

Umbilical cord-derived mesenchymal stem cells can inhibit the biological functions of melanoma A375 cells

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Abstract. Tumor tropism is an important property of mesenchymal stem cells (MSCs) that has been used in tumor-targeting therapies. However, the effects of MSCs on tumors remain controversial. The aim of the present study was to investigate the effects of MSCs on A375 melanoma cells. Umbilical cord-derived mesenchymal stem cells (UCMSCs) were co-cultured with A375 cells. MTT and Transwell assays were used to assess cell proliferation and invasion, while flow cytometry was performed to detect the apoptosis and the cell cycle distribution of A375 cells. The expression levels of kinases were assayed by western blotting and fluorescence analysis was conducted to detect cytoskeletal rearrangement. The results clearly indicated that UCMSCs could inhibit the proliferation, induce apoptosis and suppress the invasion of A375 cells. Mechanistic studies revealed decreased expression of several kinases (AKT, STAT3 and mTOR) and UCMSCs were also found to promote cytoskeletal rearrangement in A375 cells. These results confirmed that UCMSCs exert anti-tumor effects on melanoma A375 cells.

Introduction

Mesenchymal stem cells (MSCs) are multipotent progenitors that constitute a small proportion of many tissues, including bone marrow, umbilical cord, amniotic fluid, adipose tissue and fetal lung tissue (1). MSCs are described as fibroblast-like cells with the property of plastic adherence during *in vitro* culture. MSCs are able to differentiate into numerous cell lineages (such as adipocytes, osteoblasts, chondrocytes,

tenocytes and cells of visceral mesoderm) in the presence of different conditional media (2). MSCs are positive for the expression of CD105, CD73 and CD90 markers, and negative for the expression of the co-stimulatory factors CD34, CD80, CD86 and HLA-II (3). MSCs also have immunosuppressive capabilities, meaning that they can modulate the proliferation and function of all the components of the immune system. Increasing evidence indicates that MSCs inhibit the maturation of Tregs and dendritic cells and suppress the functions of memory and naïve T cells, as well as CD4⁺ and CD8⁺ T cells. Similar effects were also observed in B cells and natural killer cells (4).

To date, MSCs are considered to be optimal candidates for use in cell therapy based on these advantages. The *in vivo* immunomodulatory properties of MSCs play an important role in the maintenance of peripheral tolerance and the induction of transplantation tolerance, which can protect solid-organ grafts from being rejected (5). Transfusion of MSCs in systemic lupus erythematosus patients has been revealed to improve the levels of serological markers and stabilize renal function without the occurrence of serious adverse events (6). In addition, increasing evidence indicates that MSCs are involved in various diseases, such as type I diabetes (7), liver cirrhosis (8), cerebral palsy (9) and other autoimmune diseases (10). Collectively, these studies indicate that MSCs have potentially significant clinical applications.

However, the effect of MSCs on malignant tumors has remained controversial. Certain studies have indicated that MSCs could inhibit breast tumors (11) and lung cancer (12), whereas other studies have demonstrated that MSCs promote the metastasis of pancreatic (13) and prostate cancer cells (14). In the present study, MSCs isolated from umbilical cord were co-cultured with melanoma A375 cells, and the effect of MSCs on the function of A375 cells was investigated.

Materials and methods

Cell culture. The UCMSCs were purchased from the Sichuan Umbilical Cord Blood Stem Cell Bank (Chengdu, China). Following dissolving in a 37°C water bath, UCMSCs were maintained in Dulbecco's modified Eagle's medium (DMEM;

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Table I. List of primers in quantitative PCR detection.

Gene	Forward primer	Reverse primer	GenBank number
BCL2	CAGGTTATATCTCATCTTTGAG	GTTGAGTAACGAGCTGACCCC	KY098794
Survivin	ACCATAACCCACCACAGC	CAGTTCGTCCCTTTCCAG	DQ508249
MCL1	ATTCCAGAACAGGAGTACAGCTGT	CAGATGTACCCCTTGTTGTAGAGT	NM_021960
Bax	GACTTGAGTTGGGAGGGGAA	GAGGCTCAGCGCCAGGGCTGGG	KJ890756
Fas	TGGCAACGCTGTCCTGTG	CCTTTTGCCAGTAGATGCGAG	KR709619
TNF α	GGTGCTTGTTCCCTCAGCCTC	CAGGCAGAAGAGCGTGCTG	M10988
GAPDH	GAAGGTGAAGGTCGGAGTC	GAAGATGGTGATGGGATTTC	J04038

BCL, B-cell lymphoma; MCL, myeloid cell leukemia; TNF, tumor necrosis factor; GAPDH, glyceraldehydes-3-phosphate dehydrogenase.

