Research Article

IL-6 Promotes the Proliferation and Immunosuppressive Function of Myeloid-Derived Suppressor Cells via the MAPK Signaling Pathway in Bladder Cancer

Zhong Zheng,1 Xinyi Zheng,2 Yiwen Zhu,3 Zhixian Yao,1 Weiguang Zhao,1 Youjia Zhu,4 Feng Sun,1 Xingyu Mu,1 Yong Wang,5 Wanqing He,6 Zhihong Liu,1 Ke Wu,1 and Junhua Zheng1

1Department of Urology, Shanghai General Hospital, School of Medicine, Shanghai Jiao Tong University, Shanghai, China
2Department of Pharmacy, Huashan Hospital, Fudan University, Shanghai, China
3School of Rehabilitation Science, Shanghai University of Traditional Chinese Medicine, Shanghai, China
4The Second School of Medicine, Wenzhou Medical University, Wenzhou, China
5Department of Urology, Shanghai Jiangqiao Hospital, Jiading Branch, Shanghai General Hospital, Shanghai, China
6Student Innovation Center, Shanghai Jiao Tong University, Shanghai, China

Correspondence should be addressed to Zhihong Liu; drzhihongliu@sjtu.edu.cn, Ke Wu; doctorwuke@sjtu.edu.cn, and Junhua Zheng; zjhzztg@163.com

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Muscle-invasive bladder cancer (MIBC) is characterized by a highly complex immune environment, which is not well understood. Interleukin-6 (IL-6) is generated and secreted by multi-facets of cells, including tumor cells. This study was aimed at demonstrating that the levels of IL-6 and the number of myeloid-derived suppressor cells (MDSCs), with a positive correlation between them, increased in MIBC tissues, promoting MIBC cell proliferation, especially in patients with recurrence. In coculture analysis, MDSCs, with the stimulation of IL-6, could significantly lower the proliferation ability of CD4+ or CD8+ T lymphocytes. Further, this study demonstrated that IL-6 could upregulate the mitogen-activated protein kinase (MAPK) signaling pathway in MDSCs. The MAPK signaling inhibitor, aloesin, partially reversed the effects of IL-6 on MDSCs. These data suggested that IL-6 promoted MIBC progression by not only accelerating proliferation but also improving the immune suppression ability of MDSCs through activating the MAPK signaling pathway.

1. Introduction

Bladder cancer (BC) has the 14th and 9th highest rates of mortality and incidence among all types of cancers [1], accounting for approximately 900,000 newly diagnosed cases each year, with 250,000 deaths occurring during the same time span [2]. Nearly 75% of patients are diagnosed with non-muscle-invasive BC (NMIBC) and close to 25% with muscle-invasive BC (MIBC) [3]. Similar to other solid tumors, MIBC is classified as immunogenic cancer, with numerous tumor-infiltrating lymphocytes (TILs) in the tumor microenvironment (TME) [4]. Besides tumor cells, the TME includes fibroblasts, endothelial cells, mast cells, macrophagocytes (Mφ), TILs, and myeloid-derived suppressor cells (MDSCs) [5]. In particular, TILs contribute to the adaptive immunological response in opposition to cancer cells and exert antitumor effects in most malignant tumors, including BC, colorectal cancer,
lung cancer, and melanoma [6]. Nevertheless, the dysregulation of immunosuppressive TME is intensively associated with the development from NMIBC to MIBC [7]. It is, thus, possible that the immunosuppressive functions of TILs in MIBC tumors are prompted, explaining the poor outcomes in MIBC. However, the mechanisms underlying the peculiar pathophysiologic behavior of TILs in MIBC are unclear.

Heterogeneous MDSCs, a cluster of immature myeloid-derived cells, are both immunosuppressive and angiogenic. They result from and suppress inflammatory conditions that support tumor growth [8] by stimulating the accumulation of regulatory T lymphocytes and suppressing responses initiated by cytotoxic T cells [9]. In patients bearing solid tumors, including MIBC, the interleukin 6 (IL-6) levels are often elevated and inversely associated with tumor growth [8] by stimulating the accumulation of a plethora of relevant genes that modulate the immune status, inducing the infiltration of MDSCs. Hence, the mechanisms underlying IL-6-mediated MDSC penetration and functionality should be further explored [12].

