Research Article

Transfer of Bone-Marrow-Derived Mesenchymal Stem Cells Influences Vascular Remodeling and Calcification after Balloon Injury in Hyperlipidemic Rats

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Bone-marrow-derived mesenchymal stem cells (BM-MSCs) were found to markedly increase atherosclerotic lesion size. The aim of this paper was to investigate whether BM-MSCs contribute to vascular remodeling and calcification after balloon injury in hyperlipidemic rats. Labeled BM-MSCs were found in the lesion of hyperlipidemic rats after balloon injury. Comparing injury group, transferred BM-MSCs significantly triggered vascular negative remodeling, characterized by the changes of remodeling index (0.628 ± 0.0293 versus 0.544 ± 0.0217), neointimal area (0.078 ± 0.015 mm² versus 0.098 ± 0.019 mm²), PCNA index (23.91 ± 6.59% versus 43.11 ± 5.31%), and percentage of stenosis (18.20 ± 1.09% versus 30.58 ± 1.21%). Apparent vascular calcification was detected in medial layers at 6 weeks after balloon angioplasty, which may be associated with upregulation of bone morphogenetic protein-2 (BMP-2). Our data indicated that unselected BM-MSCs transfer may induce vascular remodeling and calcification after balloon injury in hyperlipidemic rats.

1. Introduction

Cellular therapy has been an area of intense research over few decades, which is widely used for treatment of cardiovascular disease. Bone-marrow-derived mesenchymal stem cells (BM-MSCs) were isolated from adult bone marrow, along with the ability for self-renewal and multilineage differentiation into cardiomyocytes, smooth muscle cells, endothelium, osteoblasts, and chondroblasts [1, 2]. Accumulating evidence from a growing body of animal studies demonstrated the extensive capacity of MSCs to engraft, differentiate and produce substantial functional recovery [3]. Transplantation has been considered as a potential clinical strategy for the treatment of ischemic heart diseases and heart failure.

Although a growing body of evidence from animal experiments and clinical trials indicated the beneficial therapeutic potential of MSCs in cardiovascular diseases, the safety of MSCs transplantation is a critical problem that should be kept in mind. The findings that MSCs transplantation could result in tumor formation and alteration of electrophysiological properties caused wide concerns [4, 5]. Furthermore, unexpected severe intramyocardial calcification was elicited by direct transplantation of unselected bone marrow cells into acutely infarcted myocardium [6]. It has been demonstrated that transfer of bone marrow and endothelial progenitor cells accelerated atherosclerosis and influenced the plaque stability [7]. Increasing evidence indicates a link between MSCs therapy and pathogenesis of atherosclerosis [8, 9].

Our previous studies demonstrated that oxidized LDL synergistically promoted osteodifferentiation of bone-marrow-derived MSCs in response to osteogenic inductor. In this study, we investigated the effects of transfer of unselected BM-MSCs on vascular remodeling and calcification in the hyperlipidemic rat.
2. Method

2.1. Preparation of BM-MSCs. Unselected BM-MSCs were extracted from male Sprague-Dawley (SD) rats (150–200 g) femurs by flushing the bone marrow cavities with Dulbecco’s modified Eagle’s medium-low glucose (DMEM) and cultured in DMEM supplemented with 10% fetal bovine serum (FBS). After 24 h, the medium was discarded to remove hematopoietic stem cells and nonadherent cells. Medium was changed every 1–2 days. Cells became almost confluent after 8 days and were trypsinized with 0.25% trypsin containing 1 mM EDTA for 3 min at 37°C. All experiments were performed on cells cultured for up to 3–5 passages. BM-MSCs were identified as CD90-positive and CD34-negative by flow cytometry.