Invitrogen, Carlsbad, CA, USA) supplemented with 10% fetal bovine serum (FBS; Invitrogen) at 1×10^5 cells/well in a 6-well plate (BD Falcon; BD Biosciences, Bedford, MA, USA). The medium was changed every two days, and adherent cells were harvested after two weeks using 0.25% trypsin (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA). Only UCMSCs passaged <6 times were used for co-culture since in our previous study (15) we observed that the differentiation of UCMSCs increased dramatically when they were passaged more than 6 times. Melanoma cells A375 [obtained from the American Type Culture Collection (ATCC; Manassas, VA, USA)] were cultured in RPMI-1640 (Gibco; Thermo Fisher Scientific) with 10% FBS. The co-culture ratio of UCMSCs to A375 cells was 2:1.

Cell proliferation assay. Cell proliferation was assessed by an MTT assay at 24, 48 and 72 h following co-culture.

Analysis of apoptosis and cell cycle distribution by flow cytometry. An apoptosis assay was performed by incubating the cells with 3 μ l Annexin V-FITC (Beyotime Institute of Biotechnology, Beijing, China) for 10 min and then counter-staining with 5 μ l PI, and detecting the apoptotic cells by flow cytometry. The cell cycle distribution of the A375 cells was also assessed by flow cytometry.

RNA extraction and quantitative PCR. Total RNA was extracted from A375 cells using an RNeasy Mini kit (Qiagen, Dusseldorf, Germany) at 24, 48 and 72 h post co-culture with UCMSCs. cDNA was synthesized using a ReverTra Ace qPCR RT kit (FSQ-101; Toyobo Co., Ltd., Kagoshima, Japan) with the following transcription conditions: 65°C (5 min), 37°C (15 min) and 98°C (5 min). Real Master Mix (SYBR Green; FP202; Tiangen Biotech, Beijing, China) was used for the qPCR with an iCycler iQ™ Optical Module (Beckman Coulter, Fullerton, CA, USA) under the following conditions: 95°C for 30 sec, followed by 40 cycles at 95°C for 30 sec, 58°C for 30 sec and 72°C for 30 sec. A melt curve analysis followed, consisting of increasing temperatures from 55 to 95°C in 0.5°C increments at 10-sec intervals for 40 cycles. The primers used are listed in Table I. All amplifications were performed three times.

Invasion assay. An invasion assay was performed using 24-well (8- μ m pore size) Transwell plates (Corning, Lowell,

MA, USA). The A375 cells were plated in the upper chambers, which were pre-coated with Matrigel (20%; BD Biosciences, Sparks, MD, USA), while UCMSCs were maintained in the bottom chamber. Toluidine blue (Leagene, Beijing, China) was used to stain the invaded A375 cells at 24 and 48 h post co-culture.

Western blot analysis. Collected A375 cells were washed twice with cold PBS and proteins were extracted using a protein extraction reagent (Pierce, Rockford, IL, USA) containing protease inhibitors (Roche Applied Science, Indianapolis, IN, USA). Protein concentration was assessed using the Micro BCA Protein Assay kit (Pierce). Protein samples (20 μ g) were separated by 12% SDS-polyacrylamide gel electrophoresis and transferred onto nitrocellulose membranes (Invitrogen). Following blocking, the membranes were incubated with primary antibodies (Table II) followed by horseradish peroxidase-conjugated secondary antibodies (1:5,000; cat. no. ab97040; Abcam, Cambridge, UK). Antigen-antibody complexes were visualized using an enhanced chemiluminescence reagent (Amersham Biosciences, Fairfield, CT, USA).