Mitogen-activated protein kinases (MAPKs) are serine-threonine protein kinases that modulate a large number of cellular processes, such as innate immunity, survival, apoptosis, differentiation, proliferation, and inflammation [13]. In the wake of tissue damage or pathogen infection, the initiation of pattern recognition receptors in the cell plasma and on the cellular surface of innate immune cells activates a majority of MAPK subfamilies, that is, p38 MAPK, extracellular signal-regulated kinase (ERK), and Jun N-terminal kinase subfamilies. In combination with the upregulation of interferon-regulatory factor as well as nuclear factor-κB, MAPK activation leads to the upregulation of a plethora of relevant genes that modulate the immune responses together [13]. The ERK signaling pathway mediates the transition of MDSCs. Its downregulation is correlated with MDSC differentiation. The dysregulation of MAPK signaling pathways promotes the pathogenesis of malignant tumors [14].

This study was aimed at exploring the mechanisms by which IL-6 promotes the immunosuppressive functions and proliferation of MDSCs via the MAPK signaling pathway in MIBC.

2. Materials and Methods

2.1. Patient Materials. Malignant lesions and adjacent normal controls were obtained from 200 patients (aged 53–88 years) diagnosed with MIBC who underwent radical cystectomy or transurethral resection of the bladder tumor (TURBT). Specimens were obtained from Shanghai General Hospital, Shanghai, China. The removal of tissues and subsequent analyses were approved by the ethics committee of the hospital and performed after obtaining informed consent from the patients’ relatives.

2.2. In Vivo Experiments. C57BL/6 mice (Shanghai Model Organisms Center, China), aged 6–8 weeks, were subcutaneously (s.c.) injected with MB49 cells (2 × 10⁶) and raised under pathogen-free conditions in the Shanghai General Hospital Animal Center. In the IL-6 treatment group, the mice were randomly divided into the experimental group (n = 20) and control group (n = 20). The grouping conditions in the MDSC transfer group were identical with those in the IL-6 treatment group, but it was launched after the sacrifice of mice on the 30th day in the IL-6 control group owing to the requirement of MDSCs. The BC tissues of the mice in the IL-6 control group after relevant analysis were subsequently used for MDSC separation and transfer. All the in vivo experiments were launched 10 days (the average number of days for the visible tumor node) after tumor implantation. In the IL-6 treatment group, the mice were intraperitoneally injected with phosphate-buffered saline (PBS) or recombinant murine IL-6 (aa 20-357, 20 ng/g body weight; carrier-free, R&D Systems) at 10 a.m. every day. In addition, MDSCs from BC tissues (5 × 10⁶ cells/mouse) mentioned earlier were adoptively transferred to BC mice via tail vein injection every 7 days, and PBS as a vehicle was administrated to the control group. The tumor growth rate was calculated by measuring the estimated tumor volume with a digital caliper every 5 days. The estimated tumor volume was determined using the canonical formula: volume (cm³) = 1/2 × (width)² × length. After 30 days, 10 mice randomly extracted from each group were euthanized. The tumor tissues were collected for flow cytometry and MDSC separation for further experiments, including coculture analysis, quantitative reverse transcription polymerase chain reaction (qRT-PCR), and RNA sequencing, under specific conditions. The remaining 10 mice per group were kept for survival analysis. The survival curves for all groups were established using the Kaplan–Meier criteria. The mice were considered expired when the tumor volume reached 2.5 cm³ posttreatment in concordance with the regulations released by the Institutional Animal Care and Use Committee. The animal study was scrutinized and approved by the Institutional Review Board of Shanghai Jiao Tong University.

2.3. Immunohistochemistry. The slices from the formalin-fixed paraffin-embedded blocks of MIBC specimens were obtained, and an anti-IL-6 antibody (ab233706, Abcam) was applied for immunostaining to investigate the expression of IL-6. Slices (4 μm) of MIBC were loaded on poly-L-lysine-coated slides with air drying and deparaffinization. Subsequently, peroxidases were blocked with 5% hydrogen peroxide in 50% methyl alcohol at 20°C for 20 min. Then, antigen retrieval was performed using a microwave oven for 20 min in citrate buffer solution. The slices were treated with a monoclonal rabbit anti-IL-6 antibody for 12h at 4°C after blocking nonspecific binding. They were washed with PBS and treated with biotinylated goat anti-rabbit IgG (ab150077, Abcam) for 30 min, followed by rinsing with PBS. The immune reactivity was examined
using the avidin-biotin-peroxidase complex method utilizing VECTASTAIN ABC Kit (PK-4001, Vector). Then, the slices were stained with hematoxylin, dehydrated, cleared, and mounted. The tissues manifesting brown staining in the membrane, nucleus, or cytoplasm were considered positive. The average optical density (AOD) was analyzed using ImageJ software (version 2.0.0, Rawak Software, Inc., Germany) to investigate the correlation between IL-6 levels and MDSCs.

