Role of mesenchymal stem cells and their culture medium in alleviating kidney injury in rats diabetic nephropathy

Amal Al-Shahat Ibrahim 1, Manal Mohammad Morsy 1, Safwat E. Abouhashem 2, Omnia Aly 3, Norhan A. Sabbah 4 and Nermin Raafat 4*

Abstract

Background: Diabetic nephropathy (DN) is considered as one of the most serious complications resulting from diabetes mellitus and end-stage of renal failure globally. Up to 40% of diabetic patients will develop DN. The involvement of mesenchymal stem cells (MSCs) in diabetic renal lesions management has been established in many animal models of DN. The aim is to evaluate the capability of MSCs and their culture medium (CM) to alleviate DN in streptozotocin (STZ)-induced diabetic rat model. Female albino rats were made diabetic and were further categorized into 4 subgroups of 15 each: DN group, DN group received fibroblasts, MSCs group received one dose of $1 \times 10^6$ cells of MSCs, and CM group received one dose of 500 μl of CM. In all groups, the treatment was delivered by intravenous injection (IV) into the tail vein.

Results: MSCs insinuated themselves into the injured kidney as detected by CD44 expression. Biochemical and histological results showed that MSCs and/or CM effectively attenuated DN manifestations in rat model through their possible anti-inflammatory (tumor necrosis factor-α and transforming growth factor-β1 were decreased), anti-apoptotic (Bcl2 was increased while Bax and caspases were decreased), and anti-oxidant role (malondialdehyde was decreased while glutathione and catalase were increased).

Conclusion: These results provide a potential therapeutic tool for DN management through the administration of the CM from MSCs that ameliorates the effects of diabetes. It is also possible to treat DN using CM alone thus avoiding cell transplantation.

Keywords: MSCs, Culture medium, Diabetic nephropathy, Therapeutic effect

Background

Diabetes mellitus (DM) is a worldwide public health challenge. WHO estimated that there were around 422 million people living with diabetes and that there was a rising trend in the number of people living with DM. Diabetic nephropathy (DN) is the major micro-vascular complication in diabetic patients and accounts for approximately half of all end-stage renal disease globally [1].

The key characterized features of diabetic nephropathy include thickened basement membrane of both glomeruli and tubules, mesangial cell (MC) hypertrophy, extracellular matrix accumulation, tubule-interstitial fibrosis, and finally glomerular sclerosis which is the leading cause of micro-albuminuria and renal failure [2, 3]. Over the last decade, many factors have been proposed to be involved in the pathogenesis of diabetic nephropathies such as genetic and hemodynamic factors, metabolic, oxidative stress, and cytokine signaling [2, 4].

Hyperglycemia is a discriminative pathologic element in diabetes mellitus which not only generates free radicals but also attenuates antioxidant effects via glycation...
of antioxidant enzymes [5]. The major sources of reactive oxygen species (ROS) in DN are advanced glycation end products (AGEs), autoxidation of glucose, xanthine oxidase activity, mitochondrial respiratory chain deficiencies, NADPH oxidase, and nitric oxide synthase (NOS) [6, 7]. Streptozotocin is extensively used to induce diabetes model, and its toxicity is mediated through nitric oxide (NO) generation that affects pancreatic β-cells [8].

Currently available treatments, involving glycemic and/or blood pressure control, can delay but not abolish the development of DN [9]. Therefore, there is an imperative need to develop new, effective therapeutic strategies that preserve normal renal function or halt the progression of diabetic nephropathy.

Mesenchymal stem cells are multipotent progenitor cells which are non-hematopoietic and can differentiate into many types of cells. Stem cell therapy is considered as a new promising approach for treatment of diabetes and its complications [10]. Adipose-derived MSCs (AD-MSCs) have the capability of differentiating into cells and tissues which are of mesodermal origin. They are easily harvested and abundant which makes them an ideal choice for use [11]. MSCs have a lot of advantages for use for therapeutic purposes, such as the capability to migrate to the sites where tissue injury is present, potent immunosuppressive action [12, 13], and safer after infusion of allogeneic MSCs [14, 15].

It is reported that after the injection of MSCs, they directly infiltrate the diseased kidneys [16]. Other studies have found no evidence for direct MSCs incorporation into the kidney during the repair processes [17]. Many factors play a great role in the therapeutic action of MSCs like their ability to modulate and change local environment, activation of endogenous progenitor cells, secretion of several growth factors, and their differentiation potency to repair diseased tissue [18, 19].

Vascular endothelial growth factor-a (VEGF-A) is a protein secreted by podocytes that is necessary for the survival of endothelial cells, podocytes, and mesangial cells. Human kidney biopsies showed high VEGF-A mRNA at early stages of DN, and lower VEGF-A expression in advanced DN, specifically in sclerotic glomeruli and mesangial nodules [20, 21]. The decline in VEGF-A expression as overt DN progresses is thought to be due to podocyte dropout [22, 23].

