Immune oppression array elucidating immune escape and survival mechanisms in uveal melanoma

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Abstract

• AIM: To examine the genetic profile of primary uveal melanoma (UM) as compared to UM in immune escape.

• METHODS: Dendritic cells (DC) loaded with lysates of UM cells of high metastatic potential were used to stimulate cytotoxic T-lymphocytes (CTLs). When CTLs co-cultured with the UM cells, most UM cells could be eliminated. Survival UM cells grew slowly and were considered to be survival variants and examined by a microarray analysis. These differential genes were analyzed further with Venn Diagrams and functions related to immune escape. We additionally examined transcriptional changes of manually selected survival variants of UM cells and of clinical UM samples by quantitative real-time polymerase chain reaction (qRT-PCR), and analyzed the correlation of these expressions and patients’ survival.

• RESULTS: Gene expression analyses revealed a marked up-regulation of SLAMF7 and CCL22 and a significant down-regulation of KRT10, FXYD3 and ABCC2. The expression of these genes in the relapsed UM was significantly greater than those in primary UM. UM patients with overexpression of these genes had a shorter survival period as compared with those of their underexpression.

• CONCLUSION: Gene expression, in particular of SLAMF7, CCL22, KRT10, FXYD3 and ABCC2, differed between primary UM cells and survival variants of UM cells.

• KEYWORDS: uveal melanoma; dendritic cells; immune escape; gene expression profile; SLAMF7

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INTRODUCTION

Uveal melanoma (UM) is the most common malignant intraocular tumor in adults and the second most common type of melanoma [1]. The primary tumor can be treated by irradiation or by surgical resection, the survival of patients with UM is strongly linked to the development of distant metastases, which usually arise in the liver [2]. As a malignant intraocular tumor, UM cells may benefit from the immune privilege of intraocular compartment. It has been assumed that the UM cells when metastasizing keep elements of the ocular immune privilege as protection against an attack and destruction by lymphocytes [3]. In addition, UM cells can escape the immune destruction by inhibiting the immunostimulatory function of dendritic cells (DC) [4], by impairing the T-cell function of programmed death-1 (PD-1) and programmed death-ligand 1 (PD-L1) interaction [5], and by autocrine secretion of the Fas ligand (FasL or CD95L) [6]. The development and propagation of UM cells can thus take place despite the activation of cytotoxic T-lymphocytes (CTLs) as response of the host immune system. Although the eye is an immunoprivileged site, enlarging UM can show a dense lymphocytic infiltration [7]. Interestingly, the amount of the anti-tumor immune response characterized by a dense lymphocytic tumor infiltration including T-suppressor/CTLs [8], macrophages and regulatory T-cells has been correlated with the immune escape of the tumor cells [9-10].

To explain the high immunogenicity of UM [11], previous studies analyzed the secretion of cytokines and chemokines which mediated the inflammatory responses [7], or inhibited the immune cell functions [40-46]. These studies revealed that these cytokines, chemokines and infiltrating immune cells could influence the inflammatory tumor microenvironment that not only helped the tumor to suppress immune reactions but also aided the growth of new blood vessels for supporting the tumor growth [12]. Immune cells involved in this process were Foxp3+ regulatory T-cells [10], myeloid-derived...
suppressor cells\textsuperscript{(13)}, and CD68\/+CD163\/+ M2 macrophages\textsuperscript{(14)}. However, these studies did not address the phenomenon of the immune escape of UM cells that can lead to extraocular metastases. They also have not examined the changes occurred in the survival, variants of UM cells, such as alterations in gene expression or protein translation. Little was known about the gene expression profile of the immune escape variants of UM cells. As a consequence, molecular markers that would allow a selective targeting of UM cells and which could thus interrupt the metastatic cascade have not been developed yet. The development of tumor cell escape variants due to an immunoeediting may also explain the failure of CTLs to eliminate tumor cells completely\textsuperscript{(18)}. One of the other common mechanisms for the escape from CTL associated immunosurveillance is the lack of surface expression of HLA class I molecules on UM cells and subsequent loss of the non-classical major histocompatibility complex (MHC) molecules MIC-A/B expression during the progression of UM cells \textsuperscript{(16-17)}. UM cells lost HLA class I would be immunoselected under immune pressure.

**MATERIALS AND METHODS**

**Cell Culture** The UM cell line MUM-2B was obtained from the Beijing Ophthalmology and Visual Science Key Laboratory. MUM-2B is an epithelial cell type with a high metastatic potential. It was maintained in RPMI-1640 (Gibco, Life Technologies, Grand Island, NY, USA) with 10\% fetal calf serum (Gibco, Life Technologies, Carlsbad, CA, USA) at 37\°C in a 5\% CO\textsubscript{2} incubator.