2.2. Animals. All animal experiments were approved by the Animal Care and Use Committee of Fudan University according to 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). Animals were fed with high cholesterol diet containing 10% fat, 5% cholesterol, 0.5% sodium cholate, and 0.2% propylthiouracil received vitamin D₃ (600,000 U/kg body weight) for 1 week before balloon denudation. Female SD rats were randomly allocated to three groups: (1) rats without denudation were fed with high cholesterol diet (Control, n = 8); (2) rats were subjected to balloon denudation and fed with high cholesterol diet (Injury group, n = 8); (3) rats were subjected to balloon denudation and BM-MSCs transplantation plus with high cholesterol diet (MSC group, n = 8). Rats were anesthetized with ketamine (150 mg/kg body weight IP) and heparinized with 100 U/kg heparin sodium before procedures. A well-established rat model of aortic artery balloon injury was used in this study. The abdominal aorta was isolated and stripped of adventitia. A 6 F balloon catheter was introduced through incision under the renal artery bifurcation about 1 cm and advanced into the thoracic aorta. The balloon was inflated with saline and pulled back to the abdominal aorta. After balloon denudation was performed for 3 times, saline or BMSCs labeled by DAPI were injected before removal of the catheter. After balloon denudation, rats were fed with high cholesterol diet containing 10% fat, 5% cholesterol, 0.5% sodium cholate, and 0.2% propylthiouracil for 6 weeks after balloon denudation. In addition, rats without denudation were fed with a normal diet to compare with those with a high cholesterol diet.

2.3. Analyses of Plasma Lipid. After feeding, the rats were fasted overnight and the samples were collected and centrifuged at 3000 rpm for 15 min. The plasma lipid profile was determined by measuring the contents of total cholesterol (TC) and low-density lipoprotein (LDL) by standard enzymatic procedures using reagent kits.

2.4. Histology. Paraffin sections of vascular tissue were stained with hematoxylin-eosin for morphometric analysis as previously described [10]. Histomorphometric parameters, including luminal area (LA), internal elastic lamina area (IELA), external elastic lamina area (EELA), and neointimal area were defined and determined using the software of Leica QWin Plus as previously described [11]. The chronic variation of the area surrounded by the external elastic lamina was evaluated by remodeling index compared with a reference normal segment upstream from the lesion site, which was calculated as a percentage as follows: EELA(lesion)/EELA (reference). The percentage of stenosis was calculated as (IELA–LA/IELA) × 100.

2.5. Von Kossa Stain. Von Kossa stain (Sigma-Aldrich, USA) was performed to evaluate abnormal deposits of calcium within vascular wall. Paraffin-fixed tissue sections were incubated with silver nitrate solution under a strong light. The calcium was replaced with silver deposit, visualized as metallic silver. Procedural details are described elsewhere [6].

2.6. Immunohistochemistry. Proliferative activity was determined in vascular tissue using analysis of proliferating cell nuclear antigen (PCNA) expression defined immunohistochemically. In brief, tissue sections were quenched for endogenous peroxidase and antigen-retrieved in 0.01 M citrate buffer (pH 6.0) for 15 min in a microwave oven at 100°C. The sections were blocked with normal goat serum and then incubated with mouse anti-PCNA antibody (Boster Company, China) overnight in a humid chamber at 4°C. After incubation with biotinylated goat anti-mouse IgG, sections were visualized by using avidin-biotin horseradish peroxidase visualization system. Sections were then counterstained with haematoxylin. Images were recorded using Leica Qwin Plus.

2.7. Immunofluorescence Staining. Immunofluorescence staining was performed according to standard procedures. Briefly, vascular tissue sections were deparaffinized and hydrated. After preincubation with normal goat serum, sections were incubated with rabbit anti-BMP-2 antibody (Boster Company, China) overnight in a humid chamber at 4°C followed by incubation with appropriate FITC-linked secondary antibody. Images were captured by using Leica Qwin Plus.

2.8. Statistical Analysis. Data were presented as means ± SEM. Comparison was performed using one-way analysis of variance (ANOVA) with Tukey test for post hoc analysis to determine the difference among groups. P < 0.05 was considered statistically significant.

