mortality rate from SC to IR-SS + Ex4-ADMSC was 0% (0/16), 56.25% (9/16), 31.25% (5/16), 37.5% (6/16) and 18.75% (3/16), respectively (Figure 2A). By Pairwise comparisons (without Bonferroni’s correction), the mortality rate between two different groups was (1) SC vs. IR-SS, P=0.0004, (2) SC vs. IR-SS + Ex4, P=0.016, (3) SC vs. IR-SS + ADMSC, P=0.007, (4) SC vs. IR-SS + Ex4 + ADMSC, P=0.074, (5) IR-SS vs. IR-SS + Ex4 + ADMSC, P=0.015, (6) IR-SS vs. IR-SS + Ex4, P=0.156, (7) IR-SS vs. IR-SS + ADMSC, P=0.212, and (8) IR-SS + Ex4 vs. IR-SS + ADMSC, P=0.816, respectively (Figure 2B).

The circulating level of level Ly6G+ cells, an indicator of innate inflammatory reaction, was highest in in IR-SS and lowest in SC, significantly higher in IR-SS + Ex4 and IR-SS + ADMSC, but they showed no difference between the latter two groups. Urine amount, an index of glomerular filtration integrity, was highest in IR-SS + Ex4 + ADMSC and lowest in IR-SS, but was not different among the groups of SC, IR-SS + Ex4 and IR-SS + ADMSC (Figure 1G).

The circulating level of TLR-2 was significantly higher in IR-SS and lowest in SC and IR-SS + Ex4-ADMSC, and significantly in IR-SS + Ex4 than in IR-SS + ADMSC, but it showed no difference between SC and IR-SS + Ex4-ADMSC (Figure 2D).
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Figure 5. Protein expressions of apoptotic, anti-apoptotic and oxidative-stress biomarkers in kidney parenchyma by day 3 after IR-SS procedure. A. Protein expression of cleaved caspase 3 (c-Casp 3), * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. B. Protein expression of cleaved Poly (ADP-ribose) polymerase (c-PARP), * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. C. Protein expression of mitochondrial (mito)-Bax, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. D. Protein expression of Bcl-2, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. E. Protein expression of NOX-1, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. F. Protein expression of NOX-2, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. G. Oxidized protein expression, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. (Note: left and right lanes shown on the upper panel represent protein molecular weight marker and control oxidized molecular protein standard, respectively). M.W = molecular weight; DNP = 1-3 dinitrophenylhydrazone. All statistical analyses were performed by one-way ANOVA, followed by Bonferroni multiple comparison post hoc test (at least n=7 for each group). Symbols (*, †, ‡, §, ¶) indicate significance (at 0.05 level). SC = sham-operative control; IR-SS = ischemia-reperfusion and sepsis syndrome; Ex4 = exendin-4; ADMSC = adipose-derived mesenchymal stem cell.

groups and significantly higher in IR-SS + Ex4 than in SC, IR-SS + ADMSC and IR-SS + Ex4 + ADMSC, but it showed no difference among the later three groups (Figure 2E, 2F). However, the circulating level of CD14/TLR-4 did not differ among the five groups (Figure 2G).

Light microscopic findings for assessment of kidney injury score and small vessel density in kidney parenchyma by day 3 after IR-SS procedure (Figure 3)

Light microscopy of H&E-stained kidney sections revealed that kidney injury score was highest in IR-SS and lowest in SC, significantly higher in IR-SS + Ex4 and IR-SS + ADMSC than that in IR-SS + Ex4 + ADMSC, and significantly higher in IR-SS + Ex4 than that in IR-SS + ADMSC (Figure 3A-F). On the other hand, α-smooth muscle actin staining for identifying vessel density in kidney parenchyma exhibited that the number of small vessels (i.e., the diameter ≤25 μM) exhibited an opposite pattern to kidney injury score among the five groups (Figure 3G-L).