**Preparation of Human Dendritic Cells** Human immature DCs were prepared from peripheral blood mononuclear cells which were obtained from a healthy donor \textsuperscript{(18)}. The DCs were prepared by a Ficoll/Paque density gradient centrifugation (Invitrogen, Life Technologies, Carlsbad, CA, USA). Peripheral blood mononuclear cells were seeded (9x10\textsuperscript{6} cells/3 mL/well) into 6-well plates (Corning Costar Corp., Cambridge, MA, USA) in RPMI-1640 medium. After 1.5h of incubation at 37\°C, adherent cells were used for the generation of DCs by adding granulocyte macrophage colony stimulating factor (1000 IU/mL, Peprotech, Rocky Hill, NJ, USA) and interleukin 4 (500 IU/mL, Peprotech). At least 3 to 6 million MUM-2B cells in suspension were smashed by an ultrasonic cell disrupter (Safer Co., Nanjing, China) to prepare cells as antigens \textsuperscript{(19)}. Tumor antigen protein concentrations were determined using Bio-Rad protein assay reagents (Bio-Rad Laboratories, Hercules, CA, USA)\textsuperscript{(19)}.

**T-cell Stimulation and Cytotoxic T-cell Assays Against High Metastatic Potential Uveal Melanoma Cell in Vitro**

The non-adherent lymphocytes were stimulated with autologous DCs loaded with MUM-2B lysates at a ratio of 5:1. The MUM-2B lysates and the DCs were co-cultured overnight before they were seeded with the lymphocytes. Interleukin-2 (300 IU/mL; Peprotech) was added to the cell cultures the next day and every 3d thereafter. The lymphocytes were re-stimulated with DCs every 5d for up to 2 stimulations, which consistently resulted in more than 90\% of CD3\/+ T-cells \textsuperscript{(20-22)}. Cytotoxic T-lymphocyte-mediated MUM-2B cytotoxicity was tested using the lactate dehydrogenase release assay (CytoTox 96 Non-Radioactive Cytotoxicity Assay, Promega Co., Madison, WI, USA) according to the manufacturer's protocol. Target cells were the cells of the high metastatic potential UM cell line MUM-2B. The ratio of effector and target cells was 2.5:1, 5:1, 10:1, 20:1 and 40:1.

**Flow Cytometry Analysis** For flow cytometry analysis, one million cells were stained with a solution containing 1\% phamingen stain buffer (BSA), 2 mmol ethylenediaminetetraacetic acid (EDTA), and flow antibodies for 30min at 4\°C.

We used the following antibodies: anti-CD11c (conjugated with fluorescein isothiocyanate), anti-CD83 (conjugated with phycoerythrin), HLA-DR (conjugated with peridinin-chlorophyll-protein), anti-CD86 (conjugated with phycoerythrin), anti-CD40 (conjugated with fluorescein isothiocyanate), and anti-CD80 (conjugated with phycoerythrin) (BioLegend, Inc., San Diego, CA, USA). The stimulated T-cells used anti-CD8 (conjugated with fluorescein isothiocyanate), anti-CD56 (conjugated with phycoerythrin), and CD3 (conjugated with peridinin-chlorophyll-protein), anti-CD25 (conjugated with fluorescein isothiocyanate), and anti-CD4 (conjugated with phycoerythrin) (BioLegend, Inc.). Mouse fluorescein isothiocyanate-IgG1, phycoerythrin-IgG1 and peridinin-chlorophyll-protein-IgG1 were used as isotype controls respectively. The flow cytometric analysis was carried out with a BD-FACSCalibur device (BD Biosciences, San Jose, CA, USA). The data were analyzed with the Cell Quest software (Becton Dickinson, Heidelberg, Germany). A minimum of 100 000 viable cells were measured per sample. Using forward and side scatter profile, debris and dead cells were gated out.

**Cell Proliferation Assay** In the cytotoxic experiment of killing UM cells, T cells were washed by used 1xPBS (pH=7.4) because T cells were suspension growth, so we collected these survival MUM-2B cells which were adherent growth. For un-interfusing T cells, we sorted CD3\/+ T cells and obtained single MUM cells by using MACS method \textsuperscript{(25-28)}. To perform a cell proliferation assay, both the survival and original MUM-2B cells were seeded in 96-well plates in a concentration of 1000 cells/well. The cell medium also included CTLs which were primed by DCs which were loaded by MUM-2B. The cell counting kit-8 (CCK8) assay (Biyuntian, Beijing, China) was used to determine the relative cell growth every 6, 12, 24 or 48h. Ten microliter of
CCK8 solution was added to the media for incubation for 2h at 37°C. The absorbance at 450 nm was measured by a Microplate Reader (Bio-Rad, Hercules, CA, USA).