Fluorescence detection. A375 cells were maintained on glass slides and co-cultured with UCMSCs for 48 h. Primary antibodies against E-cadherin and vimentin were applied to the cells for 4 h at room temperature. Following secondary antibody staining, A375 cells were viewed under a confocal scanning microscope (Carl Zeiss, Jena, Germany) at 488 and 525 nm.

Statistical analysis. qPCR and western blotting results were analyzed with Bio-Rad CFX manager software (Bio-Rad Laboratories, Inc., Hercules, CA, USA) and Image Lab software 3.0 (Bio-Rad Laboratories), respectively. Cell cycle distribution was analyzed by ModFit 3.2 (Verity Software House, Topsham, ME, USA). All data are expressed as the mean \pm standard error using Student's t-test. SPSS 19.0 (IBM SPSS Statistics, Armonk, NY, USA) was used for statistical analysis. Values of $P < 0.05$ and $P < 0.01$ were considered to indicate statistically significant differences compared with the control group. All figures were generated with GraphPad Prism 5 (GraphPad Software, Inc., La Jolla, CA, USA).

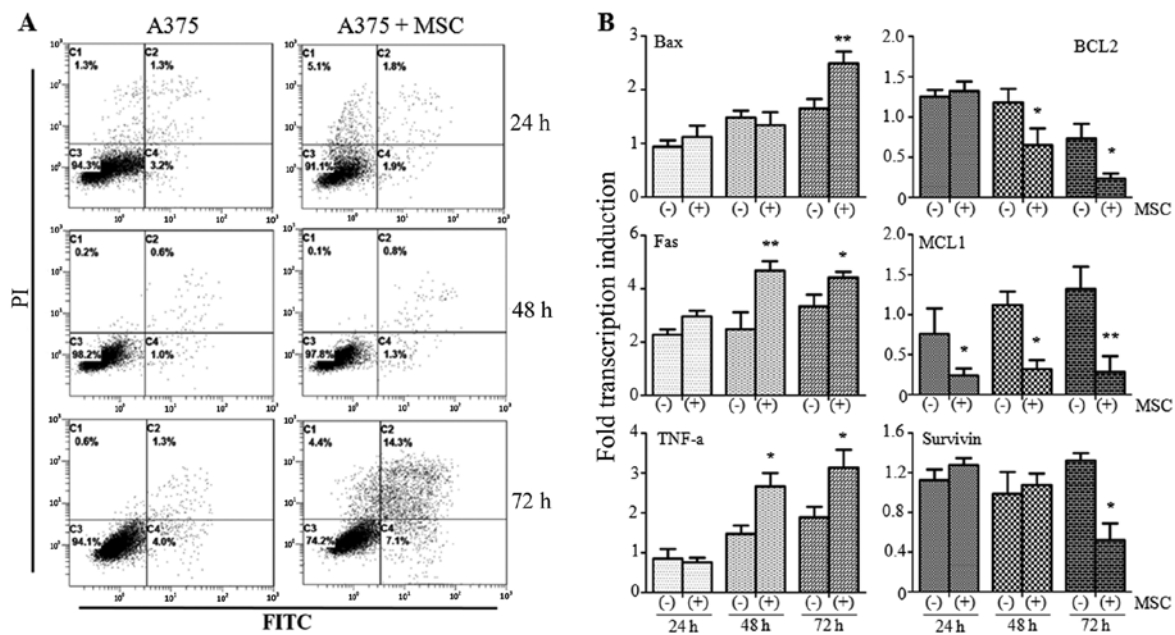


Figure 1. The apoptosis detection of A375 cells co-cultured with UCMSCs. (A) Flow cytometry for the apoptosis assay. (B) Real-time PCR detection of apoptosis-associated molecules. * $P < 0.05$, ** $P < 0.01$ vs. the control. UCMSCs, umbilical cord-derived mesenchymal stem cells.

Table II. The antibodies used in western blot analysis.

