Abstract. Notch signaling, which is driven by the Notch1 receptor, plays an essential role in the pathogenesis and stroma-mediated drug resistance of T-cell acute lymphoblastic leukemia (T-ALL). However, little is known about the roles of Notch ligands in the survival or drug resistance of T-ALL cells. In the present study, isolated mesenchymal stem cells (MSCs) from human umbilical cord (hUC) samples, termed hUC-MSCs, were used as stromal cells for the Jurkat T-ALL cell line. The role of the Notch ligand, Jagged1, was assessed in the survival of Jurkat T-ALL cells using this co-culture system. hUC-MSCs and Jurkat cells were observed to express Jagged1. Furthermore, co-culture with hUC-MSCs led to a significant upregulation of Jagged1 and a more significant overexpression of its receptor, Notch1, in the Jurkat cells, indicating that the receptor and ligand pair may play a role in the reciprocal or autonomous activation of the Notch pathway. In addition, a higher level of CD28 expression was observed in the Jurkat cells that were co-cultured with hUC-MSCs. Blocking Jagged1 expression using neutralizing antibodies restored drug-induced apoptosis in the Jurkat cells that were co-cultured with hUC-MSCs, and also increased the drug sensitivity of the Jurkat cells that were cultured alone. By contrast, direct incubation with exogenously recombinant Jagged1 produced the same protective effects in Jurkat cells as those induced by hUC-MSCs. These results indicate a significant role for Jagged1 in hUC-MSC-induced survival and the self-maintenance of the Jurkat T-ALL cell line, making it a potential target for the treatment of human T-ALL.

Introduction

T-cell acute lymphoblastic leukemia (T-ALL) is an aggressive malignant disease induced by the malignant transformation of T-cell precursors. T-ALL accounts for 10-15% of all leukemias in children and adolescents (1). The molecular mechanisms underpinning T-ALL are likely to be complex (2). A series of studies have demonstrated that the abnormal activation of the Notch1 signaling pathway plays a significant role in the pathogenesis of T-ALL (3,4).

The Notch1 gene encodes a single-pass heterodimeric transmembrane receptor, which has a fundamental function in the development of normal T cells (5). Normally, the activation of Notch signaling is triggered by Notch receptor-ligand interactions. The direct binding of a ligand from a signaling cell to a Notch receptor on the membrane of a receiving cell initiates two successive proteolytic cleavages by the TNF-α-converting enzyme (TACE) and the γ-secretase/presenilin complex. The proteolytic cleavage ultimately results in the release of the Notch intracellular domain (NICD), which translocates into the cell nucleus and interacts with the recombination signal binding protein Jκ (RBP-J). The NICD/RBP-J complex transactivates downstream target genes, including the hairy/enhancer-of-split 1 (Hes-1) gene (6). However, in T-ALL patients, mutations in the Notch1 gene are common and may lead to aberrant activation of the Notch signaling pathway plays a significant role in the pathogenesis of T-ALL (3,4).

The Notch1 gene encodes a single-pass heterodimeric transmembrane receptor, which has a fundamental function in the development of normal T cells (5). Normally, the activation of Notch signaling is triggered by Notch receptor-ligand interactions. The direct binding of a ligand from a signaling cell to a Notch receptor on the membrane of a receiving cell initiates two successive proteolytic cleavages by the TNF-α-converting enzyme (TACE) and the γ-secretase/presenilin complex. The proteolytic cleavage ultimately results in the release of the Notch intracellular domain (NICD), which translocates into the cell nucleus and interacts with the recombination signal binding protein Jκ (RBP-J). The NICD/RBP-J complex transactivates downstream target genes, including the hairy/enhancer-of-split 1 (Hes-1) gene (6). However, in T-ALL patients, mutations in the Notch1 gene are common and may lead to aberrant activation of the Notch signaling pathway that is independent of ligand binding (3). By contrast, the Notch1 proteins in the T-ALL cells also serve as surface receptors that may be triggered by Notch ligands that are expressed by specific cell types, including bone marrow stromal cells. Increasing evidence has suggested that the interaction between tumor cells and the stromal microenvironment results in the resistance to chemotherapy in leukemia and myeloma (7,8). Notch signaling has been shown to be one of the molecular mechanisms involved.
It has been shown that the signaling driven by Notch1 may inhibit apoptosis in developing thymocytes, mature T cells and T-ALL cells (9-11).

In contrast to the roles of the Notch1 receptor, the roles for Notch ligands in T-ALL biology are less clear. The known Notch ligands in mammals include Jagged1 and 2, and Delta-like (DLL)-1, 3 and 4 (6). The actions of these ligands differ in the initiation of Notch signaling and may result in a diverse or opposed biological outcome (12). The present study assessed the role of Jagged1 in the survival of Jurkat T-ALL cells when exposed to a cytotoxic drug, with or without stromal support.