Based on the previously mentioned findings, we, therefore, raised the hypothesis that cytotherapy by the use of MSCs transplantation is of benefit in reducing diabetic nephropathy. Accordingly, this study was undertaken to evaluate the capability of MSCs to alleviate DN in streptozotocin (STZ)-induced diabetic rat model, to investigate the mechanisms underlying the protective effect of MSCs in DN and to mark the signals of paracrine origin that mediate the action of MSCs.

**Methods**

**Animals and ethics**

Seventy-five healthy female albino rats (6–8 weeks) weighing 150–200 g and ten male albino rats (6 weeks) weighing 120–140 g were included in this study. They were obtained from the animal facility house of the faculty, and they were housed in a light and temperature controlled room with free access to water and food. All experimental procedures were done according to guidelines of the Institutional Review Board (IRB) for research studies (reference number is 4920/14-7-2016) and IACUC (approval number is ZU-IACUC/3/F/90/2018) for animal research.

**STZ-induction of diabetic nephropathy in rats**

To establish type 1 diabetic rat model, 50 mg/kg body weight of STZ (Sigma, St Louis, USA) was dissolved in 0.05 ml saline and injected intraperitoneal into adult rats twice 4 days apart. Diabetes was defined when a random rat blood glucose level reading of more than 300 mg/dl in the 3 continuous days at day 14 after STZ injection was done [24]. Diabetic nephropathy was confirmed 4 weeks after onset of diabetes mellitus by measuring serum urea and creatinine in blood and also by histopathological changes scarifying one rat every 2 weeks [25].

**Experimental design**

The animals were allocated into the following groups, control group which received phosphate buffered saline (PBS) at time of injection, included 15 female albino rats, and diabetic nephropathy group, comprised 60 female albino rats that were further categorized into 4 subgroups of 15 each: DN group, fibroblast group received one dose of 1 × 10⁶ HFF-1 cells (a human fibroblast cell line from American Type Culture Collection (ATCC)) were used as the negative control for migrating cells according to Domouky et al. [24], MSCs group received one dose of 1 × 10⁶ cells of MSCs per rat, and CM group received one dose of 500 μl of culture medium (MSCs were cultured with Dulbecco’s Modified Eagle’s Medium (DMEM) for 24 h. In all groups, the treatment was delivered by intravenous injection (IV) into the tail vein.

**Preparation of adipose tissue-derived mesenchymal stem cells (AT-MSCs)**

**Isolation of AD-MSCs from male rats**

AT-MSCs were isolated from the peritoneal adipose tissue excised from the abdominal region of 6–8-week-old male albino rats. Adipose tissue was
enzymatically digested with 0.075% collagenase (type I; Sigma–Aldrich, St. Louis, MO) in PBS for 1 h at 37 °C with gentle agitation. The collagenase was inactivated with an equal volume of DMEM (GIBCO/BRL) supplemented with 10% fetal bovine serum (FBS), and the infranatant was centrifuged at 1000×g for 5 min at room temperature. The resulting cell pellet was resuspended in 160 mM NH4Cl (Stem Cell Technologies Inc., Vancouver, BC), incubated at room temperature for 2 min to eliminate contaminating red blood cells, and filtered through a 100-μm nylon mesh strainer (Becton Dickinson Labware, Franklin Lakes, NJ) to remove debris [26]. The culture of isolated cells was done according to Raafat et al. [27].

For the conditioned medium-treated groups, the MSCs were cultured in DMEM without FBS for 24 h. After this period, the medium was collected, centrifuged, and filtered to remove the cells, treated with an RNase inhibitor (500 U) and administered to the rats [28].

Characterization of AD-derived MSCs
MSCs at day 14 are characterized by their adhesiveness and fusiform, star, or spindle shape. MSCs by flow cytometric evaluation express CD105, CD73, and CD29 on their cell surface, and they do not express CD34, CD45, or MHC-class II.

Blood and tissue sampling
After 12 weeks of STZ injection according to Fang et al. [25], animals were fasted for half a day, and blood samples were obtained by puncturing retro-orbital plexus under light ether anesthesia. All rats were then sacrificed by cervical dislocation, and tissue was perfused with PBS solution, containing 0.16 mg/ml heparin to remove any red blood corpuscles and clots. Both kidneys were immediately excised, dried, and weighed. The parts of the kidney were kept in 10% formalin for the histological assessment and immune-histochemistry, and then the kidneys were minced into small parts for different parameters assessment. The blood samples were left for 30–60 min for spontaneous clotting. Serum samples were separated for assessment of urea and creatinine.

PCR detection of the Sry gene
Genomic DNA was extracted from the kidney tissue using the QIAnp DNA purification kit (QIAGEN, Germany). PCR analysis for Sry gene expression in AD-MSCs recipient rats was done using primer sequence listed in Table 1 [29]. The amplification protocol was as follows: 95 °C for 5 min; then 35 cycles of 95 °C for 35 s, 50 °C for 60 s, and 72 °C for 60 s; then 72 °C for 5 min, using heated lid thermal cycler (Mastercycler® pro, Eppindor, Germany).