Antibody (dilutions)	Catalogue number	Company
AKT (1:1,000)	AF6261	Affinity Biosciences ^a
p-AKT (1:1,000)	AF0016	Affinity Biosciences
PI3K (1:500)	AF6242	Affinity Biosciences
p-PI3K (1:500)	AF3241	Affinity Biosciences
STAT3 (1:1,000)	AF6294	Affinity Biosciences
P-STAT3 (1:1,000)	AF3294	Affinity Biosciences
ERK (1:1,000)	AF0155	Affinity Biosciences
P-ERK (1:1,000)	AF1015	Affinity Biosciences
MTOR (1:1,000)	AF6308	Affinity Biosciences
P-MTOR (1:1,000)	AF3310	Affinity Biosciences
GAPDH (1:1,000)	200608	Zen BioScience ^b

^aAffinity Biosciences, Zhenjiang, China; ^bZen BioScience Co., Ltd., Chengdu, China.

Results

Apoptosis of A375 cells can be induced by UCMSCs. Flow cytometry was performed to detect the occurrence of apoptosis in A375 cells co-cultured with UCMSCs. The results indicated no obvious variation in the apoptotic rate between the A375 monoculture and A375+MSC co-culture groups at 24 h (4.5 vs. 3.7%, respectively) or 48 h (1.6 vs. 2.1%, respectively). However, the apoptotic rate increased markedly in the A375+MSC co-culture group (21.4%) compared with the A375 monoculture group (5.3%) at 72 h (Fig. 1A).

qPCR was then used to detect the levels of apoptosis-associated molecules in the different groups. Among the

apoptosis-promoting factors, the expression of Bax was found to be increased at 72 h ($P < 0.05$) while Fas was markedly increased at 48 h ($P < 0.01$) and 72 h ($P < 0.05$) post co-culture compared with the A375 monoculture group. TNF- α was also found to be increased at 48 and 72 h (both $P < 0.05$) in the presence of UCMSCs. We also investigated three anti-apoptotic molecules: BCL2 was suppressed at 48 and 72 h (both $P < 0.05$) while the expression of MCL1 was inhibited at 24 h ($P < 0.05$), 48 h ($P < 0.05$) and 72 h ($P < 0.01$) post co-culture. Survivin was only decreased after 72 h ($P < 0.05$) (Fig. 1B). Our results indicated that UCMSCs could induce apoptosis in A375 cells.

UCMSCs inhibit the proliferation of A375 cells, however, have no effect on the cell cycle. Subsequently, we detected whether the proliferation and the cell cycle of A375 cells could be influenced by UCMSCs. In the MTT assay, the proliferation did not significantly differ between the A375 monoculture and the A375+UCMSCs co-culture groups at 24 and 48 h. However, the proliferation of A375 cell was markedly inhibited ($P < 0.01$) at 72 h post co-culture with UCMSCs (Fig. 2A). Flow cytometry was then used to assess the cell cycle distribution in various groups of A375 cells. The results indicated that UCMSCs had no effect on the cell cycle at 48 or 72 h post co-culture (Fig. 2B and C).

UCMSCs suppress the invasive ability of A375 cells. As invasion is another important property of malignant tumor cells, we investigated the influence of UCMSCs on the invasive ability of A375 cells in a Transwell chamber pre-coated with Matrigel. As displayed in Fig. 3, the invaded cell numbers showed no obvious difference between the A375 and A375+MSC groups at 24 h post co-culture. However, the invasion of the A375 cells was significantly suppressed in the presence of UCMSCs after 48 h of co-culture.