2.4. IL-6 Enzyme-Linked Immunosorbent Assay. The IL-6 enzyme-linked immunosorbent assay (ELISA) was applied according to the product specification (ab178013 for humans and ab100713 for mice, Abcam). Duplicate aliquots of the supernatant were dispersed in the flat-bottom 96-well plates. The assay was implemented triply on independent samples.

2.5. Flow Cytometry and MDSC Isolation. Fresh tumor and adjacent tissues were used to prepare single-cell suspensions. The antibodies used for flow cytometry were provided by BioLegend and are listed in Table S1. The cells were examined using a FACSCalibur Flow Cytometer (Becton Dickinson). The cells were stained with carboxyfluorescein succinimidyl ester (CFSE) and analyzed using a flow cytometer (Thermo Fisher) following the manufacturer’s protocol. The cells were fixed, permeabilized, and stained with anti-IL-6-FITC (501104, BioLegend) for 30 min to detect the expression of IL-6. Data were analyzed using FlowJo software (version 10.7.1, Treestar). For MDSC flow cytometric sorting, 1 × 10^7 cells/mL single-cell suspensions made from malignant tissues derived from tumor-bearing mice were applied with conjugated antibodies shown in Table S1 for 20 min on ice in staining buffer (1% FBS in PBS). Then, the cells were rinsed with PBS, and the samples were sorted using a BD Influx. The 1.5-drop pure sort mode was selected to obtain cells with high purity. Subsequently, isolated MDSCs were maintained for experiments as follows. The cell culture passage number was no more than 10 for all isolated cells, and three replicates were performed in each passage.

2.6. RT-PCR Analyses of IL6, Arg1, and iNOS. Total RNA was obtained using the RNeasy Mini Kit (27104, Qiagen). Reverse transcription was performed using the TaKaRa RNA PCR kit (RR012A, TaKaRa) following the manufacturer’s protocols. The primers were as follows: IL6 (forward 5' - GCTTCC CTCAGGATGCTTGTT-3', reverse 5' - ATTAACGTTGGG TGCTTGCT-3'), Arg1 (forward 5' - AGTTAGAGGCC ACAAAGC-3', reverse 5' - TGTACGCTCCACCAGCAT TTCC-3'), and iNOS (forward 5' - CTGAAGATACCGGAC GGCTGCG -3', reverse 5' - CGATGAAGTCCGGGTTGCTAG -3') for humans and IL6 (forward 5' - GCCCTCTAGTGTTT CTGTGT-3', reverse 5' - ACTGCAAGGCCCAGTTTACATCC -3'), Arg1 (forward 5' - GCAATGCGAGTGGAACTGC -3'), and iNOS (forward 5' - CTGAAGATACGGCAGGCTGCG -3', reverse 5' - GATCGAGGTGGGTTGCTAG-3') for mice. The SYBR Green PCR Master Mix (4309155, Thermo Fisher) and a real-time PCR machine (Applied Biosystems) were used in the reactions. All experiments were performed in triplicate.

2.7. Immune Suppression. Lymphocytes from plasma were isolated by the density-gradient centrifugation using Ficoll-Hypaque solution (Solarbio). CD8+ microbeads (130-045-201, Miltenyi Biotec) or CD4+ microbeads (130-045-101, Miltenyi Biotec) were used for T cell sorting. The freshly isolated CD8+ T cells or CD4+ T cells were subsequently cultured in 96-well round-bottomed plates under the conditions that a complete culture medium contained 500 ng/mL soluble anti-CD28 (T9-577-T100, American Research Products) and 500 ng/mL soluble or immobilized anti-CD3 (T1P1D4+C6, American Research Products) for T cell stimulation and activation. The CD8+ or CD4+ T cell proliferation was evaluated using the CFSE Cell Division Tracker Kit (423801, BioLegend). MDSCs were added to the coculture at different ratios to CD8+ T cells or CD4+ T cells for 72 h (CD8+ T cells/CD4+ T cells to MDSCs = 4 : 1 and 2 : 1), 48 h after T cell stimulation. The cells were then collected and treated with the APC-CD8+ T cell antibody (344722 for humans and 100712 for mice, BioLegend) or APC-CD4+ T cells (357408 for humans and 100412 for mice, BioLegend) prior to flow cytometry. The cell culture passage number was no more than 10 for all isolated cells, and three replicates were performed in each passage. All experiments were performed in triplicate.