**Preparation of RNA from Original Uveal Melanoma Cell and Its Survival Cells** Both the survival and original MUM-2B cells were collected for extracting total RNA that used for microarray analysis. Total RNA was extracted by using the Trizol reagent (Invitrogen, Carlsbad, CA, USA) and it was purified with an RNA Isolation Kit (Ambion, Austin, TX, USA). The purity and concentration of RNA were determined from OD 260/280 (optical density at 260 nm/optical density at 280 nm) readings using a spectrophotometer (Nanodrop ND-1000, Thermo Specific Co., Wilmington, DE, USA). The RNA integrity was determined by a 1% formaldehyde denaturing gel electrophoresis. Only RNA extracts with RNA integrity number values >6 were included in the in further analysis.

**Gene Expression Profiling** The gene expression profiling was performed using the Affymetrix Human Transcriptome Array 2.0 array (Affymetrix Co., Santa Clara, CA, USA). The human transcriptome array was designed with 10 per exon and 4 per junction, with approximately 4.8 million probes selected from all 98 million possible 25 mers. The array contained 7 million probes, so that the comprehensive transcriptome detected more than 0.28 million transcripts including more than 0.24 million coding transcripts and more than 0.04 million non-coding transcripts.

**Microarray Data Analysis** The Affymetrix HTA 2.0 arrays were hybridized according to Affymetrix recommendations using the Ambion WT protocol (Life Technologies, Carlsbad, CA, USA) and the Affymetrix labeling and hybridization kits. One hundred ng of total RNA were processed in parallel with an external microarray quality control (MAQC) A RNA to control the robustness of data. The labeled DNA mean yield was 7.19 μg (minimum: 6.27 μg; maximum: 7.57 μg). The Affymetrix GeneChip® Human Transcriptome 2.0 microarrays (HTA2) were hybridized with 4.7 μg of labeled DNA. The raw data, transcript data and the exon data were controlled with the Expression console (Affymetrix) at the Institute Curie microarray core facility (Santa Clara, CA, USA). The Affymetrix HTA2 dataset was analyzed by the Transcriptome Analysis Console (TAC) 2.0 (www. Affymetrix.com)23,24. We performed an unpaired Student’s t-test to compare gene intensities between the primary MUM-2B cells and the survival variants of MUM-2B cells. Genes were considered to be significantly differentially expressed when the fold-change was ≥2 and when the P-value was <0.05 (unadjusted P-value). The results were considered statistically significant for unadjusted P-values <0.05 and for fold-changes of ≥2 for the expressing analysis. After bioinformatic analysis of the microarray data, a manual inspection using the Molecular Annotation System was conducted to select differential genes, its gene ontology and its pathways. The data was logarithmically (log2) transformed and the median centered by genes using the adjust data function of the CLUSTER 3.0 software was then further analyzed with hierarchical clustering with average linkage.

For screening the candidate genes, these different expressed genes were analyzed further with Venn Diagrams and functions related to tumor metastatic and immune escaping25. **Target Genes Validation by Quantitative Real–time Polymerase Chain Reaction** We screened the candidate genes which were statistically significant for unadjusted P-values <0.05 and for fold-changes of ≥2 for the expressing analysis. *i.e. SLAMF7, CCL22, AQP9, KRT10, FXYD3 and ABC2C_, and validated further by quantitative real-time polymerase chain reaction (qRT-PCR). A cDNA synthesis was carried out using a PrimeScript II 1st strand cDNA Synthesis Kit (TaKaRa, Dalian, China). A qRT-PCR was performed using the ABI 7300 sequence detection system (Applied Biosystems, Foster City, CA, USA) in a volume of 20 μL containing 10 μL GoTaq® qPCR Master Mix (Promega, Madison, WI, USA) and 0.2 μm of each primer of the tested target genes. SLAMF7 fw: 5’-ATGTTCTA CGAGCACCTGTCAAACGC-3’, rev: 5’-GGGCTTTCCAG GTATAAATCACATC-3’, CCL22 fw: 5’-TCTGGGTTTCTT GCCTGGGATG-3’, rev: 5’-CCAAGAATCTGCAGAGACT GTGA-3’, AQP9 fw: 5’-AGAGACGGCTAGCGAAAGAAACCCC-3’, rev: 5’-GATAGTGATGACCCCTCCCAAAAACGG-3’, KRT10 fw: 5’-CGTGGACAGAACAGAGTGTCGCT-3’, rev: 5’-GCAGGCTTGGTGATGTTTGAA-3’, FXYD3 fw: 5’-AG ATGTCCGCTGCCGAAGATTTCCC-3’, rev: 5’-TTCCATT TGGGGGTTTGGTC-3’, ABC2C fw: 5’-GCTGGGAAAGTCA TCCCTCACAAC-3’, rev: 5’-GGTAGGACCAATGGAAAG CAATA-3’, Actin fw: 5’-ACTTAGTGATGCACCTTACCC-3’, rev: 5’-GTCACCTTACGTTCAC-3’. The reaction profile consisted of 40 cycles at 95°C for 2min, at 55°C for 30s, and at 72°C for 30s. A dissociation stage was performed at the end of the reaction consisting of 200 cycles of 7s with the temperature increased at 0.2°C cycle to demonstrate the specificity of the amplification. The expression analysis was performed in triplicate for each sample. The housekeeping actin was used as the normalization control. The fold difference for each sample was obtained using the equation 2^-dCt, where Ct is the threshold cycle (the cycle number at which the fluorescence generated within a reaction crosses the threshold) and dCt equals the mean Ct of the sample gene minus the mean Ct of actin.