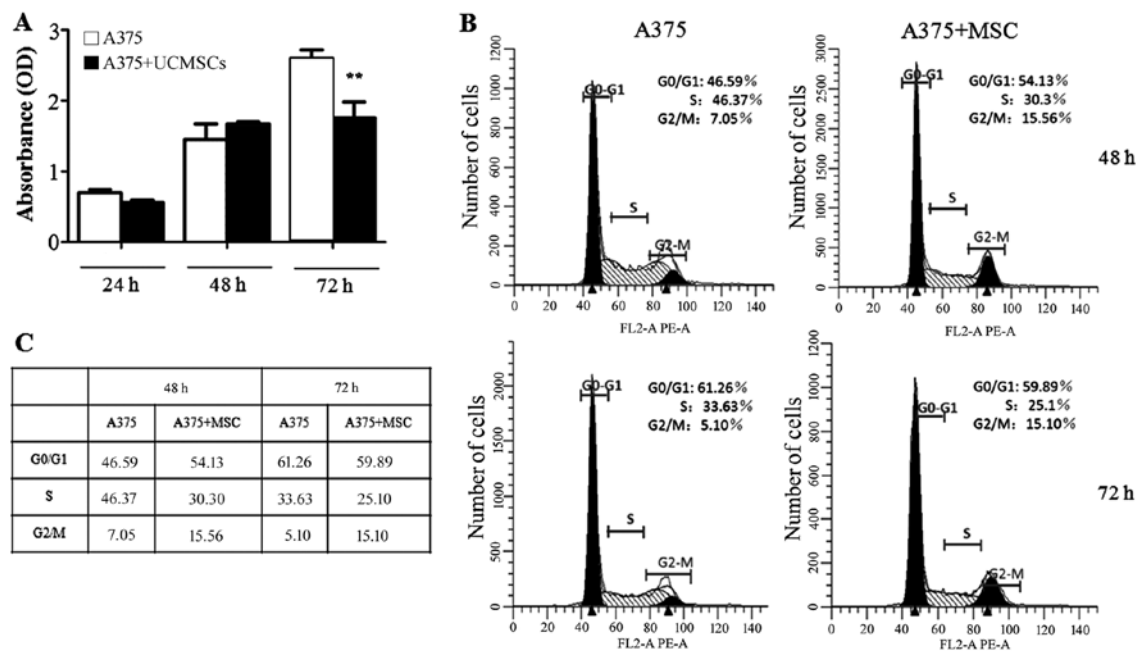


Figure 2. Proliferation and cell cycle assay by flow cytometry. (A) Proliferation assay by MTT. (B) Cell cycle detection (C). Cell cycle analysis. **P<0.01 vs. the control.

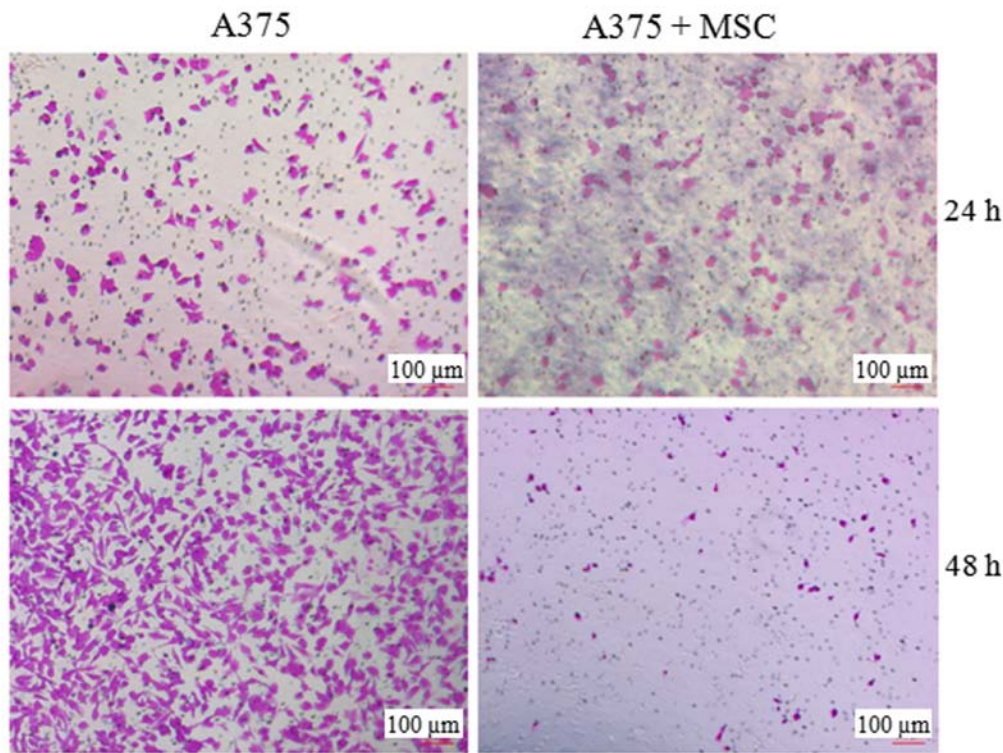


Figure 3. Detection of the invasion of A375 cells co-cultured with UCMSCs. Magnification, x100. UCMSCs, umbilical cord-derived mesenchymal stem cells.