Reverse transcription PCR for renal localization of MSC
Total RNA was isolated from the renal tissue with Qia-genRNaseasy Mini kit. RNA was reverse-transcribed to cDNA using a Primescript RT Reagent Kit (Takara, Shiga, Japan). The primers sequences used for CD105 and CD34 are listed in Table 1. PCR was performed for 35 cycles with each cycle comprising 20 s at 94 °C, 30 s at 55 °C for CD105 and 60 °C for CD34 and β-actin, and 1 min at 72 °C [30].

Biochemical analysis
Kidney function assessment
Urea level was estimated in accordance to Fawcett and Scott’s method (1960) [31], and serum creatinine level was measured in accordance to Henry’s method (1974)

### Table 1
| Target   | Ref seq accession | Sequence | Product size (bp) |
|----------|------------------|----------|-------------------|
| Sry      | NM_012772        |          | 402               |
| CD105    | NM_007932        |          | 374               |
| CD34     | NM_133654        |          | 744               |
| Bax      | NM_017059.2      |          | 120               |
| Bcl-2    | NM_016993.1      |          | 144               |
| Caspase-3| NM_012922.2      |          | 180               |
| Caspase-8| NM_022277.1      |          | 163               |
| β-actin  | NM_031144.3      |          | 150               |

Bcl-2 B cell leukemia/lymphoma, Bax Bcl-2 associated X protein
[32] by using reagent kit obtained from the Diagnostics Company (Egypt).

Renal oxidant-antioxidant markers assessment
Malondialdehyde (MDA) [33], reduced glutathione (GSH) [34], and catalase (CAT) [35] on renal tissues were assessed in accordance to the respective manufacturer’s instructions (Biodiagnostic Tahreer St., Dokki, Giza, Egypt).

Measurement of tumor necrosis factor-α (TNF-α) and transforming growth factor-β1 (TGF-β1)
The content of protein in kidney homogenates (Vertishear tissue homogenizer) was determined by Bradford’s method (1976) [36] by using crystalline BSA as standard. The TNF-α and TGF-β1 levels were quantified by quantitative enzyme-linked immunosorbent assay (ELISA) kits according to the manufacturer’s instructions (TNF-α rat ELISA kit, ab100785 and TGF-β1 rat ELISA kits, ab119558, Abcam, UK). Results are mentioned as picogram per milligram protein.

Quantitative real-time PCR (qRT-PCR) for apoptotic markers gene expression
Expression levels in cDNA were determined by Stratagene, MX3000P quantitative PCR System (Agilent Technologies) and analyzed using the MxPro QPCR Software (Agilent technologies). Table 1 lists the primer sequences used for Bax, Bcl2, caspase-3, 8, and β-actin [37, 38].

The PCR was performed on 25 μl containing 12.5 μl 2× QuantiFast SYBR Green PCR Master Mix, 1 μM of each primer, and 5 μl cDNA with the following conditions: 95 °C for 5 min, then 40 cycles at 95 °C for 10 s, and combined annealing and extension 60 °C for 30 s. All kits were supplied by (QIAGEN, Valencia, CA, USA). Samples were assessed in duplicates to make sure the reproducibility and accuracy of the obtained results.

Histological and immunohistochemical analysis
The kidneys were dissected, washed with phosphate buffer saline, and fixed in 10% formalin solution. Tissue samples were dehydrated in gradual grades of alcohol, cleared in xylolthen embedded in paraffin sections. Five micrometer thickness sections were obtained via using a microtome, mounted on slides, and subjected to hematoxylin and eosin stain (H&E) [39], periodic acid Schiff reaction (PAS) counterstained with hematoxylin [40], Mallory’s trichrome stain [41], immunohistochemical staining for detection of CD44 antigen using rat monoclonal anti CD44 antibody “Neomarkers, Labvision, USA,” as a marker for mesenchymal stem cells [42], and immunohistochemical staining for VEGF [43].

The avidin-biotin-peroxidase complex technique was done as follows for immunohistochemical detection of CD44 and VEGF; paraffin sections were deparaffinized and rehydrated. The endogenous activity of peroxidase was blocked by 3% solution of hydrogen peroxide for 5 min. Then, the kidney sections were incubated for 30 min with primary antibody. The primary antibody used was a monoclonal anti-CD44 (Lab Vision Corporation, Thurmont, California, USA) and monoclonal antibody for detection of VEGF (anti-VEGF, Lab Vision, USA). The secondary antibody (biotinylated goat anti-rabbit) was applied after washing the sections with phosphate buffer. The slides were rinsed in PBS then incubated with labelled avidin-biotin peroxidase. Avidin-biotin-peroxidase complex (Histostain SP kit, Zymed Laboratories Inc, San Francisco, USA) is used for immunodetection of the bound primary antibody. The binding site was visualized after adding diaminobenzidine (DAB) which was used as a chromogen, which is converted into a brown precipitate and Meyer’s hematoxylin as a counterstain.