**Clinical Tumor Samples and Controls** Fresh tumor samples were obtained from 10 UM patients at the time of eye removal or tumor resection. There were 3 patients ("relapsing UMs") who had undergone laser therapy and radiotherapy of the UMs prior to enucleation of the eyes.
Figure 1 Flow cytometric results of DCs and their activated CTLs A: DCs were cultured by inducing peripheral mononuclear blood cells which were derived from healthy donors. DCs were loaded with lysates of MUM-2B and matured. They then highly expressed CD11c+/HLA-DR+, and the co-stimulatory molecules CD40+, CD80+, CD83+ and CD86+. B: Before DCs stimulated T-lymphocytes which were derived from the same healthy donor. The phenotypes of T-lymphocytes had CD3+, CD8+, and CD3+/CD56+. C: DCs loaded with lysates of MUM-2B cells, stimulated proliferation and activation of T-lymphocytes which were derived from the same healthy donor. The resulting CTLs highly expressed CD3+, CD8+, and CD3+/CD56+, while in contrast CD4+ was expressed to a lower degree.

the remaining 7 patients, the tumor had not been treated prior to the enucleation. The tissue samples were immediately frozen in liquid nitrogen and stored at -80°C until processing. The diagnosis of UM was confirmed by a surgical pathologist for all patients. The UM samples were analyzed for target genes and protein expression by performing qRT-PCR and Western blotting.

Statistical Analysis The statistical analysis was performed using a commercially available statistical analysis program (SPSS 21.0, IBM-SPSS; Chicago, IL, USA). Data were presented as mean ± standard deviation from three separate experiments. Statistical significances of differences of means throughout this study were calculated by Student's t-test in comparing data between two groups and ANOVA one-way test in comparing data from more than two groups. Correlations between the expressions of five genes were evaluated by bivariable analyses using log rank test of Kaplan-Meier methodology. Differences between groups were considered to be statistically significant with a P-value of <0.05.

Ethics Statement All experiments and procedures were performed in accordance with the appropriate guidelines. All procedures were approved by Capital Medical University. Informed consent was obtained from all volunteers before being enrolled in the study.

RESULTS Production of Dendritic Cells Loaded with Uveal Melanoma Cell Line (MUM –2B) Elicits Specific Cytotoxic T-lymphocytes in Vitro To determine the effect of the vaccination of the DCs, DCs were isolated from peripheral mononuclear blood cells of a healthy donor. They were primed in vitro using autologous DCs pulsed with MUM-2B lysates. The DCs then highly expressed the DC markers of CD11c and HLA-DR and the co-stimulatory molecules of CD80, CD86, CD83 and CD40 (Figure 1A). The lymphocytes were re-stimulated with DCs every 5d. The stimulated T cells were then tested for cells phenotypes by flow cytometry [26]. Before DCs stimulated T-lymphocytes which were derived from the same healthy donor. The phenotypes of T-lymphocytes were low level correspondingly (Figure 1B). But the stimulated T cells contained CD3+ in 94.9% of the cells, CD8+ in 79.6% of the cells, and CD3+/CD56+ in 37.5% of the natural killer T (NKT) cells, while only 28.6% of the cells expressed CD4+, and 9% of the
cells were regulatory T-cells (CD4+/CD25+) (Figure 1C). It showed that DCs loaded with MUM-2B lysates stimulated the proliferation and activation of CD8+ T-lymphocytes. The stimulated cells were then tested for their cytotoxic ability against high metastatic potential UM cell (MUM-2B) by a cytotoxicity assay of the CTLs. We included cytotoxic T-lymphocytes which were stimulated by DCs pulsed with MUM-2B lysates (UM-DC-CTLs), CTLs stimulated by DCs without antigens (NON-DC-CTLs), and non-stimulated T-lymphocytes. With the increase of Effector/Target (E/T) ratio, the cytolytic against MUM-2B cells increased \(^{[23-24]}\). It showed that the cytotoxicity of UM-DC-CTLs was significantly higher than the cytotoxicity of NON-DC-CTLs and the cytotoxicity of T-cells (Figure 2A). Moreover the cytotoxicity of NON-DC- CTLs did not differ significantly from the cytotoxicity of T-cells against the survival UM cells.