UCMSCs inhibit the expression of several AKT/STAT3/PI3K/mTOR pathway components in A375 cells. To identify the mechanism by which the functions of A375 cells were attenuated by UCMSCs, western blotting was performed to detect the protein expression of several important kinases at 48 and 72 h post co-culture. We found that all tested kinases (AKT/p-AKT, PI3K/p-PI3K, STAT3/p-STAT3, ERK/p-ERK and mTOR), except ERK (P<0.05), demonstrated no significant variation

between the A375 monoculture and A375+MSC co-culture groups at 48 h (Fig. 4). Following 72 h of co-culture, UCMSCs led to the downregulation of the expression of AKT (P<0.05), p-AKT (P<0.01), p-PI3K (P<0.05), p-STAT3 (P<0.01), ERK (P<0.01), p-ERK (P<0.05), mTOR (P<0.05) and p-mTOR (P<0.05) (Fig. 4). These results indicated the important role of the AKT/STAT3/PI3K/mTOR pathway in the regulation of the function of A375 cells by UCMSCs.

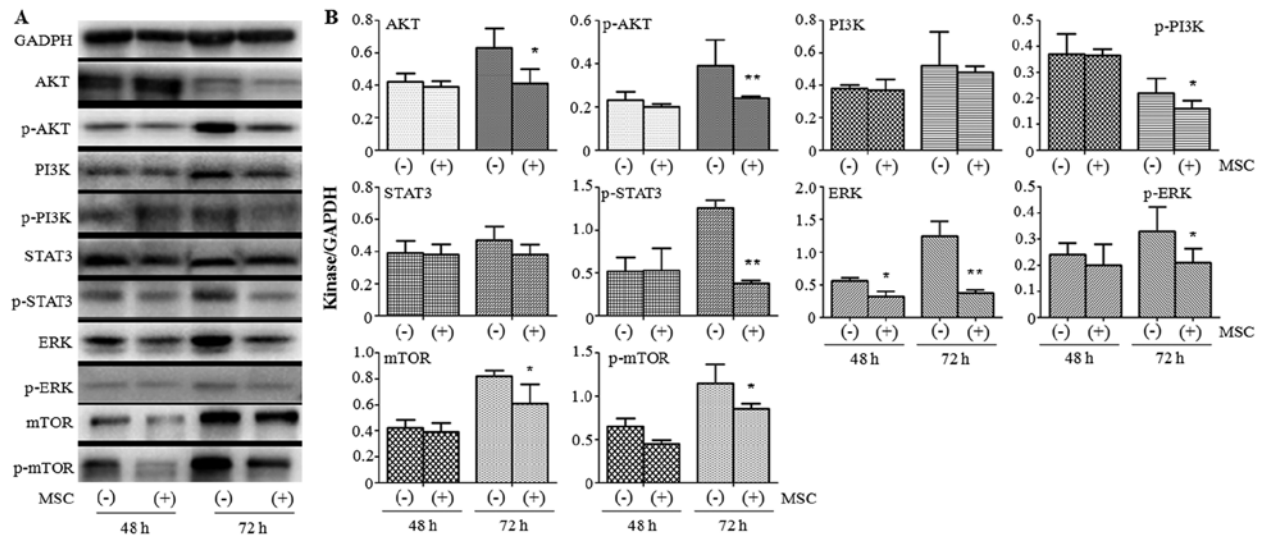


Figure 4. Expression of kinases of A375 cells co-cultured with UCMSCs. (A) Western blot analysis. (B) Data analysis. *P<0.05, **P<0.01 vs. the control. UCMSCs, umbilical cord-derived mesenchymal stem cells.