Quantitative morphometric study
Morphometric measurements were performed via using Leica Q500 Imaging analysis System. The following parameters were measured: urinary space diameter (USD) (μm) which is the distance between visceral and parietal layer of the glomerulus, outer diameter of the proximal convoluted tubules (PCD) (μm), outer diameter of the distal convoluted tubule (DCD) (μm), and the area percent (AP) (μm²) occupied by collagen fibers within the glomeruli and around the tubules. Measurements of USD, PCD, and DCD were assessed in PAS-stained slides while AP was calculated in Mallory stained sections. All parameter measurements were detected in magnification of × 400 for 25 randomly selected glomeruli, proximal, and distal convoluted tubules in each animal.

Statistical analysis
The results for quantitative variables are mentioned as mean ± standard deviation. The means were compared in a one-way analysis of variance and multiple comparison post hoc tests. Differences between groups were considered significant when P values were < 0.05 and highly significant when P values were < 0.001. Statistical analyses were done via the use of SPSS, version 10 (SPSS Inc).

Results
Characterization of AD-MSCs
AD-MSCs were characterized by their adhesiveness and fusiform shape, spindle, and fibroblast-like cells. Flow cytometric analysis of cell surface markers on gated cells showed positive surface expression of CD29, CD73, and CD105 (PE labelled) while the mainstream of cells showed negative expression of CD34, CD45, and MHC-
class II (FITC labelled). This surface marker expression patterns corresponded to AD-MSC in accordance with the International Society of Cellular Therapy system. The kidney sections of AD-MSC group showed a CD44 positive immunoreactive spindle-shaped cell exhibiting intense brown colored cytoplasmic granules. The positive CD44 cells are located in-between tubules or incorporated with the cells of the kidney tubules (Fig. 1).

Sry gene (402 bp) was expressed in male rats from which AD-MSCs were isolated. It was also detected in renal cells of female rats that were injected by male-derived AD-MSCs in MSCs group. The Y-chromosome marker was neither detected in the control group nor in the DN group nor in the CM group. MSC-specific marker CD105 gene (374 bp) was expressed in renal tissue, but CD34 gene (744 bp) was not detected. These results indicate that our AD-MSCs were consistent with recognized MSCs.

### Table 2 Kidney functions assessment in the different studied rats’ groups

| Studied groups (number of rats) | Kidney function tests |
|---------------------------------|-----------------------|
|                                | Serum creatinine (mg/dl) | Serum urea (mg/dl) |
| Control (n = 15)                | 0.63 ± 0.08            | 24.66 ± 2.86       |
| DN (n = 15)                     | 2.17 ± 0.32<sup>a</sup> | 56.84 ± 4.68<sup>a</sup> |
| Fibroblast (n = 15)             | 2.2 ± 0.23<sup>a</sup>  | 60.13 ± 0.9<sup>a</sup> |
| AD-MSC (n = 15)                 | 1.05 ± 0.04<sup>b,c</sup> | 38.40 ± 1.09<sup>b,c</sup> |
| CM (n = 15)                     | 1.00 ± 0.02<sup>b,c</sup> | 35.23 ± 1.02<sup>b,c,d</sup> |

<sup>F</sup><sup>241.42 511.12</sup><br><sup>P</sup><sup>&lt; 0.001 0.001</sup><br>

One-way ANOVA, post hoc tests, <sup>P &gt; 0.05</sup> no significant differences, <sup>P &lt; 0.05</sup> significant differences, and <sup>P &lt; 0.001</sup> highly significant differences

<sup>a</sup>Significant difference vs control group
<sup>b</sup>Significant difference vs DN group
<sup>c</sup>Significant difference vs fibroblast group
<sup>d</sup>Significant difference vs AD-MSC group

**Effect of MSCs/CM on kidney function tests**

Treatment with MSCs/CM showed a significant decrease in serum creatinine and urea levels (<sup>P &lt; 0.01</sup> and <sup>P &lt; 0.05</sup>, respectively) as compared to DN rats or fibroblast injected rats. Table 2 shows the kidney function tests in the studied groups.

**Effect of MSCs/CM on markers of oxidative stress in renal tissue**

MDA was significantly (<sup>P &lt; 0.001</sup>) increased in renal tissue of diabetic nephropathy rats as compared to non-diabetic rats. There was a highly significant (<sup>P &lt; 0.001</sup>) decrease in the levels of GSH and CAT in renal tissue as compared to normal control group. MSC/CM treatment of diabetic nephropathy rats showed a significant decrease (<sup>P &lt; 0.01</sup>) in MDA levels and a significant increase in GSH and CAT (<sup>P &lt; 0.05</sup> and <sup>P &lt; 0.01</sup>, respectively) as compared to DN or fibroblast injected rats (Table 3).

**Effect of MSCs/CM on TNF-α and TGF-β levels in renal tissue**

Diabetic nephropathy and fibroblast injected groups showed a highly significant (<sup>P &lt; 0.001</sup>) increase in inflammatory markers such as renal TNF-α and TGF-β levels as compared to normal control rats’ kidneys. MSC/CM treated rats showed a significant reduction in TNF-α (<sup>P &lt; 0.001</sup>) level and TGF-β (<sup>P &lt; 0.01</sup>) (Table 4). There was an insignificant difference between MSCs and CM group indicating that the action is paracrine.