The protein expression of inflammatory and anti-inflammatory biomarkers in kidney parenchyma by day 3 after IR-SS procedure (Figure 4)

The protein expressions of MMP-9, IL-1ß, TNF-α, TLR-2 and TLR-4, five indicators of inflammation, were highest in IR-SS, lowest in SC, and significantly higher in IR-SS + Ex4 than in IR-SS + ADMSC and IR-SS + Ex4 + ADMSC (Figure 4A-E). However, protein expressions of IL-4 and IL-10, two indices of anti-inflammation, showed progressive increases from SC to IR-SS + Ex4 + ADMSC, suggesting a phenomenon of intrinsic response to inflammatory stimulation (Figure 4F, 4G).
The protein expressions of apoptotic, anti-apoptotic and oxidative-stress biomarkers in kidney parenchyma by day 3 after IR-SS procedure (Figure 5)

The protein expression of cleaved caspase 3, cleaved PARP and mitochondrial Bax, three indicators of apoptosis, were highest in IR-SS, lowest in SC, significantly higher in IR-SS + Ex4 and IR-SS + ADMSC than in IR-SS + Ex4 + ADMSC, and significantly higher in IR-SS + Ex4 than in IR-SS + ADMSC (Figure 5A-C). Conversely, the protein expression of Bcl-2, an indicator of anti-apoptosis, showed an opposite pat-
tern to the apoptotic biomarkers among the five groups (Figure 5D).

The protein expressions of NOX-1, NOX-2 and oxidized protein, three indicators of oxidative stress, exhibited an identical pattern to apoptosis among the five groups (Figure 5E-G).

Inflammatory cell infiltration and angiogenesis expressions in kidney parenchyma by day 3 after IR-SS procedure (Figures 6, 7)

IF microscopic finding showed that the KIM-1, a kidney injury biomarker predominant expression in renal tubules, was highest in IR-SS, low-
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Figure 8. Assessment of ZO-1+ and podocin+ cells in kidney parenchyma by day 3 after IR-SS procedure. (A-E) Immunofluorescent microscopy (400x) for ZO-1+ cells (green color) in kidney parenchyma. (F) Analytical result of number of ZO-1+ cells, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. The red color with Dil stained nucleus (yellow arrows) in (D) and (E) indicated the dye-labeling ADMSCs. (G-K) Immunohistochemical microscopy (400x) for podocin+ cells (gray color) in kidney parenchyma. (L) Analytical result of number of podocin+ cells, * vs. other groups with different symbols (†, ‡, §, ¶), P<0.0001. Scale bars in right lower corner represent 20 µm. All statistical analyses were performed by one-way ANOVA, followed by Bonferroni multiple comparison post hoc test (at least n=7 for each group). Symbols (*, †, ‡, §, ¶) indicate significance (at 0.05 level). SC = sham-operative control; IR-SS = ischemia-reperfusion and sepsis syndrome; Ex4 = exendin-4; ADMSC = adipose-derived mesenchymal stem cell.

IF microscopy also showed that the cellular expression of vWF (Figure 7A-F), an indicator of endothelial function, exhibited an opposite pat-
tern to inflammatory cells among the five groups. On the other hand, the IF microscopy identified that the cellular expression of CXCR4, an indicator of endothelial progenitor cell/angiogenesis, significantly progressively increased from SC to IR-SS + Ex4-ADMSC (Figure 7G-L).
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Microscopic findings of podocyte components and renal tubules at day 3 after IR-SS procedure (Figures 8-10)

IF analysis displayed that the expression of ZO-1, a tight junction-associated protein that offers a link between the integral membrane proteins and the filamentous cytoskeleton in podocytes, was highest in SC, lowest in IR-SS, significantly higher in IR-SS + Ex4-ADMSC than in IR-SS + Ex4 and IR-SS + ADMSC, and significantly higher in IR-SS + ADMSC than in IR-SS +...
Ex4 (Figure 8A-F). Consistently, IHC staining showed that the expression of podocin, a component of podocyte foot process revealed an identical pattern to ZO-1 among the five groups (Figure 8G-L).