Stromal cells derive from their mesodermal precursors, mesenchymal stem cells (MSCs), which are non-hematopoietic progenitor cells that are located in the bone marrow and a number of other tissues (13,14). Currently, bone marrow is the main source of MSCs. However, the aspiration of bone marrow involves invasive procedures and the yield of bone marrow-derived MSCs (BM-MSCs) decreases significantly with the age of the donor (15). The umbilical cord is an excellent alternative to bone marrow as a source of MSCs for experimental and clinical needs (16). However, data on the application of umbilical cord-derived MSCs is limited. In the present study, human umbilical cord-derived MSCs (hUC-MSCs) were used as stromal cells to evaluate their function in the drug resistance of T-ALL cells.

**Materials and methods**

**Cell culture.** The human T-ALL cell line, Jurkat, was cultured in suspension in RPMI 1640 medium (Gibco BRL, Grand Island, NY, USA) supplemented with 10% fetal bovine serum (FBS; Sijiqing, Hangzhou, China) and 1% penicillin/streptomycin (Gibco BRL). The cells were maintained at 37°C in a humidified chamber with 5% CO2 and routinely subcultured every 2-3 days, ensuring that the cell density in the culture did not exceed 1x10^6 cells/ml.

hUC-MSC cultures were established from the umbilical cords of healthy donors using the direct plastic adherence method after informed consent had been obtained. The study was approved by the ethics committee of the School of Life Science and Biopharmaceutics of Guangdong Pharmaceutical University (Guangzhou, China). Briefly, the umbilical cord samples were sheared into 2-3-cm long segments and minced into 1-2-mm fragments, plated separately in 6-cm polystyrene tissue culture dishes and maintained in DMEM/F12 medium (Gibco BRL) at 37°C in a humidified atmosphere with 5% CO2. The non-adherent tissues were removed on day seven and the culture medium was changed every 3-4 days thereafter. Approximately three weeks later, when well-developed colonies of fibroblast-like cells had appeared (80-90% confluent), the cultures were washed, harvested with 0.25% trypsin (Gibco BRL) and passed through a 100-µm sterile mesh to remove any residual tissue pieces. The filtered cells were then seeded in larger flasks for further expansion. The hUC-MSCs at passages 3-8, displaying a homogeneous mesenchymal immunophenotype and multipotent differentiation potential into adipocytic, osteoblastic and chondrocytic lineages, were used for the experiments.

**Polymerase chain reaction (PCR).** Total RNA was extracted from the hUC-MSCs and Jurkat cells using the TRIzol reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions. Reverse transcription was carried out using the PrimeScript II 1st strand cDNA synthesis kit (Takara, Otsu, Japan) with 1 µg total RNA as a template and oligo dT as a primer. All semiquantitative PCR experiments were performed using the same serially-diluted cDNA batches as templates. Amplification was performed at 95°C for 5 min followed by 38 cycles of 94°C for 30 sec, 56°C for 30 sec and 72°C for 30 sec, then a final extension at 72°C for 7 min. The amplified fragments were analyzed using electrophoresis on a 2% agarose gel. The gene-specific primers that were used for PCR are listed in Table I. The PCR of human β-actin was performed as a control.

**Detection of signaling molecules using flow cytometry.** The Jurkat cells were harvested and prepared for flow cytometry following co-culture with hUC-MSCs. In brief, the hUC-MSCs were plated into 6-well plates at 2x10^5 cells per well to form a confluent monolayer. Following this, 2x10^5 Jurkat cells were added to each well of the adherent hUC-MSCs or cultured alone for 72 h. The co-cultured Jurkat cells were then separated from the hUC-MSCs by careful pipetting with ice-cold PBS. For the flow cytometry, the cells from the various cultures were washed and adjusted to a concentration of 5x10^6 cells/ml in PBS. Aliquots of 100 µl cell suspension were then added into separate tubes. Fc receptors were blocked using the Fc Receptor Blocking reagent (Miltenyi Biotec, Bergisch Gladbach, Germany) for 15 min at 4°C. Surface antibodies were added and incubated for 30 min at 4°C in the dark. The unbound antibodies were removed by washing the cells twice in PBS and the cells were resuspended in 500 µl PBS for the final flow cytometric analysis on a Gallios cytometer (Beckman Coulter, Brea, CA, USA). The antibodies that were used were allophycocyanin (APC)-conjugated anti-CD45 (ebioscience, San Diego, CA, USA), carboxyfluorescein (CFS)-conjugated anti-Jagged1 (R&D systems, Minneapolis, MN, USA), phycoerythrin (PE)-conjugated anti-Notch1 (R&D systems), PE-conjugated anti-CD28 (eBioscience) and non-specific isotype-matched antibodies.