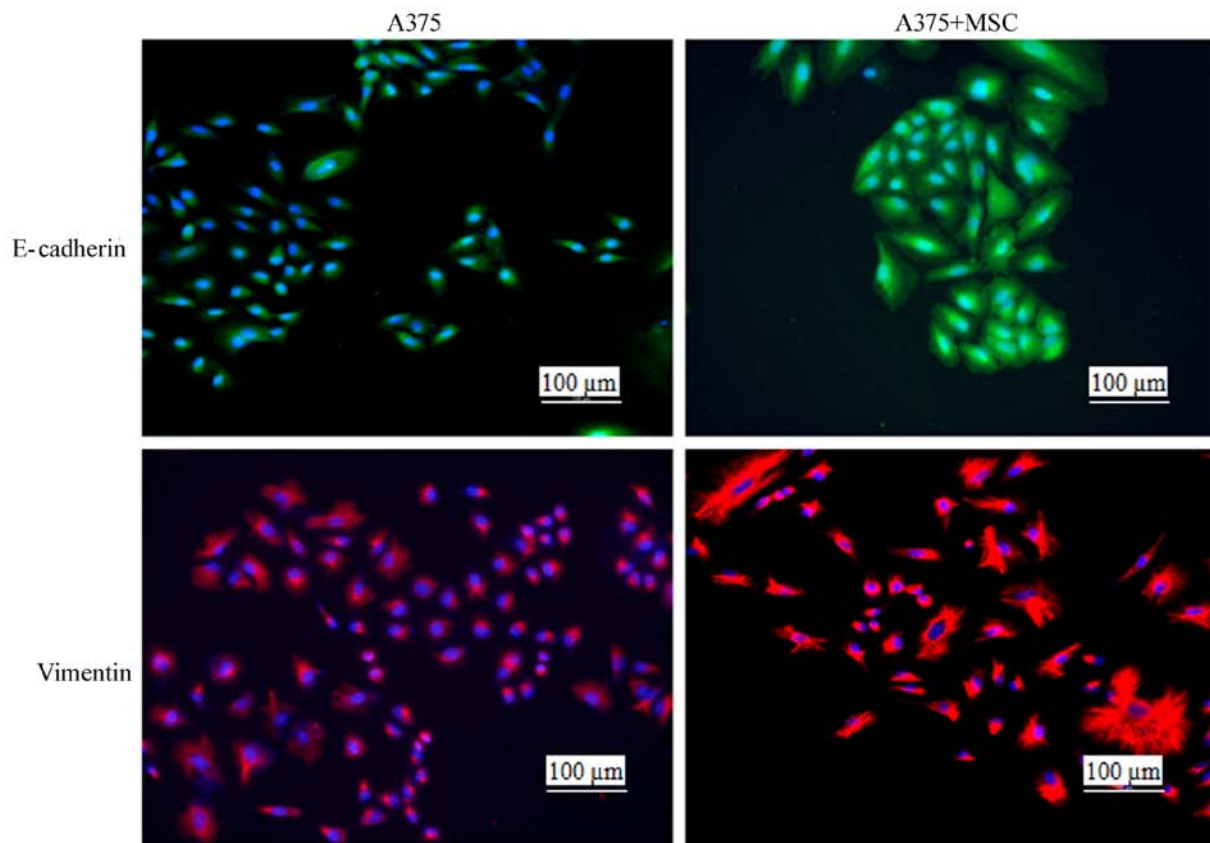


Figure 5. Cytoskeletal rearrangement of the A375 cells co-cultured with UCMSCs. Magnification, x100. UCMSCs, umbilical cord-derived mesenchymal stem cells.

UCMSCs promote cytoskeletal remodeling in A375 cells. The invasion of malignant tumor cells involves decreased adherence and enhanced migration properties, and these changes are part of the epithelial-mesenchymal transition (EMT) (16). During the EMT process, cytoskeletal molecules (including E-cadherin, N-cadherin, laminin and vimentin) play important regulatory roles (17). In the present study, we detected

the occurrence of cytoskeletal rearrangement by staining for E-cadherin and vimentin at 48 h post co-culture with UCMSCs. The results indicated that the cytoskeletal rearrangement was obvious in the A375+MSC co-culture group compared with the A375 monoculture group (Fig. 5), indicating that in the presence of UCMSCs, the invasion of A375 cells may be regulated by certain cytoskeletal molecules.