Additionally, IHC staining showed that changes in the expression of fibroblast specific protein 1 (FSP-1), which is mainly situated in kidney interstitials, showed an opposite pattern compared with ZO-1 among the five groups (Figure 9A-F). On the other hand, IF demonstrated that the synaptopodin, a podocyte foot process component, displayed an identical pattern to ZO-1 among the five groups (Figure 9G-L). Furthermore, IHC staining revealed that snail, predominantly accumulated in the tubular nuclei (Figure 10A-F), and WT-1, predominantly in podocytes (Figure 10G-L), showed an opposite pattern to ZO-1 among the five groups.

Discussion

This study investigated the therapeutic effect of Ex4-ADMS in a preclinical setting of IR-SS and yielded several striking implications. First, the present study successfully created an experimental model that mimicked the clinical setting of AKI complicated by SS, which is an extremely common scenario in daily clinical practice. Additionally, based on this experimental model, we delineated that IR-SS damaged the kidney through highly complex pathomechanisms. Second, the histopathological findings (i.e., destructive glomerular and renal tubular architectures were identified by kidney injury score) explained why remarkably high urine protein to creatinine ratio and circulating levels of BUN and creatinine were found in animals with acute renal IR-SS compared to SC animals. Third, the results of the present study highlighted that Ex4-ADMSC therapy contributed a marked additive therapeutic effect for protecting kidney against IR-SS injury, inviting prospective clinical trials to evaluate the potential impact of this combined regimen in patients with AKI complicated by SS, especially DM patients.

An essential finding in the present study is that the circulating levels of BUN and creatinine, two important indices of AKI/renal dysfunction, were notably increased in animals with IR-SS at day 3 compared with sham-operated controls. Additionally, the ratio of urine protein to creatinine, another renal functional indicator, was comparable to that of creatinine level, whereas, the urine amount revealed an opposite pattern to creatinine level among the five groups; this indicates the fragility of kidney to damage in various disease entities, including IR-SS.

It is well recognized that AKI/acute kidney IR injury and IR-SS occur commonly in the clinical setting. Most previous studies [23, 25-28] utilized the experimental model of AKI/acute kidney IR injury rather than the experimental model of IR-SS to investigate the outcomes or effects of different management strategies. To the best of our knowledge, this is the first experimental model that mimicked the clinical setting of AKI/acute kidney IR injury complicated by SS. Importantly, the results of the present study demonstrated that combined treatment with Ex4 and ADMSC was superior to either alone for protecting kidney against IR-SS.

The inflammatory reaction and overwhelming immune response associated with generation of ROS/oxidative stress have been identified as the two processes with crucial roles for organ damage leading to inevitable functional deterioration and poor acute outcomes [9-13, 29]. Another essential finding in the present study was that the cellular and protein levels of inflammatory reactions were markedly upregulated in IR-SS animals compared to SC. In addition, the generation of ROS/oxidative stress was markedly augmented in an identical way between the two groups. These findings could partially explain why the renal function (i.e., increased circulating levels of creatinine and BUN) was remarkably impaired and the kidney architecture affected detrimentally (i.e., an increase in kidney injury score) in IR-SS animals. Importantly, these parameters in IR-SS animals were significantly reversed by Ex4 and ADMSC treatment and further significantly reversed after receiving Ex4-ADMSC treatment.