**Effect of Uveal Melanoma Cell Proliferation by Immune Function of Dendritic Cells Activating Cytotoxic T-lymphocytes**

To test the proliferation of the UM cells under immune suppression by the DCs activated CTLs, we examined the MUM-2B viability at different time points \(^{[23-24]}\). We found that the growth of the MUM-2B cells decreased gradually. Viability was lowest at two days after co-culture with the CTLs, and then slightly increased for the next few days (Figure 2B, 2C). The proliferation of the primary MUM-2B cells differed significantly \((P<0.01)\) from the proliferation of the MUM-2B cells under immune suppression (Figure 2B, 2C). The MUM-2B cells still survived at two days at the greatest E/T ratio, and their growth recovered by 67% at 8d later (Figure 2C).

**Gene Expression Signature Discriminates Between Original Uveal Melanoma Cells and Its Survival Cells**

In the gene expression analysis, samples were clustered into two groups, \(\leq\) the group of survival variants of MUM-2B cells and the group of the original MUM-2B cells as normal controls. The survival variants of MUM-2B cells were obtained from surviving UM cells at a E/T ratio of 5/1 (group C1), a E/T ratio of 10/1 (group C2), and a E/T ratio of 20/1 (group C3). The mean expression of each gene was compared between the three survival variants of MUM-2B cell groups and the control group. With a \(P\)-value of \(<0.01\) used as a cut-off, 824 genes were overexpressed and 69 genes were underexpressed with a \(\geq2.0\) fold-change in merge of the C1/N, C2/N and C3/N group \((P<0.01)\) (Figure 3).

**Gene Function Analysis and Differentially Expressed Immune Response Genes in Survival Uveal Melanoma Cells**

The analysis of gene functions of differentially expressed immune response genes in the survival variants of
UM cells versus the original UM cells revealed genes in the field of antigen processing and antigen presentation (109 genes, \(P=0.00\)), response to stress (226 genes, \(P=0.00\)), regulation of a defense response (70 genes, \(P=0.00\)), chemotaxis (27 genes, \(P=3.81 \times 10^{-4}\)), regulation of inflammatory response (40 genes, \(P=0.00\)), T-cell activation (29 genes, \(P=1.97 \times 10^{-14}\)), regulation of regulatory T cell differentiation (6 genes, \(P=1.33 \times 10^{-6}\)), regulation of interleukin-6 production (22 genes, \(P=1.97 \times 10^{-4}\)), regulation of interleukin-8 production (11 genes, \(P=3.25 \times 10^{-7}\)), regulation of interleukin-10 production (5 genes, \(P=2.33 \times 10^{-3}\)), and positive regulation of transforming growth factor beta production (2 genes, \(P=0.04\)).

Differential genes related to antigen processing and presentation included 109 genes which were differentially expressed (\(P<0.05\)) in merge of C1/N, C2/N and C3/N group (Table 1). These genes were up-regulated in all survival variants of MUM-2B cells. There were three genes involved in positive regulation (\(CD74\)) and negative regulation (\(CCR7, CD47\)) of DC antigen processing and presentation pathways. There were 226 differentially expressed genes which were associated with response to stress (\(P<0.05\)) (Table 1). Most of these genes are involved in the defense response, immune response and cell communication. The genes \(AQP9\) and \(SLAMF7\) were the

Table 1 The summary of significant overexpression and underexpression genes which were differently expressed in MUM-2B suppressed by CTLs activated by DCs

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genes with the most marked increase. Down-regulation of KRT10 and ABCC2 related to drug-resistance of cancer medication.

Differential genes related to inflammation (cytokines and chemokines) were genes encoding macrophage- and T-lymphocyte-attractant chemokines and cytokine receptors (CCL2, CCL5, CCR1, CCR4, CCL18, CCL22, IL6 etc.), molecules related to cytokine-signaling pathways (TNFRSF18, TNFAP6, PIK3CG, JAK2, PIK3AP1), proteins implicated in the extravasation of leukocytes from blood to tissues (CSF1, ITGAI, ICAMI, P2RX7) and molecules that contribute to inflammation (LYZ, HCK, CXCL10, and Down-regulation of FXYD3) (Table 1).