3. Result

3.1. Identification of BM-MSCs Prepared for Transplantation. For transplantation, unselected BM-MSCs were isolated from bone marrow according to standard technique. A few small, bright, round cells were present in the primary culture, which gradually disappeared by the second generation or later (Figure 1(a)). BM-MSCs were characterized as
Figure 1: Identification of BM-MSCs and the profile of lipid level in cholesterol-rich fed rats. Phase contrast microscopy depicted that a few small, bright, round cells were present in the primary culture (a). Phase contrast microscopy revealed that BM-MSCs were characterized as a fibroblast-like spindle-shaped morphology (b). Baseline of serum lipid content before received cholesterol-rich diet did not significantly differ among three groups. Cholesterol-rich diet significantly increased total serum cholesterol (c), and LDL (d) levels after angioplasty in rats. The data were expressed as means ± SEM. n = 8 per group. *P < 0.05 versus normal diet group.

a fibroblast-like spindle-shaped morphology (Figure 1(b)). More than 95.69% of cells isolated from bone marrow were positive for CD90, and negative for the early hematopoietic marker CD34, indicating specific phenotype of BM-MSCs markers.

3.2. The Profile of Lipid Level in Rats with Cholesterol-Rich Diet. Baseline of serum lipid content before received cholesterol-rich diet did not significantly differ among three groups. The addition of cholesterol to the diet significantly increased total serum cholesterol, and LDL levels. Procedures of balloon angioplasty and BM-MSCs transplantation did not affect plasma lipid levels in this study (Figures 1(c) and 1(d)).

3.3. Effect of BM-MSCs on Vascular Remodeling after Balloon Angioplasty. To address vascular remodeling in vivo, histological analysis of vascular wall was assessed at 6 weeks after balloon angioplasty. Ruptured internal elastic lamina was present at the site of lesion, whereas splitting of the intima was replaced with significant neointimal formation after balloon angioplasty in injury group and MSC group. Data from histomorphometric parameters depicted that LA, IELA, EELA, and remodeling index significantly decreased after balloon angioplasty (LA: 1.632 ± 0.043 mm² versus 1.335 ± 0.056 mm² versus 1.133 ± 0.074 mm²; IELA: 1.632 mm² ± 0.043 mm² versus 1.414 mm² ± 0.073 mm² versus 1.185 mm² ± 0.086 mm²; EELA: 2.403 mm² ± 0.067 mm² versus 1.509 ± 0.058 mm² versus1.308 ± 0.044; remodeling index: 1 versus 0.628 ± 0.0293 versus 0.544 ± 0.0217; control versus injury group versus MSC group, resp., P < 0.05), whereas significant neointimal area and percentage of stenosis were increased, compared with control (neointimal area: 0 versus 0.078 ± 0.015 mm² versus 0.098 ± 0.019 mm², P < 0.05; percentage of stenosis: 0 versus 18.20 ± 1.09% versus 30.58 ± 1.21%; control versus injury group versus MSC group, resp., P < 0.05). BM-MSCs transplantation significantly aggravated vascular negative remodeling when compared with injury group, characterized by the decrease of LA, IELA, EELA, and remodeling index and increased neointimal area and percentage of stenosis (Figure 2).
3.4. Effect of BM-MSCs on Neointimal Formation after Balloon Angioplasty. To investigate the neointimal formation in vivo, the PCNA index was evaluated. PCNA-stained nuclei were detected in the neointimal and medial layers of vascular wall at 6 weeks after vascular denudation (Figure 3). Data demonstrated that BM-MSCs resulted in a significant increase in the neointimal area (0.078 ± 0.015 mm² versus 0.098 ± 0.019 mm², $P < 0.05$), PCNA index (23.91 ± 6.59% versus 43.11 ± 5.31%, $P < 0.05$), and percentage of stenosis (18.20 ± 1.09% versus 30.58 ± 1.21%, $P < 0.05$) compared with injury group.

3.5. Effect of BM-MSCs on Vascular Calcification after Balloon Angioplasty. To further characterize vascular calcification in
lesions, von Kossa staining was performed. Typical of calcification, revealing black metallic deposit, was obviously visualized in medial layers in MSC group, compared control and injury group (Figure 4(a)). In view of association between bone morphogenetic protein-2 (BMP-2) and vascular calcification, the expression of BMP-2 was determined by Immunofluorescence staining. Results indicated that BMP-2 was strongly expressed in the neointimal and medial layers of vascular wall (Figure 4(b)). These results indicated that BM-MSCs-induced vascular calcification might be associated with upregulation of BMP-2. To track the fate of BM-MSCs, BM-MSCs were prelabeled by DAPI. Labeled BM-MSCs were homing in the intimal layers at 6 weeks after angioplasty, indicating a role of BM-MSCs in the pathological changes of vascular remodeling and calcification (Figure 4(c)).