**Apoptosis analysis.** To induce apoptosis, the Jurkat cells were cultured alone or co-cultured with hUC-MSCs for 72 h as described previously and then exposed to dexamethasone (Sigma, St Louis, MO, USA; final concentration 1 µM) for an additional 24 h. The blocking experiments were performed by incubating the hUC-MSCs and Jurkat cells with neutralizing monoclonal antibodies against human Jagged1 (R&D Systems; 1 µg/ml) prior to their inoculation into culture plates. Recombinant human Jagged1 proteins (R&D systems; 1 µg/ml) were used to stimulate the Jurkat cells directly. Apoptotic cell death was detected by Annexin V/propidium iodide (PI) staining using the MEBCYTO apoptosis kit (MBL, Nagoya, Japan). Briefly, the Jurkat cells from the various cultures were harvested, washed and immunolabeled with...
APC-conjugated anti-CD45. The cells were then washed and resuspended in 85 µl binding buffer, followed by incubation with 10 µl Annexin V-FITC and 5 µl PI at room temperature for 15 min in the dark. Following incubation, 400 µl binding buffer was added and the cell samples were measured using flow cytometry.

Statistical analysis. All statistical calculations were performed using the GraphPad Prism software (GraphPad Software, Inc., La Jolla, CA, USA). The data are presented as the mean ± SD. When applicable, Student’s unpaired t-test, a one-way ANOVA and Holm-Sidak tests were used to determine significance. P<0.05 was considered to indicate a statistically significant difference.

Results

Characterization of hUC-MSCs. Fibroblast-like cells were successfully isolated from hUC tissues using the direct plastic adherence method in the present study (Fig. 1A). The cells formed whirlpool-like arrays when a confluent monolayer had developed (Fig. 1A and B). The flow cytometry analysis demonstrated that the hUC-MSCs showed good homogeneity and expressed MSC markers CD73, CD90, CD105, CD44 and CD29, but were negative for CD34, CD45, human leukocyte antigen (HLA)-DR and CD14 (Fig. 2). The same cells showed multilineage differentiation potential, as assessed by culturing in adipogenic, osteogenic or chondrogenic medium (Fig. 3).

Table I. Primers for the PCR analysis.

| Genes      | Primers, 5’-3’ targets, bp |
|------------|---------------------------|
| Notch1     | F: CTACCTGTCAACGTGGCCT    |
|            | R: CGCAGAGGGTTGATTTGCTT  |
|            |                           | 357                          |
| Jagged1    | F: CTCATCAGCCGTTCTCAAC    |
|            | R: GGCACAACACTTTAATCGC    |
|            |                           | 297                          |
| DLL1       | F: TATCCGCTATCCAGGCTGTC   |
|            | R: GGTGGGCAGGTACAGGAGTA   |
|            |                           | 297                          |
| DLL4       | F: AAGGCTGCGCTACTCTTACC   |
|            | R: ATCTCTCTGGTCTTACAGC    |
|            |                           | 538                          |
| Hes-1      | F: ATCACACAGGATCCGGAGCT   |
|            | R: TGACACTGGCGGAGTCGC     |
|            |                           | 300                          |
| β-actin    | F: CTACAATGAGCTGCGTGGG    |
|            | R: CCGTGAGGATCTTCAATG     |
|            |                           | 314                          |

PCR, polymerase chain reaction; F, forward primer; R, reverse primer; DLL, Delta-like; Hes-1, hairy/enhancer-of-split 1.
Expression of Notch ligands by hUC-MSCs. To assess a possible role for the hUC-MSCs in inducing Notch signaling in the Jurkat T-ALL cells, the expression of Notch ligands Jagged1, DLL1 and DLL4 were examined in the hUC-MSCs by PCR using gene-specific primers, with β-actin as an internal control (Table I). This analysis revealed that transcripts for Jagged1 and DLL4 were detected in the hUC-MSCs, while the transcript for DLL1 was undetectable (Fig. 4A). In addition, Jagged1 was relatively highly expressed by the hUC-MSCs at the mRNA level.

Table II. Expression of Notch-related molecules by Jurkat cells cultured alone or co-cultured with hUC-MSCs.

| Jurkat cells | Jagged1 | Notch1 | CD28 |
|--------------|---------|--------|------|
| Alone        | 0.6±0.2 | 1.2±0.5 | 5.7±2.0 |
| Co-culture   | 1.6±0.4 | 2.6±0.7 | 17.6±3.5 |

Student’s t-test P<0.05 P<0.05 P<0.01

The results are expressed as the mean ± SD of the MFI values (n=3) of the Jurkat cells that were cultured alone or co-cultured with hUC-MSCs for 72 h. hUC-MSCs, human umbilical cord-derived mesenchymal stem cells; MFI, mean fluorescence intensity.