**Effect of MSCs/CM on apoptotic and anti-apoptotic markers in renal tissue**

Bcl2 gene expression was significantly decreased while Bax, caspase-3, and caspase-8 were significantly increased in the DN or fibroblast groups as compared to the control group (<sup>P &lt; 0.001</sup>). Administration of either MSCs or CM to DN rats has a reversed response as there was highly significant increase in Bcl2 gene expression and decreased Bax, caspase-3, and caspase-8 gene expression (<sup>P &lt; 0.001</sup>). There was also non-significant difference in those genes expression between MSCs/CM and control group (Fig. 2).

**Histological and immunohistochecmical results**

H&E stained sections of the control group showed normal glomeruli lined with visceral and parietal cells with normal Bowman’s space in between. The tubules,
proximal convoluted tubules (PCT) and distal convoluted tubules (DCT), showed normal histological appearance (Fig. 3a). H&E stained sections of the diabetic group showed that the glomeruli appeared to be distorted with expanded Bowman’s space; some of the glomeruli showed foci of hyalinosis and eosinophilic deposits (Fig. 3b). Tubular atrophy with flattening of epithelial cells, broken membranes, and loss of brush border was observed in PCT also; DCT showed disorganization and atrophy with dilated lumina (Fig. 3c). Blood vessels showed marked thickening (hyalinization) with lymphocytic infiltration (Fig. 3d). H&E stained sections of fibroblast group exhibited shrunken distorted glomeruli. The tubules had vacuolated cytoplasm and darkly stained nuclei. Blood vessels showed thickening (hyalinization) (Fig. 3e). H&E stained sections of MScs or CM administration to DN rats revealed marked improvement of renal architecture when compared to the DN group or fibroblast group. The glomeruli and tubules appeared nearly similar to normal control group except that there were minimal focal tubular lesions which involve few tubules.

**Mallory staining findings**
Examination of renal cortex sections from control group showed thin blue collagen fibers around renal glomeruli and tubules (Fig. 4a) while DN group (Fig. 4b) or fibroblast group (Fig. 4c) revealed excess blue stained collagen fibers around tubules and around and within the glomeruli, and foci of interstitial fibrosis were noticed. Sections of MScs or CM administration to DN rats demonstrated thin blue collagen fibers around renal glomeruli and tubules (Fig. 4d, e).

**Periodic acid Schiff reaction (PAS) staining findings**
The control group showed a positive reaction for PAS in the form of the thin basement membrane of Bowman’s capsule and thin tubular basement membrane. Also, a positive reaction was observed in the luminal brush border of PCT (Fig. 5a). The kidney tissue of diabetic group (DN) (Fig. 5b) or fibroblast group (Fig. 5c) showed a thick basement membrane of Bowman’s capsule and diffuse thickening of the tubular basement membrane. Loss of brush border of PCT was also noted. By PAS-stained sections of MScs or CM administration to DN rats revealed improvement in the form of the relatively thin basement membrane of the Bowman’s capsule and the tubules. Most of PCT retained their brush border (Fig. 5d, e).

**The immunohistochemical reaction for VEGF**
The control group showed a strong positive reaction for vascular endothelial growth factor (VEGF) expression in glomeruli and tubules (Fig. 6a). In the diabetic group (Fig. 6b) or fibroblast group (Fig. 6c), faint or no reaction

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### Table 3 Oxidative stress markers in the different studied rats’ groups

| Studied groups | MDA (nmol/g tissue) | GSH (mg/g tissue) | CAT (U/g tissue) |
|----------------|---------------------|------------------|------------------|
| Control (n = 15) | 45.13 ± 7.24         | 95.87 ± 2.13      | 8.62 ± 0.76      |
| DN (n = 15)     | 105.32 ± 6.1 a       | 75.81 ± 2.09 a    | 5.83 ± 0.82 a    |
| Fibroblast (n = 15) | 102.42 ± 1.3 a     | 77.17 ± 1.22 a    | 4.3 ± 0.09 b     |
| AD-MSC (n = 15) | 63.31 ± 6.45 ab c   | 84.89 ± 1.72 ab c | 7.86 ± 0.45 b c  |
| CM (n = 15)     | 60.45 ± 4.15 ab c   | 82.06 ± 2.52 ab c | 6.90 ± 0.35 b c d|
| F               | 572.97              | 355.06            | 197.49           |
| P               | < 0.001             | < 0.001           | < 0.001          |

*One-way ANOVA, post hoc tests, P > 0.05 no significant differences, P < 0.05 significant differences, and P < 0.001 highly significant differences
MDA malondialdehyde, GSH reduced glutathione, CAT catalase

aSignificant difference vs control group
bSignificant difference vs DN group
cSignificant difference vs fibroblast group
dSignificant difference vs AD-MSC group