Differential genes related to T-lymphocytes included those encoding for molecules that participate in the signal transduction through the T-cell receptor TCR (LCPI, LC2, XCL1, and PTTPN22), proteins involved in T-cell activation and proliferation (IL7R, CD84 and P2RX7), natural killer cell mediated cytotoxicity markers (SLAMF7, PTTPN6, TGFBI1 and IDO1), and T-cell mediated cytotoxicity markers (PTPRC, ITGAL, ICAMI, GNYL, CCL2) (Table 1). Genes expressed in activated immune cells that were negatively regulated by the TGFB-β pathway (XCL1 and A1P6AP2), the interleukin-10 pathway (XCL1, PLA2G7, IDO1, CD48 and CD40LG), the interleukin-8 pathway (BTN3A1, CD40LG, TLR8, PPA2B, CD80 and DDO11), the interleukin-6 pathway (PLA2G7, TAPI, BTN3A1, TNFAP3, HLA-B, and CD40LG) and regulatory T-cell pathway (CTLA4, TGFBI, CD28, HLA-G, and FOXP3). These negative regulatory pathways participated in mediating tumor survival.

Validation of the Differentially Expressed Genes by Quantitative Real-time Polymerase Chain Reaction

Based on the gene ontology and pathways analysis of differential genes, we selected six candidate target genes by screening further of Venn Diagrams and gene functions. These candidate target genes were three up-regulated genes (SLAMF7, CCL22 and AQP9) and three down-regulated genes (KRT10, FXYD3 and ABCC2) (Table 1). Table 2 showed that SLAMF7, CCL22 and AQP9 were still highly expressed in survival variants of MUM-2B at E/T ratios of 5/1 (CTL5), 10/1 (CTL10) and 20/1 (CTL20). KRT10, FXYD3 and ABCC2 were expressed to a lower degree in the survival variants of MUM-2B cells, but only the expression of KRT10 showed a significant difference to the expression in the primary MUM-2B UM cells.

Differential Genes Validation in Clinical Uveal Melanoma Samples by Quantitative Real-time Polymerase Chain Reaction

For the validation of the differential genes we collected UM cells from 10 patients with UMs samples (Table 3). We found that the expression of SLAMF7 was significantly higher in the relapsing UM cells than in the primary UM cells (Table 4; Figure 4A). The expression of CCL22 was higher in the relapsing UM cells than in the primary UM cells and in the UM tumor tissue (Table 4; Figure 4B). The expression of AQP9 did not differ significantly between the groups. The expression of KRT10 was significantly (P<0.01) lower in primary tumors than in relapsing tumors. In addition, the expression of KRT10 differed between the various subtypes of primary UM (Table 3; Figure 4C). The expression of FXYD3 was significantly (P<0.001) higher in the relapsing UM cells than in the primary UM cells (Table 4; Figure 4E). ABCC2 was expressed significantly higher in the relapsing UM cells than in primary UM (Table 4; Figure 4F).

Differential Genes Expression Correlated with Patients’ Survival

Twenty UM patients were collected and their samples were detected expression of SLAMF7, CCL22, KRT10, FXYD3 and ABCC2 by qRT-PCR. The cut-off value of these genes was 2^ΔΔCt=0.23 of SLAMF7, 2^ΔΔCt=0.69 of CCL22, 2^ΔΔCt=0.45 of KRT10, 2^ΔΔCt=4.53 of FXYD3 and
Figure 4 The fold difference of target genes obtained in UM tissues

A: SLAMF7 was up-expressed in relapsing tissues and significantly higher than in primary UM tissues; B: CCL22 was expressed highly in relapsing tissues and significantly higher than in primary UM tissues; C: The expression of AQP9 in mixed type UM tissues was highest in these tissues. However, the expression of AQP9 hadn't significantly difference among all tissues; D: The expression of KRT10 in relapsing tissues was highest among UM tissues. KRT10 also was expressed highly in relapsing tissues and significantly higher than in primary UM tissues; E: The expression of FXYD3 was the highest in relapsing tissues. The expression of FXYD3 in relapsing UM tissues had evidently difference with in primary UM tissues. Additionally, the expression of FXYD3 in relapsing UM tissues has relevance with mixed type UM; F: ABCC2 also was up-expressed in relapsing tissues and significantly higher than in primary UM tissues.

