Research Article

Conditioned Medium from Adipose-Derived Stem Cell Inhibits Jurkat Cell Proliferation through TGF-β1 and p38/MAPK Pathway

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Received 6 March 2019; Revised 10 August 2019; Accepted 26 August 2019; Published 19 December 2019

Academic Editor: Sebastião Roberto Taboga

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Background. Since the first report on the immunomodulatory and immunosuppressive properties of Adipose-Derived Stem Cells (ADSCs), many studies have elucidated the underlying molecular mechanism of their suppressive activity on mixed lymphocyte reaction (MLR). However, a gap exists in our understanding of the molecular mechanism of ADSC-conditioned medium (ADSC-CM) on MLR.

Methods. ADSCs were isolated from Human Adipose Tissues, and Enzyme-linked Immunosorbent Assay (ELISA) was used to identify the concentration of transforming growth factor β1 (TGF-β1) in ADSC-CM. The transcript abundance of TGF-β1, as well as that of insulin-like growth factor binding protein 3 (IGF-BP3), was evaluated using qRT-PCR on Jurkat cells cultured in ADSC-CM for 24 hours. The proliferation of the Jurkat cells was assessed using cell cycle assay. Western blotting was performed to identify potential signaling molecules involved in the ADSC-CM-induced inhibition of Jurkat cell proliferation.

Results. The findings confirm that the isolated ADSCs demonstrate classic ADSC characteristics. The level of TGF-β1 was found to be low in ADSC-CM, as assessed by ELISA. Jurkat cells grown in ADSC-CM show reduced gene expression of TGF-β1 and IGF-BP3 compared with that of the control group. Furthermore, western blotting of ADSC-CM grown Jurkat cells that were blocked at the G0/G1 stage indicates that ADSC-CM decreases the protein expression of pP38 in a dose-dependent manner.

Conclusion. ADSC-CM can inhibit Jurkat cell proliferation through the TGF-β1-p38 signaling pathway.

1. Introduction

Mesenchymal stem cells (MSCs) are recognized for their immunomodulatory abilities [1–3]. ADSCs, due to their easiness of isolation from an ample tissue source, are a favorable source of MSCs used for the treatment of an extensive variety of diseases [4]. ADSCs do not show the main histocompatibility complex-II expression [5], and their immunosuppressive actions are regulated by prostaglandin E2 (PGE2) [6]. Both preclinical and clinical studies have revealed that allogeneic transplantation of ADSCs is capable of regulating graft-versus-host diseases [7, 8]. Studies have documented that many targeted diseases exhibit significant improvements after ADSC xenotransplantation [9].

Nuclear factor-κB (NF-κB) exists as a constituent of dormant cytoplasmic complexes that are attached to the members of the inhibitor of κB (IκB) family and plays a crucial and evolutionarily role in immunity [10]. Previous studies have shown that ADSCs can modulate T and dendritic cell bioactivity through the NF-κB signaling pathway, in a manner similar to that of the MSCs [11]. Furthermore, paracrine secretions by ADSCs, including cytokines such as IL-10, PGE2, and indoleamine-2,3-dioxygenase (IDO), also have an important role in immunity [6]. Although much research has been focused on the immunosuppressive activity of ADSCs, their effect on the molecular changes during MLR has remained unexplored. Thus, the current study demonstrates the effect of ADSC-CM on Jurkat cells.
2. Materials and Methods

2.1. Isolation and Cultivation of Human ADSCs. Human adipose tissue samples were collected from donors who underwent liposuction. The acquired adipose tissues were minced 3 times using PBS and digested with 0.1% collagenase IV (Roche Diagnostic, Germany) for 1 h. Next, the suspension obtained was centrifuged to obtain the ADSCs. The ADSCs were cultured in Dulbecco’s modified Eagle medium (DMEM) comprising 10% fetal bovine serum, 100 U/mL penicillin/streptomycin (Gibco, USA) at 37°C, in a 5% CO₂ incubator. Once the ADSCs reached 80–90% confluence, they were incubated in DMEM/F12 medium for 24 h. After this incubation, the medium was collected, centrifuged at 300 × g for 5 min, and then stored at −80°C. The culture medium of the cells at passages 2–6 was used for the rest of the experiments. Approval was obtained from the Ethics Committee of the School of Medicine at Shanghai Jiao Tong University. Informed consent was obtained from all the patients.

2.2. Jurkat Cells. Jurkat cells, purchased from ATCC, were cultured in RPMI medium comprising 10% fetal bovine serum at 37°C in a humidified incubator containing 5% CO₂.