Previous [30-32] and our recent [15] studies have shown that the presence of intact podocytes and functional integrity between podocytes and their components (i.e., primary processes, secondary processes and foot processes) play major roles in establishing a size-selective barrier to protein loss. An important finding in the present study was that, when we looked at the cellular level of microscopic findings, the integrity [i.e., Figures 8-10] of ultrastructural components of podocytes was sub-

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Substantially devastated in IR-SS animals as compared with SC. Additionally, apoptotic biomarkers in kidney parenchyma were also substantially higher in IR-SS animals than in the SC. These molecular-cellular perturbations could explain why the ratio of urine protein to creatinine was remarkably increased in IR-SS animals compared to SC. As expected, these molecular-cellular perturbations in IR-SS animals were notably suppressed by Ex4 or ADMSC treatment and further suppressed by Ex4-ADMSC treatment.

Abundant data have shown that angiogenesis plays a crucial role in the restoration of blood flow to ischemic regions, to improve IR-caused/ischemia-related organ dysfunction [20, 21, 23-25, 33]. Stem cell therapy enhances angiogenesis, as has been clearly identified by copious previous studies [20, 21, 23-25, 33]. Furthermore, Ex4 has anti-oxidative stress effects as well as angiogenesis capacity through upregulating the circulating levels of endothelial progenitor cells, angiogenesis factors and vascular density in ischemic regions [14, 17]. A principal finding in the present study, at cellular and tissue levels, was that angiogenesis biomarkers as well as vessel density were significantly higher in IR-SS + Ex4 and in IR-SS + ADMSC animals, and even more significantly higher in IR-SS + Ex4 + ADMSC animals as compared to IR-SS animals. These findings reinforce previous studies [14, 17, 20, 21, 23-25, 33] and help explain why renal function and renal parenchyma were notably preserved in the former three groups of animals than in the latter group of animals.

Study limitation

This study has limitations. First, the purpose of the study was designed to investigate the acute phase of IR-SS that mimicked daily clinical practice. Long-term outcomes from IR-SS were therefore out with the scope of this present study. Second, despite the extensive investigations in the present study, the precise mechanism of Ex4-ADMSC therapy that improves outcomes in rodents after IR-SS is still not entirely clear.

Conclusion

In conclusion, the results of the present study demonstrated that Ex4-ADMSC therapy effectively protected the integrity of kidney parenchyma and renal function in the setting of IR-SS in rodents.

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Disclosure of conflict of interest

None.

Authors' contribution

Pei-Hsun Sung and Hsin-Ju Chiang: design, data acquisition, analysis, and drafting the manuscript. Christopher Glenn Wallace, Chih-Chao Yang, Yen-Ta Chen, Kuan-Hung Chen, and Chih-Hung Chen: laboratory assay and troubleshooting. Pei-Lin Shao, Yung-Lung Chen, Sarah Chua, Han-Tan Chai, Yi-Ling Chen, and Tien-Hung Huang: data acquisition, analysis, and interpretation. Hon-Kan Yip and Mel S. Lee: conceived of the study, participated design, coordination, and helped to draft the manuscript.

Abbreviations

Ex4, exendin-4; ADMSC, adipose-derived mesenchymal stem cell; IR, ischemia-reperfusion; SS, sepsis syndrome; AKI, acute kidney injury; RIFLE, Risk, Injury, Failure, Loss and End-stage; DM, diabetes mellitus; ROS, reactive oxygen species; GLP-1, glucagon-like peptide-1; MSC, mesenchymal stem cells; CLP, cecal ligation and puncture; Ly6G, Lymphocyte antigen 6 complex locus G6D; PBMC, peripheral blood mononuclear cell; TLR, toll-like receptor; BUN, blood urine nitrogen; ZO-1, zonula occludens-1; WT-1, Wilm’s tumor suppressor gene 1; KIM-1, kidney injury molecule-1; vWF, Von Willebrand factor; CXCR4, C-X-C chemokine receptor type 4; FSP-1, fibroblast specific protein-1; MMP-9, matrix metalloproteinase-9; TNF-α, tumor necrosis factor-α; NF-κB, nuclear factor-κB; PARP, Poly-ADP-ribose polymerase; NOX, NADPH oxidase.

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