Table 4 The fold difference of target gene expression between UM tumor tissue and normal choroidal tissues

<table>
<thead>
<tr>
<th>Genes</th>
<th>Spindle-cell type (n=4)</th>
<th>Mixed cell type (n=5)</th>
<th>Epitheloid cell type (n=4)</th>
<th>Tumor relapse</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Yes (n=7)</td>
</tr>
<tr>
<td>SLAMF7</td>
<td>0.08±0.10</td>
<td>0.13±0.10</td>
<td>0.08±0.00</td>
<td>0.77; 0.50</td>
</tr>
<tr>
<td>CCL22</td>
<td>0.38±0.31</td>
<td>0.13±0.09</td>
<td>0.22±0.29</td>
<td>0.16; 0.38</td>
</tr>
<tr>
<td>AQP9</td>
<td>3.77±2.28</td>
<td>16.01±24.87</td>
<td>1.40±1.52</td>
<td>0.73; 0.37</td>
</tr>
<tr>
<td>KRT10</td>
<td>0.12±0.02</td>
<td>0.03±0.03</td>
<td>0.19±0.26</td>
<td>0.01; 0.74</td>
</tr>
<tr>
<td>FXYD3</td>
<td>0.10±0.33</td>
<td>1.45±1.12</td>
<td>2.22±2.31</td>
<td>0.71; 0.57</td>
</tr>
<tr>
<td>ABCC2</td>
<td>0.02±0.02</td>
<td>0.02±0.02</td>
<td>0.03±0.04</td>
<td>0.99; 0.80</td>
</tr>
</tbody>
</table>

1P: Mixed cell type vs Spindle-cell type; 2P: Epitheloid cell type vs Spindle-cell type.
2.0) = 0.07 of ABCC2, so distinguish overexpression from underexpression. There were these genes overexpression of 7 UM patients and these genes underexpression of other 13 UM patients. The survival time showed significant difference between both patients by Kaplan-Meier survival curves (log-rank test, \( P = 0.000 \) ) (Figure 5). All patients of these genes overexpression occurred UM relapse and liver metastasis. The result elucidated that these genes could be used as targets to locate diagnostic, prognostic and therapeutic biomarkers for UM.

**DISCUSSION**

In our experimental study, we cultured mature DCs derived from peripheral blood mononuclear cells of healthy donors. DCs were pulsed with lysates of UM cells of high metastatic potential (MUM-2B cell line) and were used to stimulate CTLs. The mature DCs highly expressed co-stimulatory molecules, and the activated CTLs highly expressed more than 90% CD3+, more than CD70% CD8+, and more than 30% CD3+/CD56+NKT-cells (Figure 1A, 1B). When mixed with the primary UM cells, the CTLs killed the majority of the MUM-2B cells within two days (Figure 2A, 2B), while about 10% of the UM cells survived and slowly proliferated under the CTLs associated immunization for 10d (Figure 2C). One may assume that the surviving UM cells escaped the immune response activated by the DCs. We then used microarray samples of these survival variants of UM cells and compared them with microarray samples of the original MUM-2B cells and analyzed differences in expressed genes and their molecular function and pathways which were closely related to the survival (Figure 3).

The survival variants of UM cells differentially expressed 109 genes associated with the antigen processing and presentation pathways which were involved in the positive regulation and negative regulation of DCs (Table 1).

Important molecules were CD47 and CCR7 for the negative regulation, and MHC class II molecules that induced antigen-specific CD4+ T-cell tolerance. CD47 is generally considered as an anti-phagocytic signal expressed by cancer cells to prevent macrophages and DCs from attacking these tumor cells. We found 226 genes which were differentially expressed and which were associated with the response to stress involving defense response, immune response, and cell communication (Table 1). Interestingly, we detected that AQP9 and SLAMF7 were up-regulated in the survival variants of UM cells as compared with the corresponding primary UM cells. The unique property of AQP9 is associated with an increased metabolic demand, such as spontaneous labor and inflammation, which plays a pivotal role in various arsenide-mediated cytocidal effects on cancer cells. SLAMF7 inhibits natural killer cell activation and cytotoxicity by limiting antibody-dependent cell-mediated cytotoxicity.

Additionally, the down-expression level of KRT10 and ABCC2 was the first and second gene that was involved in drug-resistance of cancer. PTEN improves cisplatin-resistance of human ovarian cancer cells through up-regulating KRT10 expression. ABCC2 is associated with lower differentiation of cancer cells and their resistance to the cisplatin, moreover its overexpression is responsible for the drug-resistance of cancer stem cells.