2.8. Western Blot Analysis. The cells were completely lysed in proteinase-containing radioimmunoprecipitation assay buffer inhibitor (4693132001, Complete Roche). Protein quantification was applied using the protein assay reagent (5000001, Bio-Rad). Further, 10 μg of total protein was used for subsequent western blot (WB) analysis. The lysates were electrophoresed and transferred to Hybond nitrocellulose paper (FFN08, Beyotime). The nonspecific binding was blocked with 5% skimmed milk in PBS with 0.1% Tween-20 (P0220, Beyotime). For target protein examination, the blots were treated with the primary anti-human or anti-mouse antibodies at 4°C overnight. The blots were rinsed with PBS and treated with the HRP-conjugated goat anti-rabbit IgG antibody (ab6702, Abcam) for 60 min at 20°C. The target proteins were detected using the ECL WB substrate kit (ab65623, Abcam), and blots were applied to Amersham Imager 600 (GE Healthcare Life Sciences). The densitometry readings/intensity ratio of each band was analyzed by using ImageJ software (version 2.0.0, Rawak Software, Inc., Germany). All experiments were performed in triplicate. All relevant antibodies are listed in Table S2.

2.9. RNA Sequencing. Total RNA was obtained and enriched for mRNA using MicroPoly A Purist (AM1919, Ambion). The RNA quantity and quality were assessed using Bioanalyzer 2100 (Agilent Technologies). RNA sequencing was performed using the Illumina HiSeq 2000
Figure 1: Continued.

(a) Normal BC vs BC IL-6 (pg/ml) showing a significant difference indicated by ****.

(b) Comparison of normal and BC tissue images with 50μm scale.

(c) Recurrence BC IL-6 (pg/ml) with ** indicating a significant difference.

(d) Recurrence BC tissue images with 50μm scale.

(e) Survival probability over time in days for different expression levels: High expression (n = 134) vs Low/medium expression (n = 397) with p = 0.00065.

(f) Tumor volume in cm³ over time (days) with NC and IL-6 expression levels.
platform (Illumina) for each sample. Digital gene expression profiles were used to analyze the differential gene expression between groups. The pathway enrichment was analyzed to further elucidate the cellular mechanisms.

2.10. Statistical Analysis. The results were expressed as means ± standard deviation. Analyses of variance and unpaired or paired-sample Student’s t-tests were used for in vitro and in vivo experiments.

3. Results

3.1. IL-6 Levels Were Elevated in MIBC Cells and Associated with Cancer Progression. MIBC tissues (n = 200) were collected and compared with the normal urothelium (n = 200), which included 11 recurrent MIBC tissues to test the IL-6 expression level in MIBC cells. ELISA showed that the IL-6 level was higher in MIBC tissues than in adjacent noncancer tissues (Figure 1(a)). Immunochemistry showed that IL-6 staining was more obvious in MIBC tissues than in adjacent tissues (Figure 1(b)). Also, IL-6 levels were compared between recurrent and primary MIBC. ELISA showed that the IL-6 levels were higher in recurrent MIBC tissues than in primary MIBC tissues. Immunochemistry also showed that IL-6 staining was more obvious in recurrent MIBC tissues than in primary MIBC tissues (Figures 1(c) and 1(d)). According to TCGA database, high IL-6 expression was associated with a worse prognosis in patients with MIBC than in patients with low IL-6 levels (Figure 1(e)). In vivo, MIBC mouse models were treated with IL-6 for the analyses of tumor growth rates and tumor weights. IL-6 promoted MIBC growth (Figures 1(f)–1(g)). Mice with MIBC treated with IL-6 showed shorter survival times compared with untreated control mice (Figure 1(h)). These results showed that IL-6 levels were elevated in MIBC tissues, especially in cases of recurrence, and promoted MIBC progression by accelerating proliferation.