Discussion

In the present study, we co-cultured melanoma A375 cells with MSCs isolated from umbilical cord to study the role of UCMSCs in regulating the functions of A375 cells. Although UCMSCs had no effect on the cell cycle of A375 cells, this study clearly indicated that UCMSCs inhibited the cell proliferation and invasion, as well as induced the apoptosis of A375 cells. In the mechanistic analysis, the expression levels of AKT/PI3K/STAT3/mTOR pathway components were down-regulated following UCMSC co-culture. Furthermore, marked cytoskeletal rearrangement was observed in the A375+UCMSC co-culture group. Collectively, these results confirmed that UCMSCs can exert an antitumor effect on A375 melanoma cells.

The effect of MSCs on malignant tumor cells remains controversial. The reported discrepancies may be due to the heterogeneity of MSCs or the timing and dose of MSC treatments, or the involvement of other unknown molecules and mechanisms. Liu *et al* (18) reported that UCMSCs could inhibit the growth of human cholangiocarcinoma HCCC-9810 cells in xenograft models, and that UCMSC-conditioned medium suppressed cell proliferation (inhibition rate: 6.21 vs. 49.86%) and induced cell apoptosis (9.3 vs. 48.1%) in a dose- and time-dependent manner. In an immunoblot analysis, the same authors also found that p-PDK1, p-AKT, β -catenin, cyclin-D1 and c-Myc mediated the functional changes of HCCC-9810 cells induced by UCMSCs (18). In another study, pancreatic cancer cells were co-implanted with MSCs in NOD/SCID mice, and it was demonstrated that MSC-derived myofibroblast-like cells could maintain the stemness of tumor-initiating stem cells among pancreatic cancer cells. The mechanistic analysis indicated that the Notch-signaling pathway appeared to contribute to the regulation of stemness by MSCs (13). Wang *et al* (19) explored the effect of the fusion of MSCs with esophageal carcinoma cells, and observed that MSCs markedly decreased tumor cell growth, increased apoptosis and suppressed tumorigenicity. They also observed that the expression of DUSP6/MKP3 in the MAPK pathway increased and the exogenous overexpression confirmed the growth suppression. Sun *et al* (20) engineered MSCs stably transfected with TNF-related apoptosis-inducing ligand (TRAIL) and co-cultured them with hepatoblastoma HepG2 cells directly, or applied MSC-conditioned media to the HepG2 cells. The survival rate of HepG2 cells was markedly decreased by the co-culture conditions, and TRAIL was indicated to serve an important role in promoting cell death. All of these studies revealed the uncertainty regarding the effect of MSCs on malignant tumors.

However, the present study clearly demonstrated the anti-tumor effect of MSCs on A375 melanoma cells in a co-culture system. We not only observed inhibition of proliferation, induction of apoptosis and suppression of invasion, but also further investigated the possible mechanism of the MSC-mediated antitumor effect. Certain important kinases (including AKT, PI3K and STAT3) were downregulated in A375 cells upon MSC co-culture, and this finding was consistent with previous research. Of note was the observation of the rearrangement of the cytoskeleton of A375 cells co-cultured with MSCs. Based on this, we hypothesized that MSCs could inhibit tumor invasiveness by regulating the EMT. However, further research

is required to confirm this hypothesis. The limitation of the present study was that only one cell line was included, since this was just a preliminary study concerning the role of UCMSCs in regulating malignant tumor functions. Our following study will include tumor cell lines from lung, cervical, breast and prostate cancer.

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Availability of data and materials

The datasets used during the study are available from the corresponding author upon reasonable request.

Authors' contributions

WW carried out fluorescence detection, statistical analysis and drafted the manuscript. LL was responsible for completing all the cell related experiments. FC carried out PCR experiments and western blot analysis. YY conceived of the study and participated in its design and coordination and helped to draft the manuscript. All authors read and approved the manuscript and agree to be accountable for all aspects of the research in ensuring that the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate

All experimental protocols have been approved by the Ethics Committee of West China Hospital, Sichuan University (Chengdu, China).

Consent for publication

Not applicable.

Competing interests

The authors state that they have no competing interests.

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