The gene expression profiles revealed that the survival mechanism of UM cells was interrelated with signal transduction, such as T-cell activation and proliferation, and natural killer cell mediated cytotoxicity markers, and T-cell mediated cytotoxicity markers (Table 2). Negatively regulated pathways in activated T-cells involved in the TGF-\( \beta \) pathway, interleukin-10 pathway, tinterleukin-8 pathway, interleukin-6 pathway and regulatory T-cell pathway among others. In particular natural killer cells are well-developed and can contribute to the immune surveillance of tumors, thus SLAMF7, PTPN6, TGFBI, and IDO1 inhibit the natural killer cell proliferation and the natural killer cell mediated cytolsis.

There were 29 differential cytokines and chemokines which were found to join the immune inflammation (Table 1). The presence of CTLs in UMs may be a necessary but not sufficient condition for tumor rejection. In fact, the dense infiltration of lymphocytes in UMs persists after immunoedition, probably recruited or locally expanded by the elevated expression of these specific factors in the UM microenvironment. High-expression of CCL22 and down-expression of FXYD3 in our study was of particular interest, because CCL22 belongs to macrophage-derived chemokines and regulatory T-cell cell attracting chemokines, which generate a microenvironment that may favor survival and
growth of malignant cells, and overexpression of FXYD-3 may play an important role in the tumorigenesis and development by transforming growth factor-β signaling through ZEB1/δEF1 in human cancers[43-44].

Based on the gene ontology and pathways analysis of differential genes as described above, we selected three up-regulated differential genes (SLAMF7, CCL22 and AQP9) and three down-regulated differential genes (KRT10, FXYD3 and ABC2) which were validated in the survival variants of MUM-2B cells and in the original MUM-2B cells again (Table 3). Six differential genes were detected in the UM tumor tissue samples of 10 patients (Table 4; Figure 4). In addition, the expression of both SLAMF7 and ABC2 were significantly higher in the relapsing UM tissues than in the primary UM cells. Similarly, the expression of both CCL22 and KRT10 were higher in relapsing UM cells than that in primary UM cells (Table 4; Figure 4B, 4D). The expression of FXYD3 also was higher in relapsing tissues than in the primary UM tissues (Table 4; Figure 4E). In contrast, the expression of AQP9 did not differ significantly between the groups. We found that KRT10, FXYD3 and ABC2 were down-regulated under immune suppression by DC stimulated CTLs, but they were up-regulated in survival variants of UM cells. Three genes are responsible for drug-resistance of cancer cells [34-35,45-47], so that they can be up-regulated in the relapsing UM tissues, although the mechanisms have remained unclear yet. We collected relapsing UM samples of these patients who had undergone radiotherapy and laser therapy, so the expression of three genes mediated radiotherapy-resistance of UM [34-35,45-47].

Five candidate differential genes, SLAMF7, CCL22, KRT10, FXYD3 and ABC2 were possibly associated with the survival UM cells. These results may suggest that SLAMF7, CCL22, KRT10, FXYD3 and ABC2 may participate in the survival variants of mechanisms of UM. Future studies may examine their molecular functions at the cell level and in vivo to uncover their molecular mechanism in immune escaping. We found that these genes overexpression impacted the survival time of UM patients severely, moreover these patients produced UM relapse and liver metastasis. Potential limitations of our study should be mentioned. First, as for any cell culture study, the question arises how much the results can be transferred into the clinical situation. Second, the relapsing UM were pre-treated by radiotherapy or laser therapy, so that it was not possible to clearly distinguish between direct effects of these treatments or effects caused by the survival variants of tumor cells. Third, a main point about UM is that CTLs are most likely not the most important contributors to the killing of the tumor cells in the eye, due to the intraocular immune privileged conditions. They may, however, be relevant in the liver for the metastases [40]. Another factor of importance is the killing of the tumor cells by natural killer cells. Natural killer cells especially focus on cells that have lost HLA Class I antigens, and if cells become less lysable by CTLs, they may become more lysable by natural killer cells [40].

In conclusion, gene expression differed between primary UM cells and survival variants of UM cells. Since the present study did not provide data indicating whether the differentially expressed genes in the MUM-2B cells that survived the culture with CD8+ T cells conferred protection from CD8+ T cells, future studies could include cytotoxicity assays with MUM-2B cells that survived a culture with CD8+ T cells, MUM-2B cells that were engineered to overexpress the genes of interest, and surviving MUM-2B cells in which these genes of interest were silenced. Future studies may also address whether the differences in the gene expression found in our study may be helpful to early detect survival variants of UM cells and may be helpful to develop targeted treatment strategies which may be more specifically focused on the survival variants of UM cells in contrast to the primary UM cells. Addressing the survival pathways which were associated with the survival of survival variants of UM cells under immunization in our study could provide targets to disrupt the survival cascade in UM.

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REFERENCES


Immune escape and survival mechanisms in uveal melanoma