3.2. Number of MDSCs Increased in MIBC Tissues and Promoted Cancer Progression. MDSCs are a heterogeneous cluster of immune cells deriving from the myeloid lineage with an immune suppression role in MIBC. MDSCs were identified as CD45+CD11b+HLA-DR-CD33+ by flow cytometry as previously studied [15], and the frequency of MDSCs was found to be higher in MIBC tissues than in the normal urothelium (Figures 2(a) and 2(b)). The increase in the number of MDSCs was greater in recurrent MIBC than in primary MIBC tissues (Figures 2(c) and 2(d)). In vivo, MDSCs were adoptively transferred into mice with MIBC, and cancer progression was evaluated. MDSCs increased tumor growth rates (Figure 2(e)). The tumor weights were higher in the MDSC-treated group compared with the control group (Figure 2(f)). Based on survival time, MDSC transfer in the MIBC mouse model was related to a worse prognosis (Figure 2(g)). Taken together, MDSCs increased in MIBC tissues and promoted cancer progression.

3.3. IL-6 Promoted MDSC Proliferation in MIBC Tissues. IL-6 is closely related to several immune responses, including MDSC-induced immune suppression. A total of 100 MIBC tissues were collected to explore the relationship between IL-6 and MDSCs. Immunohistochemistry and

![Picture](https://via.placeholder.com/150)
Figure 2: Continued.
Flow cytometry were used to evaluate IL-6 levels, and flow cytometry was used to quantify MDSCs. The IL-6 levels were positively related to MDSCs (Figures 3(a) and 3(b)). In vitro, MDSCs were sorted from mouse MIBC tissues and treated with or without IL-6. As summarized in Figure 3(c), the CCK-8 assay showed that the rate of proliferation was higher in IL-6-treated MDSCs compared with the control cells (Figure 3(d)). CFSE further demonstrated that IL-6 promoted the proliferation of MDSCs (Figure 3(e)). The flow cytometry images also showed that IL-6 increased the proportion of MDSCs in both humans and mice (Figures 3(f) and 3(g)). These results showed that IL-6 accelerated MDSC accumulation by promoting proliferation.

3.4. IL-6 Improved the Immune-Suppressive Activity of MDSCs. MDSCs were sorted from human and mouse MIBC tissues to explore whether IL-6 influenced the immune-suppressive effects of MDSCs. MDSCs treated with or without IL-6 were collected and prepared for follow-up experiments (Figure 4(a)). RT-PCR showed that IL-6 treatment increased Arg1 and iNOS levels (Figures 4(b)–4(e)). MDSCs were cocultured with CD8+ T cells or CD4+ T cells at different ratios (T cells : MDSCs = 1 : 4 or 1 : 2, respectively). IL-6-treated MDSCs significantly impeded the proliferation of CD8+ T cells and CD4+ T cells (Figures 4(f)–4(i)). Accordingly, IL-6 promoted the proliferation of MDSCs and immune-suppressive activity.

3.5. IL-6 Activated the MAPK Signaling Pathway in MDSCs. MDSCs were collected from human and mouse MIBC tissues to explore the specific mechanism underlying the effect of IL-6 on MDSCs. MDSCs were treated with IL-6 for 48 h and used for RNA sequencing (Figures 5(a) and 5(b)). Differentially expressed genes were enriched for several signaling pathways, especially the MAPK signaling pathway (Figures 5(c) and 5(d)). To confirm this result, WB analysis was performed to evaluate the key proteins with or without phosphorylation in the MAPK signaling pathway (MEK2, ERK, and MNK1). IL-6 treatment induced the phosphorylation of MEK2, ERK, and MNK1 in MDSCs from both human and mouse MIBC tissues (Figures 5(e)–5(i)). These results indicated that IL-6 promoted the proliferation and immune-suppressive effect of MDSCs by upregulating the MAPK signaling pathway.