4. Discussion

The salient findings from our current study indicated that BM-MSCs transplantation triggered vascular remodeling and calcification after balloon angioplasty in hyperlipidemic rats. These results highlighted the potential importance of complication caused by BM-MSCs which should keep in mind in the therapeutic application.

Up to now, clinical trials for the treatment of ischemic heart diseases and heart failure using bone marrow cells were rapidly increased. Cell therapy has been regarded as a potential clinical strategy for cardiovascular diseases. However, intensive concerns about deleterious effects received significant attention, such as tumorigenicity, arrhythmia, and unregulated differentiation. Studies from Losordo reported for the first time that severe intramyocardial calcification was detected after transplantation of unselected BM cells in acute myocardial infarction [6]. The vascular remodeling and calcification were observed after balloon injury in hyperlipidemic rats. It is plausible that unselected BM-MSCs therapy could lead to unexpected complication in coronary heart disease patients with hyperlipidemia. Actually, patients with coronary heart diseases complicated by hyperlipidemia were clinically prevalent. Particularly, intracoronary delivery of BM-MSCs was performed after successful percutaneous coronary intervention in most of clinical trials. Thus, the gross adverse effect observed in the study could be clinically significant.

Increasing evidence indicated a link between bone-marrow-derived cells and the pathogenesis of atherosclerosis. Circulating myeloid calcifying cells were identified and found to contribute to ectopic vascular calcification in type 2
diabetes [12]. Previous studies demonstrated that transferred BM-MSCs were homing to the vascular lesion and markedly increased atherosclerotic lesion size [7]. In this study, the fate of transferred BM-MSCs were traced and found to present at a low density in the vascular sections. However, the homing BM-MSCs may be underestimated, as quenching of the DAPI cannot be completely excluded at 6 weeks after balloon injury. It was indicated that BM-MSCs were potentially involved in atherosclerotic calcification.

The mechanisms by which transferred BM-MSCs modulate vascular calcification remain unclear. Compelling evidence suggested that vascular calcification was regulated by bone calcification regulatory factors locally expressed in blood vessels [13–15]. Paracrine BMP-2 caused the induction of osteoblast-like cells in intima and accelerated the atherosclerotic calcification in vivo [16]. BMP-2 was found to promote vascular calcification via increased phosphate uptake and induction of osteogenic phenotype modulation in smooth muscle cells [17]. Our observation that strongly expression of BMP-2 was detected in the neointimal and medial layers indicated a potential role of BMP-2 in the BM-MSCs-induced calcification. Our data demonstrated that ox-LDL was synergistically promoted osteodifferentiation of BM-MSCs co-cultured with vascular smooth muscle cells in the presence of osteogenic inducer (unpublished data). It was suggested that BM-MSCs together with balloon injury and hyperlipidemia contribute to the formation of vascular calcification.

Figure 4: Effect of BM-MSCs on vascular calcification after balloon angioplasty and the fate of transferred BM-MSCs. (a) Von Kossa staining was performed to detect vascular calcification in lesions black metallic deposit was obviously visualized in medial layers in MSC group, compared control and injury group. (b) Immunofluorescence staining was performed to determine the expression of BMP-2. Results indicated that upregulated BMP-2 was strongly expressed in the neointimal and medial layers of vascular wall in MSC group, compared Control and Injury groups. (c) Labeled BM-MSCs were homing in the intimal layers at 6 weeks after angioplasty. \( n = 8 \) per group.
Taken together, we demonstrated an important role of transferred BM-MSCs in vascular remodeling and calcification. These findings call for caution in potential adverse effects with respect to BM-MSCs transfer in the future clinical trials.

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Authors’ Contribution

J. Liao and X. Chen contributed equally to this work.

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