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### Table 4 TNF-α and TGF-β assessment in the different studied rats’ groups

| Studied groups | Inflammatory markers |
|----------------|---------------------|
|                | TGF-β1 (pg/mg protein) | TNF-α (pg/mg protein) |
| Control (n = 15) | 80.63 ± 10.42         | 18.09 ± 0.86         |
| DN (n = 15)     | 206.17 ± 22.30 a      | 43.14 ± 1.13 a       |
| Fibroblast (n = 15) | 210.12 ± 11.32 a     | 40.22 ± 1.33 a       |
| AD-MSC (n = 15) | 101.05 ± 6.04 ab c    | 25.40 ± 0.29 ab c    |
| CM (n = 15)     | 98.15 ± 5.24 ab c     | 23.92 ± 0.17 ab c    |
| F               | 591.81               | 2132.72             |
| P               | < 0.001              | < 0.001             |

*One-way ANOVA, post hoc tests, P > 0.05 no significant differences, P < 0.05 significant differences, and P < 0.001 highly significant differences

aSignificant difference vs control group
bSignificant difference vs DN group
cSignificant difference vs fibroblast group

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Fig. 2 mRNA gene expression of Bcl2, Bax, caspase-3, and caspase-8 when normalized to β-actin gene in group I (control), group II (DN), group III (fibroblast), group IV (AD-MSCs treated), and group V (CM). Asterisk (*) indicates high significant difference vs control group, number sign (#) indicates high significant difference vs DN group, and commercial at (@) indicates high significant difference vs fibroblast group (P value < 0.001).

Fig. 3 H&E stained renal cortex sections at magnification × 400, bar 25 μm. a Control group shows normal structure of glomeruli (G) with Bowman’s spaces (S) which is lined by visceral (V) and parietal (P) cells. Proximal tubules (PC) and distal tubules (DC) are also showed. b-d Renal cortex of diabetic nephropathy group has distorted glomeruli with expanded Bowman space (arrow). The glomeruli showed foci of hyalinosis and eosinophilic deposits (curved arrow) while tubules show dark stained nuclei (arrows heads). c Some tubules are atrophied (TA); others are dilated (DT), and some are obliterated (OT). d Blood vessels have marked hyalinization (arrow head). Dark nuclei and a vacuolated cytoplasm (arrow) are seen within tubules. e Fibroblast group shows distorted shrunken glomeruli (G); blood vessels showed hyalinosis (arrow), and tubules show dark stained nuclei (arrow head). f ADMSC group shows marked improvement in the glomeruli (G), proximal tubules (PT), and distal tubules (DT). g Culture media group shows the glomeruli (G), proximal tubules (PC), and distal tubules (DC) appear nearly similar to that of the control.
for vascular endothelial growth factor (VEGF) expression in glomeruli and tubules was found. The MSC or CM administration to DN rats revealed a positive reaction for VEGF expression in glomeruli and tubules (Fig. 6d, e).

Morphometric results
The USD, DCD, and AP were highly significantly increased in DN and fibroblast groups compared to the control group ($P < 0.001$). Administration of ADMSCs or CM to DN rats showed highly significant decrease in the USD, PCD, and DCD ($P < 0.001$). No significant statistical difference was found between ADMSCs or CM groups and control group ($P > 0.05$) (Table 5). As regards the PCD, there was a significant increase in DN and fibroblast as compared with that of control ($P < 0.05$). In ADMSC or CM groups, there was a significant decrease in the PCD compared to DN or fibroblast groups ($P < 0.05$); however, no significant difference was found between ADMSC and CM groups ($P > 0.05$).

Discussion
Diabetic nephropathy represents one of the serious complications of diabetes mellitus [44]. There was no uncertainty that MSCs played an active role in the treatment of DN. As the nephrons have a mesenchymal origin, so MSCs are considered as suitable and striking candidates which motivate repair of renal tissue. The stromal connective cells are capable of signaling with differentiation into collecting ducts and nephrons [45]. In this study, MSCs were used to determine their possible anti-inflammatory, anti-apoptotic, and anti-oxidant role in amelioration of renal function in the DN model. Fibroblast cell line was used as the negative control for migration of MSCs. MSC and fibroblast share similar morphology and cell surface markers, so we had to be sure that the therapeutic effects of MSC are due to its unique characteristics and not due to its similarity in some features to fibroblasts, so we gave fibroblasts injection to a DN group. MSCs were actually insinuating themselves into the injured kidneys as detected by CD44 expression in the kidney tissue by immunohistochemistry. Similar finding has been registered by Morigi et al., who reported homing of mice MSCs in the injured kidney [16].

The kidney of DN group showed expanded Bowman’s space and deposition of eosinophilic material plus distorted glomeruli with hyalinosis. These changes go hand in hand with Najafian et al. who reported that the earliest sign in DN is microalbuminuria which occurred with
histological changes such as deposition of extracellular matrix and glomerular basement membrane thickening [46]. Also, Raparia et al. stated that the characteristic structural changes in DN, including thickening of glomerular basement membrane, hypertrophy, and proliferation of the mesangial cell [47]. Tubules in the same group DN group showed atrophy, broken membranes, and loss of brush border of PCT. The above results are in accordance with Kim et al. [48] and Zafar et al. [49].