3.6. IL-6 Promoted MDSC Proliferation and Immune-Suppressive Effects by Activating the MAPK Signaling Pathway. Aloesin was used to block the activation of the MAPK signaling pathway so as to confirm the role of the MAPK subfamilies in MDSC proliferation and immune-suppressive effects. The CCK-8 assay and flow cytometry results showed that aloesin partially attenuated the IL-6-induced increase in proliferation (Figures 6(a) and 6(b)). RT-PCR showed that IL-6 treatment increased Arg1 and iNOS levels, while IL-6 could reverse this phenomenon (Figures 6(c)–6(f)). MDSCs were cocultured with CD8+ T cells or CD4+ T cells at specific ratios (MDSCs : T cells = 1 : 4). IL-6-treated MDSCs significantly impeded the proliferation of CD8+ T cells and CD4+ T cells, which was partially reversed by aloesin (Figures 6(g)–6(j)). These results elucidated that IL-6 promoted the proliferation of MDSCs and immune-suppressive effects, and these effects could be reversed by targeting the MAPK signaling pathway. Taken together, IL-6 promoted the proliferation and immunosuppressive function of MDSCs via the MAPK signaling pathway in MIBC (Figure 7).
Figure 3: Continued.
Figure 3: IL-6 promoted MDSC proliferation in MIBC tissues. (a, b) Correlation between IL-6 (AOD and MFI) and MDSCs in MIBC tissues. (c) Experimental design of MDSC extraction and IL-6 treatment. (d) CCK-8 assays were used to test MDSCs with or without IL-6 treatment. (e) MDSC proliferation rates were evaluated according to fluorescence attenuation. (f, g) Flow cytometry images and statistical analysis of MDSCs from humans and mice with or without IL-6 treatment. Mean ± SD, ***P < 0.005, and ****P < 0.001. AOD: average optical density; MFI: median fluorescence intensity.
Figure 4: Continued.
4. Discussion

IL-6 is highly pleiotropic and has a wide range of functions in various diseases [16]. The TME of most malignant lesions shows immune cell enrichment with immunosuppressive phenotypes, often via the immunomodulatory activities of IL-6-family cytokines. The members of this family have been identified as diagnostic or prognostic biomarkers of the response to therapy and disease activity in various cancers [17]. Aberrant IL-6 family overexpression and downstream receptor signaling pathway stimulation are frequent cytokine events in carcinomas and are often correlated with poor clinical outcomes [18]. The tumorigenesis-promoting effects of IL-6 cytokine family members are mediated via direct effects on proliferation, survival, migration, invasion, and metastasis in cancer cells and indirect effects on immunosuppression, modulation of inflammation, and angiogenesis in the interstitial stromal cells, which form the local TME [19]. IL-6 is expressed in numerous BC cell lines and is more abundant in the specimens of patients with MIBC than those with NMIBC [20]. Both BC cell lines and primary BC tissues express IL-6 and IL-6 receptor protein [21]. IL-6 clinically serves as an alternative marker of host immunity in patients with BC [22]. The canonical activation of immunity in BC elicits the production of IL-6, leading to BC-associated immune malfunction [23]. Typically, serum IL-6 level is significantly higher in higher-grade patients than in low-grade patients; furthermore, it is linked to patients with recurrent BC [24]. Patients with elevated IL-6 levels have a high risk of distant metastases and lymphatic metastases [25].

Figure 4: IL-6 improved the immune-suppressive effects of MDSCs. (a) Experimental diagram of MDSC extraction and IL-6 treatment. (b–e) RT-PCR was used to evaluate Arg1 and iNOS levels in MDSCs with or without IL-6 treatment. (f–i) Summary of the suppressive effect of MDSCs by coculture with CD8$^+$ T cells and CD4$^+$ T cells at 1 : 4 and 1 : 2 ratios. Mean ± SD, *$P < 0.05$, **$P < 0.01$, ***$P < 0.005$, and ****$P < 0.001$. ns: no statistical significance.
**Figure 5: Continued.**

(a) Human

(b) Mice

(c) Human

(d) Mice

(e) Human

(f) Mice

Signaling pathways

-Log10 (p value)