2.3. Flow Cytometry. Following harvesting and washing 3 times in PBS, human ADSCs were incubated with the following FITC-conjugated antibodies: CD29, CD44, CD45, CD90, CD105, CD31, and CD34 (Santa Cruz Biotechnology, USA) at 37°C for 30 minutes in the dark. Flow cytometry was used for detection (BD Biosciences, USA).

2.4. Adipogenic and Osteogenic Differentiation. Human ADSCs (10⁵ cells/well) at passages 2–6 were sowed in 0.1% gelatin-coated six-well plates (Cyagen Bioscience, China) and were permitted to achieve 80–90% confluency. Adipogenic differentiation was prompted by growing cells in a medium containing 0.5 μmol/L dexamethasone, 0.5 mmol/L 3-isobutyl-1-methylxanthine, 0.1 mmol/L rosiglitazone, and 100 IU insulin for 2 weeks (Cyagen Bioscience). Osteogenic differentiation was accomplished by growing cells in a medium containing 0.1 μmol/L dexamethasone, 50 μmol/L ascorbic acid, and 10 mmol/L β-glycerophosphate for 3 weeks (Cyagen Bioscience). The medium was changed every 3 days. At the end of differentiation, 4% paraformaldehyde in PBS was used to fix the cells for 15 min at room temperature. Next, the cells were stained using Oil Red O and Alizarin Red S, following the supplier’s protocol.

2.5. Indirect Coculture Conditions. In order to formulate the conditioned medium, human ADSCs from 6 individuals were cultured in DMEM/F12 medium for 24 hours. The cell-free conditioned medium (CM) was centrifuged, carefully removed, and stored at −80°C until further use. The control group of cells was cultured in DMEM/F12 medium for further experiments. Further, 25%, 50%, and 75% cell-free CM in DMEM/F12 were used as the experimental groups.

2.6. ELISA. The quantity of TGF-β1 in the conditioned medium was determined using an ELISA kit, following the supplier’s protocol (R&D Systems, USA).

2.7. Quantitative Real-Time Polymerase Chain Reaction (qRT-PCR). qRT-PCR was performed as previously described [12]. In brief, total RNA was extracted from Jurkat cells following 24 hours of culture with various concentrations of ADSC-CM, using an RNA isolation kit (Takara Bio, Japan). The A260/A280 ratio was used to evaluate RNA purity, which was between 1.8 and 2.0. The following primers were used for gene amplification: TGF-β1 → forward AAG GAC CTC GGC TGG AAG TG and reverse CCG GGT TAT GCT GGT TGT A; IGF-BP3 → forward CCC TCA ACC AAG AAT G and reverse AAC AAG TAG GAC TCC ACC TT. In order to define relative gene expression, the outcomes of 3 autonomous reactions were used. GAPDH expression was used to normalize the findings obtained. Results from three independent experiments each from three technical replicates were used.

2.8. Cell Cycle. 24 h after growth in ADSC-CM, Jurkat cells were collected and washed once using PBS, then fixed in 70% alcohol, overnight. Cell cycle analysis was conducted using the Cell Cycle Kit (Qihai Biotechnology, China), as per the supplier’s protocol. Flow cytometry analysis was conducted using a flow cytometer (Beckman Coulter) and ModFit LT v2.0 software. Results from three independent experiments each from three technical replicates were used.

2.9. Western Blotting. Total protein was extracted from Jurkat cells after 24 or 48 h of coculture with RIPA lysis buffer, as previously published [13]. The nuclear proteins were extracted via NE-PER Nuclear and Cytoplasmic Extraction Reagents (Pierce Biotechnology, Rockford, USA) from the control and experimental groups. Protein samples (40 μg each) were loaded on SDS-polyacrylamide gel, electrophoresed, and then transferred onto PVDF membranes. The protein bands were captured using an enhanced chemiluminescence (ECL) detection kit (Amersham Biosciences, UK). The following primary antibodies were used: phospho-Akt, Akt, phospho-Erk, Erk1/2, phospho-JNK, JNK, phospho-p38, and p38, while GAPDH was used as the loading control. Results from three independent experiments were used.

2.10. Statistical Analysis. All data are displayed as the mean ± standard deviation (SD). Statistical analyses were conducted using SPSS version 19.0. Statistical significance was calculated using one-way ANOVA. Post hoc analysis was conducted using Tukey’s multiple comparison test. A p value of <0.05 was considered to reflect statistical significance.