In the current study, the DN rat injected with fibroblast showed similar biochemical, histological, and morphometric results to DN group. These results support that fibroblast had no effect in improvement of diabetic nephropathy. Histological examination of the DN group after stem cell injection revealed that the histology of renal corpuscles retained partially the normal appearance of the control group. This result is in concomitant with Wong et al. who stated that MSCs can regenerate, differentiate, and/or protect mesangial cells [50]. Also, the tubules retain normal structure, and this is in agreement with Qian et al. [51] and Wu et al. [52] who demonstrated the ability MSC differentiation into tubular cells. So, MSCs have been described and considered as being renotropic [16, 53], and Lv et al. revealed that the reno-protective effect of MSCs is through reduction of inflammation and fibrosis [54].

Fifty percent replacement of proximal tubular cells with donor cells was observed by Ling et al. [55]. Rookmaker et al. declared that BM-MSCs participate in regeneration of the highly specialized glomerular microvasculature by homing to injured glomerular endothelium and differentiate into endothelial cells [56]. They confirmed previous observations that BM-MSCs can replace injured mesangial cells [57] and injured glomeruli [19].

The morphometric analysis of this study exhibited statistically significant differences of USD, PCD, DCD, and AP between DN or fibroblast groups to control group. However, DN rats treated with ADMSC or CM showed a significant reduction in all these parameters compared to DN and fibroblast groups. Administration of ADMSC or CM to DN rats assisted in restoring the morphometric parameters to normal levels. Morphometric results are parallel to histopathological results.

The histological and biochemical results of the CM group showed similar results to the MSCs group. CM enhanced the recovery of renal function in DN, suggesting that MSCs participate in a regenerative cytokine environment. It could be hypothesized that the mechanisms that mediate the defensive effects of MSCs could be paracrine [58]. These results provide a rationale for the potential therapeutic tool for DN treatment because the effects of the administration of the CM from MSCs...
ameliorate the effects of diabetes, and it is possible to avoid cell transplantation in the treatment of DN.

Duffield et al. stated that functional repair is derived from the contribution of BM-DC to regenerative cytokine production [59]. BM-MSCs seemed to contribute to 3 to 22% of cells regenerating renal tubular [60] and glomerular cell populations [61] that is the majority of reparative cells were derived from intrinsic kidney cells.

Oxidative stress plays a chief role in the pathogenesis of DN in both types of diabetes mellitus, type 1 and 2 [62, 63]. Evans et al. demonstrated that the main promoting factor for the development of DN is the increase of oxidative stress, not hyperglycemia itself, and this increase leads to endothelial damage [64]. In the current work, the MSCs and/or CM effectively attenuated the increased levels of lipid peroxidation product (MDA) and improved the decreased concentration of antioxidant parameters such as GSH and CAT in DN. Hyperglycemia contributes to the increased level of advanced glycation end products (AGEs) in diabetic renal

Table 5 Morphometric analysis of urinary space proximal tubules and distal tubules diameters and area percent of connective tissue surrounding the tubules and within the glomeruli in the different studied groups

| Studied groups | Urinary space diameter (USD) (µm) | Proximal tubules diameter (PCD) (µm) | Distal tubules diameter (DCD) (µm) | Area percent (AP) (µm²) |
|----------------|----------------------------------|-------------------------------------|------------------------------------|-------------------------|
| Control (n = 15) | 7.843 ± 1.66                     | 38.34 ± 5.77                        | 21.68 ± 2.76                       | 18.5 ± 2.83             |
| DN (n = 15)     | 14.33 ± 2.54a                    | 44.5470 ± 5.96                      | 30.38 ± 2.65a                      | 36.34 ± 4.48a           |
| Fibroblast (n = 15) | 14.948 ± 2.14a                  | 44.25 ± 5.13                        | 30.63 ± 2.43a                      | 36.43 ± 4.16a           |
| AD-MSC (n = 15) | 8.04 ± 1.32bc                    | 39.21 ± 5.33                        | 20.84 ± 2.57bc                     | 20.55 ± 2.36bc          |
| CM (n = 15)     | 8.66 ± 1.40bc                    | 39.77 ± 5.81                        | 20.32 ± 2.61bc                     | 19.90 ± 2.46bc          |
| F               | 13.048                           | 3.623                               | 44.32                              | 98.41                   |
| P               | 0.000                            | 0.011                               | 0.000                              | 0.000                   |

One-way ANOVA, post hoc tests, P > 0.05 no significant differences, P < 0.05 significant differences, and P < 0.001 highly significant differences

*Significant difference vs control group

*Significant difference vs DN group

*Significant difference vs fibroblast group

Fig. 6 Immunohistochemical staining of VEGF of renal cortex sections at magnification × 400, bar 25 µm. a Control group shows positive reaction for VEGF expression in the tubules and glomeruli. b Diabetic nephropathy (DN) group shows negative reaction for VEGF expression in the tubules and glomeruli. c Fibroblast group shows negative reaction for VEGF expression in the tubules and glomeruli. d AD-MSCs group and e culture medium group show positive reaction for VEGF expression in the tubules (arrow) and glomeruli (arrow head)
glomeruli [65], which plays a crucial role in formation of reactive oxygen species (ROS). STZ also significantly elevates the activity of xanthine oxidase, which is considered as another important cause of ROS [7]. Shah et al. showed that ROS could activate nuclear factor-Kappa beta (NF-κB) which upregulate TGF-β1 in STZ-induced diabetic kidney dysfunction [66].