<table>
<thead>
<tr>
<th>Human</th>
<th>MAPK signaling</th>
<th>C9orf72-mediated autophagy</th>
<th>Pentose phosphate pathway</th>
<th>Extrinsic apoptotic</th>
<th>ITGA/B-FAK-RAC signaling</th>
<th>FAS-JNK signaling</th>
<th>Benzene degradation</th>
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<tr>
<td>Mice</td>
<td>MAPK signaling</td>
<td>FAS-JNK signaling</td>
<td>ATP6-mediated transcription</td>
<td>Pentose phosphate pathway</td>
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Among IL-6 cytokine family members, IL-6 has the most well-defined effect on the TME by prompting chronic inflammatory responses, regulating tumor vasculogenesis and the outgrowth of heterogeneous cells, and preventing Th1 cell-mediated antitumor immunocytotoxicity [26]. These protumorigenic effects are mediated by the recruitment, retention, activation, and function of TILs and reflect the various innate and adaptive immune responses regulated by IL-6 in immunity and chronic inflammatory diseases [27]. In melanoma, however, IL-6 exerts antitumor effects in the early stages of disease progression and later promotes tumor angiogenesis and the recruitment of immunosuppressive myeloid cells, such as MDSCs, to the TME [28]. Additionally, the tumor-promoting effects of IL-6 involve increases in MDSCs and Th17 cells, suppression of DCs and cytotoxic T cells, and phenotypic switching of tumor-associated macrophages (TAM) from an antitumor M1 phenotype to an immunosuppressive M2 phenotype [26]. In gastric cancer, serum-derived IL-6 activates and induces MDSCs, which express arginase 1, via the PI3K-Akt signaling pathway, thereby suppressing cytotoxic T cell functionality [29]. In hepatocellular cancer, the interaction between MDSCs and IL-6 promoted the generation of a chemoresistant phenotype [30]. However, in BC, low IL-6 levels could inhibit the activation of STAT3, promoting the formation of MDSCs [31]. The IL-6-induced experiments were simply performed on MDSCs in the serum of patients with BC. The serum MDSCs might not completely represent MDSCs in the TME of BC where elusive cell-cell communications are abundant, and MDSCs can be immunomodulated and undergo various alterations. The present analysis showed that IL-6 could stimulate the proliferation of MDSCs from the BC microenvironment in a time- and dose-dependent manner. Also, it indicated that further analysis, such as dose-dependent experiments for IL-6-induced MDSC alteration, should be implemented and more details should be elucidated.

**Figure 5:** IL-6 activated the mitogen-activated protein kinase (MAPK) signaling pathway in MDSCs. (a, b) Heat map of RNA sequencing results to evaluate differentially expressed genes in MDSCs with or without IL-6 treatment. (c, d) Differentially expressed genes were enriched for several signaling pathways. (e, f) Phosphorylation of critical proteins in the MAPK signaling pathway (MEK2, ERK, and MNK1) was evaluated by western blotting. (g–i) Intensity ratio of each band. Mean ± SD, *P < 0.05, **P < 0.01, and ***P < 0.005.
Figure 6: Continued.
with patients with low frequencies of these myeloid cells [33]. Peripheral MDSCs have a high correlation with tumor progression and grade, and low MDSC levels in circulation are significantly associated with an improved prognosis and outcomes in BC [34].

This study investigated the mechanisms by which IL-6 activated MDSCs, identifying the important role of the MAPK signaling pathway in MDSC proliferation in MIBC. Nevertheless, the precise interactions between IL-6 and MAPK are still unclear. Bongartz et al. reported that IL-6-induced MAPK activation could be categorized into a Gab1-independent early phase and a Gab1-dependent late phase. Gab1-independent MAPK activation in the early phase is vital for the follow-up launch of the Gab1-dependent expansion of MAPK activation, with the binding of the IL-6 receptor complex and SH2 domain-containing phosphatase 2 (SHP2). The successive synergetic recruitment of Grb2 and SHP2 to Gab1 is crucial for the Gab1-dependent augmentation of IL-6, eliciting MAPK signaling pathway activation in the late phase and the upregulation of various genes [35]. Additionally, MAPK cascade activation caused by IL-6 relies critically on the recruitment of SHP2 to Y759 of phosphorylation in the cytoplasmic territory of gp130 [36].