3. Results

3.1. Characterization of the ADSCs. As previously published [14], the human ADSCs indicated positive staining for the following mesenchymal stem cell surface markers: CD29 (100%), CD44 (99.4%), CD105 (85.2%), and CD90 (99.8%), while they indicated negative staining for the following hematopoietic stem cell surface markers: CD31 (0.1%).
CD34 (0.1%), and CD45 (0.1%) (Figure 1(a)). The human ADSCs also revealed a classic fibroblast-like morphology (Figure 1(b)). We inspected the multipotent differentiating capability of ADSCs using both adipogenic and osteogenic assays. ADSCs differentiate into an adipogenic phenotype when grown in an adipogenic medium for 2 weeks, as determined through exposure to Oil Red O staining (Figure 1(c)). When cultured for 3 weeks in an osteogenic medium, the ADSCs exhibit mineralization, which was evident after staining with Alizarin Red S. This indicates the existence of calcium deposits (Figure 1(d)). These outcomes indicate that the isolated ADSCs present typical ADSC characteristics.

3.2. Scarce TGF-β1 Expression in ADSC-CM. As previous studies have reported, TGF-β1 plays a vital role in cell proliferation, growth, and differentiation [15]. Based on our experimental results (Figure 2), the concentration of TGF-β1 was found to be only 72.5 ± 5.88 pg/mL, which is too low to have any effect on the cell bioactivity of the secretions from ADSCs into ADSC-CM.

3.3. ADSC-CM Decreases the Gene Expression of TGF-β1 and IGF-BP3 in Jurkat Cells. IGF-BP3, an IGF binding protein can bind to the TGF-BV receptor (TβR-V) and mediate TGF-β-induced growth inhibition in concert with the TGF-β1 receptor (TβR-I) and TGF-βII receptor (TβR-II) [16]. The results of this study indicate that ADSC-CM can sugges-

![Figure 1](image1.png)

**Figure 1:** Categorization of human ADSCs. (a) Representation of ADSCs using flow cytometry. ADSCs powerfully express CD29, CD44, CD90, and CD105, but did not express CD31, CD34, or CD45. (b) ADSCs display a fibroblast-like morphology. Cells were promoted to differentiate into adipocytes (c) and osteoblasts (d); bars = 100 μm.

![Figure 2](image2.png)

**Figure 2:** Concentration of TGF-β1 in ADSC-CM. ELISA was performed to evaluate TGF-β1 levels.

![Figure 3](image3.png)

**Figure 3:** ADSC-CM Represses Cell Proliferation. Following 24 h of culturing Jurkat cells in the conditioned medium, the number of cells in the G0/G1 phase was found to have increased, along a decrease of cells in the G2/M phase (Figure 4).

3.4. ADSC-CM Decreases the Protein Expression of Jurkat Cells. The phosphorylation levels of p38 MAPK decreased moderately after culture with Jurkat cells, at different concentrations of ADSC-CM. However, no noteworthy effect
on the stimulation of Akt, ERK1/2, and JNK was observed (Figure 5).

4. Discussion

The restoration or substitution of injured organs remains to be an imperative and perplexing community health issue. Organ grafting may be the most effective method of curing defective or damaged organs. However, graft rejection continues to be the biggest obstacle for successful and permanent transplantation [17]. There have been a substantial number of research studies that have concentrated on the approach of enduring organ graft transplantations and attenuation of immune rejections. The role of lymphocytes, especially T cells, in graft rejection is vigorous, as demonstrated by experiments on nude mice; these mice did not dismiss allogeneic skin grafts since they were deficient in CD4+ as well as CD8+ functionality [18]. Nevertheless, reestablishing this functionality using adoptive relocation of regular T cells triggers the dismissal of the skin graft [18]. Hence, a difficulty in allogeneic grafting is the activating factor of CD4+ and CD8+ T cell immune responses [19, 20]. Dendritic cells (DCs), macrophages, polymorphonuclear cells, angiogenic mediators, and cytokines also stimulate dismissal [21]. Many immunosuppressive agents that are used to suppress graft rejection cannot effectively inhibit immune responses and elicit many side effects [22].

Scientists have tried to identify an effective way to suppress immune reaction over many years. Since the discovery of MSCs, studies have found that MSCs can move to the location of tissue damage, induce inflammation, and exert an anti-inflammatory effect by regulating dendritic cells, natural killer cells (NK cells), T cells, and B cells [23]. MSCs can also promote regulatory T (Treg) cells and preserve their function [24]. ADSCs, which share the same biological properties with MSC, are promising MSCs for combating immune rejection. The first report on immunomodulatory, as well as the immunosuppressive properties of ADSCs, was published in 2005; to be precise, in vitro experimentations have indicated that ADSCs do not aggravate alloreactivity and that they are capable of repressing MLR [25]. Since numerous studies have revealed the prominence of soluble factors, amongst which

Figure 3: ADSC-CM inhibits gene expression of Jurkat cells. qRT-PCR was used to determine the expression of TGFβ1 and BP3 in Jurkat cells cultured in the absence or presence of ADSC-CM for 5 days. Gene expression is shown relative to that of control levels (*p < 0.05).