Upregulation of Notch1, Jagged1 and CD28 in Jurkat cells following contact with hUC-MSCs. The expression of the Notch-related genes in the Jurkat cells was further analyzed. PCR analysis showed that the Jurkat cells expressed the Notch1 receptor and its ligand, Jagged1 (Fig. 4B), suggesting that the receptor and ligand pair may play a role in T-ALL cells. Hes-1, one of the main downstream molecules of the Notch pathway, was also expressed in the normally-cultured Jurkat cells (Fig. 4B), suggesting that Notch signaling is constitutively active in these cells. Flow cytometry was then used to assess the expression of Notch1, Jagged1 and CD28 in the Jurkat cells. As shown in Fig. 5, at basal conditions, the Jurkat cells expressed CD28 and moderate levels of Jagged1 and Notch1. Notably, following contact with the hUC-MSCs, an upregulation in the expression of all the molecules was observed in the Jurkat cells (Fig. 5; Table II), indicating their involvement in the functional interaction between the hUC-MSCs and the Jurkat T-ALL cell line.

hUC-MSCs inhibit drug-induced apoptosis in Jurkat cells. To study the capability of the hUC-MSCs to support leukemia cell survival, the Jurkat cells were cultured alone or co-cultured with the hUC-MSCs at a 10:1 ratio for 72 h and then exposed to dexamethasone for an additional 24 h. When observed using light microscopy, the Jurkat cells in the co-culture system showed an improved cell morphology compared with those that were cultured alone (Fig. 1B and C). As assessed by Annexin V/PI

Figure 3. Multilineage differentiation capacity of hUC-MSCs. For (A) adipocytic and (B) osteoblastic differentiation, the hUC-MSCs were plated in 6-well plates at 2x10^4 cells per well, treated with specific induction media and confirmed after a 3-week culture using (A) Oil Red O staining and (B) the calcium-cobalt sulfide method, respectively. (C) Chondrocytic differentiation was identified using Alcian blue staining following a 3-week culture with chondrocytic medium, which were added to a pellet of 2.5x10^5 MSCs centrifuged at 150 x g for 5 min. hUC-MSCs, human umbilical cord-derived mesenchymal stem cells.

Figure 4. Expression of Notch-related genes in the hUC-MSCs and Jurkat cells. PCR was performed for the indicated transcripts from (A) hUC-MSCs and (B) Jurkat cells, respectively. Three serial dilutions of template cDNA are shown for each primer pair. hUC-MSCs, human umbilical cord-derived mesenchymal stem cells; Hes-1, hairy/enhancer-of-split 1; PCR, polymerase chain reaction.

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staining (Figs. 6 and 7), the Jurkat cells that were in contact with the hUC-MSCs underwent far less apoptosis induced by dexamethasone than those that were cultured alone. These data suggested that the hUC-MSCs were able to maintain the viability of the Jurkat T-ALL cells by preventing apoptosis.

**Discussion**

The interactions between hematological malignant cells and the elements of the stromal microenvironment play a key role in...
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The contemporary expression of the Notch1 receptor and addition to the constitutive expression of the Notch1 receptor. Jagged1 was also expressed by the Jurkat T-ALL cell line, in hUC-MSC-induced survival of the T-ALL cells. By contrast, the addition of recombinant Jagged1 protein enhanced the survival of the Jurkat cells that were treated with dexamethasone. The results of the blocking and stimulating experiments implied that Jagged1 contributed to hUC-MSC-induced drug resistance and to the self-maintenance of the Jurkat T-ALL cells.

In order to identify certain targets that are involved in the prevention of apoptosis mediated by hUC-MSCs, CD28 expression was assessed in the Jurkat cells in the present study. CD28 is one of the co-stimulatory molecules that are expressed by T cells (31). In the present study, the high expression level of CD28 in the Jurkat T-ALL cell line was more apparent following contact with the hUC-MSCs. CD28 has been identified as a direct target of Notch signaling (32), and has also been shown to be associated with the enhanced survival of immature (33) and activated (34) T cells. Therefore, the role of CD28 in the drug resistance of T-ALL warrants further investigation.

In conclusion, the present data indicate that the hUC-MSCs induced the drug resistance of the Jurkat T-ALL cell line. Jagged1, one of the Notch ligands, contributes to this phenomenon, which may also play a role in the self-maintenance of T-ALL cells and thus be a potential target for the treatment of human T-ALL. The evaluation of additional T-ALL cell lines, as well as primary T-ALL cells, using this co-culture system is necessary to expand these observations and to lay a theoretical basis for the development of new therapeutic strategies for T-ALL in the future.

Acknowledgements

The authors would like to thank Dr Danliang Chen, Department of Gynecology and Obstetrics, First Affiliated Hospital of Jinan University, for assisting with the umbilical cord sample collection. This study was funded by the National Natural Science Foundation of China (grant no. 31100664).

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