TGF-β1 is a fibrogenic growth factor, locally produced in the kidney. It participates in progressive renal tubular atrophy by induction of apoptosis in tubular epithelial cells [67]. In this study, the upregulated TGF-β1 expression in DN was blunted by MSCs and/or CM. This finding was proved by Mallory trichrome as excess fibrosis appeared in the glomeruli and around tubules in DN group while this fibrosis was markedly decreased in MSCs and/or CM group. We suggest that MSCs and/or CM could hinder hyperglycemia-mediated and ROS-induced activation of NF-κB signal cascade as well as upregulation of TGF-β1 in DN.

We tried to focus on the role of TNF-α in diabetic circumstances. It can activate cell survival, extrinsic apoptosis signalling pathways, and stimulation of the kidney epithelial cells to secrete chemo-attractant factors and cell adhesion factors [68]. Moreover, it enhances local production of ROS and alteration of the glomerular capillary barrier function with resulting proteinuria [69, 70]. TNF-α expression levels were lower in MSCs and/or CM treated groups as compared with the DN group in the present study.

TNF-α initiates caspase-8 activation in the cytosol through death receptor, TNF-R1. Caspase-8 activates either type I or type II pathways of programmed cell death [71]. Caspase-8, in the pathway of type I, directly activates caspase-3, resulting in apoptosis [72, 73]. In type II pathway, the caspase-8 promotes cell death depending on mitochondria apoptotic pathways via Bcl-2 family protein in the mitochondria [74].

There was a significant elevation in TNF-α levels, caspase-3, and caspase-8 expression levels in DN compared to the control group. MSC and/or CM inhibit this TNF-α apoptotic pathway by decreasing TNF-α levels and caspase-3 and caspase-8 mRNA levels in DN model. Similar results were recorded for TNF-α [75] and TGF-β [76].

ROS enhances oxidative stress with a disturbance in the balance between pro and anti-apoptotic proteins. When the ratio of Bax/Bcl-2 is elevated, the protective and defensive effects of Bcl-2 are disrupted, particularly on the mitochondrial membrane with increased permeability allowing cytochrome c leakage into the cytosol and activate caspase-3 and 9 [77, 78]. In our study, we found the decreased expression of Bax and increased expression of Bcl-2 in MSCs and/or CM therapy of DN.

**Conclusion**

MSC and/or CM have minimized DN supporting the hypothesis that MSCs may exert a paracrine effect on the damaged tissue. These paracrine effects may be mediated mainly by the RNA-like factors released by MSCs into the medium because the protective effect of the CM was blunted by RNase during a long period of incubation. Therefore, in future clinical applications, it may be possible to treat this DN using CM alone, thus avoiding cell transplantations.

**Abbreviations**

AD: Adipose tissue; AGEs: Advanced glycation end products; AP: Area percent; CAT: Catalase; CM: Culture medium; DCD: Distal tubules diameter; DN: Diabetic nephropathy; GSH: Reduced glutathione; MDA: Malondialdehyde; MSCs: Mesenchymal stem cells; NF-κB: Nuclear factor-Kappa beta; NOS: Nitric oxide synthase; PCD: Proximal tubules diameter; ROS: Reactive oxygen species; STZ: Streptozotocin; TGF-β1: Transforming growth factor-β1; TNF-α: Tumor necrosis factor-α; USD: Urinary space diameter; VEGF-A: Vascular endothelial growth factor-

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**Authors’ contributions**

AAI prepared the idea and designed the study; SEA did the data statistical analysis; NR, OA, and NAS performed all the laboratory investigations and interpreted the research data regarding each studied group, and MMM with AAI performed all histological studies. All authors wrote, read, and approve the final manuscript.

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This study has been approved by Faculty of Medicine, Zagazig University Institutional Review Board (IRB) for research studies (reference number is 4920/14-7-2016) and Zagazig University IACUC (approval number is ZU-IACUC/3/F/90/2018) for animal research.

**Consent for publication**

Not applicable

**Competing interests**

All authors declared that there are no competing interests.

**Author details**

1. Human Anatomy & Embryology Department, Faculty of Medicine, Zagazig University, Zagazig, Egypt. 2. Urology Department, Faculty of Medicine, Zagazig University, Zagazig, Egypt. 3. Medical Biochemistry Department, National Research Centre, Cairo, Egypt. 4. Medical Biochemistry Department, Faculty of Medicine, Zagazig University, Zagazig 44512, Egypt. 5. Human Anatomy & Embryology Departments and Animal house of the faculty for facilitating the performance of this work.

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