Figure 4: ADSC-CM inhibits Jurkat cell proliferation. The effect of ADSC-CM on the cell cycle profiles was assessed using flow cytometry. (*p < 0.05).

Figure 5: ADSC-CM interferes with intracellular signaling in vitro. In order to understand the fundamental mechanism of the antiproliferative effect of ADSC-CM, protein levels in Jurkat cells were determined using western blotting.
the utmost recurrently recognized is PGE2, it is believed that the immunosuppressive effects of ADSCs may not require cell-cell contact [26]. Additionally, precise prevention of indoleamine 2,3-dioxygenase or nullification of leukemia inhibitory factor has also been found to eliminate immunosuppression caused by ADSCs [27, 28]. Further, we attempted to find out the mechanism by which the lymphocyte reaction leads to the release of secretomes of ADSCs.

The findings of the present study identified that ADSCs contain CD29, CD44, CD105, and CD90 markers. At the same time, isolated human ADSCs were found to exhibit multilineage differentiation potential and self-renewal ability. Their fibroblast-like morphology was in accordance with that of previous reports [29].

TGF-β1 plays a crucial role in cell proliferation and growth. In order to exclude the effect of TGF-β1, we used ELISA to detect the concentration of TGF-β1 in ADSC-CM. Furthermore, studies show that TGF-β1 does not play a pivotal role in the suppression mediated by the passaged ADSCs [25]. As shown in Figure 2, the concentration of TGF-β1 in the ADSC-CM was found to be extremely low (75 ± 5 pg/mL); thus, it could not have interfered with the results of subsequent experiments. TβR-V serves as the IGF-BP3 receptor that mediates IGF-independent growth inhibition caused by IGF-BP3 in responsive cells. Our results indicate that the gene expression of TGF-β1 in ADSC-CM decreases significantly (Figure 3).

Cell cycle assay was also used to assess cell proliferation and growth conditions. The results of the assay indicate that ADSC-CM-treated Jurkat cells are stuck at the G0/G1 stage of the cell cycle, which corresponds with the RT-PCR results to a certain extent.

Along with analyzing proliferation and growth, we also wanted to identify the possible mechanism involved. Hence, western blotting was used to analyze possible mechanisms underlying the antiproliferative effect of Jurkat cells. Our results reveal that the level of p38 decreases significantly as the concentration of ADSC-CM increases, indicating that phosphorylation of p38 is important in ADSC-CM-mediated immunosuppression. However, the depletion of culture medium components in ADSC-conditioned medium may influence the results. Thus, further studies to explore the certain molecular effect of the proliferation of T cells in CM remain to be resolved.

Organ graft dismissal is an imperative issue in regenerative medicine. The refusal procedure takes place due to stimulation of T cells, either due to the direct, indirect, or semidirect alloantigen recognition cascades, along with the vigorous involvement of accessory B and NK cells, which abolish donor cells. Efforts to induce tolerance, as well as prolong the endurance of transplants, have not been fully successful. Supplementary alternative approaches, including stem cell-based therapies, are known to be more effective. Treatment with stem cells has an abundant potential in the field of bioengineering, due to the ease of obtaining cells for restoration or even replacement of injured tissue or organs. Hence, ADSCs, which are very similar to MSCs, are easy to obtain and show immense therapeutic potential for repairing damaged tissues.

5. Conclusion
To conclude, the results of the current study have revealed that ADSC-CM is able to inhibit Jurkat cell proliferation through the TGF-β1/p38 signaling pathway. However, more functional measurements are required before ADSC-CM can be used as a tolerogenic strategy in the near future.

Data Availability
The flow cytometry, PCR, ELISA, and WB data used to support the findings of this study are included within the article. If there is any other information needed, the corresponding author would provide the information on request.

Ethical Approval
All experiments were approved by the Ethics Committee of Shanghai 9th People’s Hospital, Shanghai Jiao Tong University, School of Medicine.

Conflicts of Interest
There is no conflict of interest.

Authors’ Contributions
Xiu Xia Wang and Yinmin Wang contributed equally to this article and are co-first author.

Acknowledgments
We would like to thank the Laboratory of Tissue Engineering at Shanghai 9th People’s Hospital, Shanghai Jiao Tong University, for providing equipment and technical guidance necessary to complete the work and the support from the Natural Science Foundation of China (81401613).

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