ERK, a member of the MAPK family, is a survival-promoting factor in immunocytes, indicating that its activity may harbor immune-stimulating or immune-suppressive effects relying on the cell types penetrating the solid tumor [37]. ERK can activate the amplification of CD8+ TILs and promote the secretion of lytic granules and cytokines, upgrading the cytotoxicity of TILs in the TME [38]. The critical role of immature MDSCs in immune evasion was mediated by ERK in the present study. Generally, a subset of MDSCs shared numerous similarities with TAM; however, the upregulation of the MAPK pathway could polarize macrophages from the

Figure 6: IL-6 promoted MDSC proliferation and immune-suppressive effects by activating the MAPK signaling pathway. (a) CCK-8 assays were used to evaluate MDSCs with or without IL-6 and/or aloesin treatment. (b) MDSC proliferation rates were evaluated according to fluorescence attenuation after IL-6 and/or aloesin treatment. (c–f) RT-PCR was used to evaluate Arg1 and iNOS levels in MDSCs with or without IL-6 and/or aloesin treatment. (g–j) Summary of the suppressive effect of MDSCs by coculture with CD8+ T cells and CD4+ T cells at 1:4 ratios. Mean ± SD, *P < 0.05, **P < 0.01, and ***P < 0.005.
M2 phenotype to M1 phenotype and lead to the inhibition of the suppressive effects of MDSCs by the stimulation of LPS [39]. STAT3 inhibition could decrease MAPK and lead to the apoptosis of MDSCs [40]. In BC, tumor cells induced MDSC aggregation and expansion in the TME via the CXCL2/MIF-CXCR2 signaling pathway and the elevated phosphorylation of p38, ERK, and p65 [41].

In summary, the data suggested that IL-6 promoted MIBC progression by not only accelerating proliferation but also improving the immune suppression ability of MDSCs via activating the MAPK signaling pathway.

**Abbreviations**

- BC: Bladder cancer
- NMIBC: Non-muscle-invasive bladder cancer
- MIBC: Muscle-invasive bladder cancer
- TIL: Tumor-infiltrating lymphocyte
- TME: Tumor microenvironment
- MDSC: Myeloid-derived suppressor cell
- IL-6: Interleukin 6
- DC: Dendritic cell
- MAPK: Mitogen-activated protein kinase
- ERK: Extracellular signal-regulated kinase
- ELISA: Enzyme-linked immunosorbent assay
- CFSE: Carboxyfluorescein diacetate succinimidyl ester
- RT-PCR: Reverse transcription-polymerase chain reaction
- HPRT: Hypoxanthine phosphoribosyltransferase
- SD: Standard deviation
- CCK-8: Cell Counting Kit-8
- ARG1: Arginase 1
- iNOS: Inducible nitric oxide synthase
- MEK2: Mitogen-activated protein kinase 2
- MNK1: Mitogen-activated protein kinase-interacting kinase 1
- PI3K: Phosphoinositide 3-kinase
- AKT: Protein kinase B
- Stat3: Signal transducer and activator of transcription 3
- Gab1: GRB2-associated-binding protein 1
- SHP2: SH2 domain-containing phosphatase 2
- LPS: Lipopolysaccharide
- CXCL2: C-X-C motif chemokine ligand 2
- CXCR2: C-X-C chemokine receptor 2
- AOD: Average optical density
- MFI: Median fluorescence intensity.

**Data Availability**

The detailed procedures of methods and figures are attached. The data used to support the findings of this study are available from the corresponding author upon request.

**Additional Points**

*Précis.* IL-6 strengthens the immunosuppressive ability of MDSCs via the MAPK signaling pathway in MIBC.

**Ethical Approval**

This work was approved by the Ethical Committee of Shanghai Jiao Tong University according to the Chinese Ethical Regulations. All animal experiments complied with the Shanghai Jiao Tong University of Medicine Policy on the Care and Use of Laboratory Animals.

**Consent**

The authors confirmed that we have obtained written consent from the patients to publish the manuscript.
Disclosure

The funders had no role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

Conflicts of Interest

The authors declare that they have no competing interests.

Authors’ Contributions

Junhua Zheng was responsible for the conceptualization and funding acquisition. Ke Wu was responsible for the supervision and resources. Zhihong Liu was responsible for the project administration and funding acquisition. Zhong Zheng was responsible for the methodology, software, and validation. Xinyi Zheng was responsible for the formal analysis and investigation. Yiwen Zhu was responsible for the data curation and visualization. Zhixian Yao was responsible for the software and writing of the original draft. Weiguang Zhao was involved in the writing of the original draft. Youjia Zhu was involved in the review and editing of the manuscript. Feng Sun was involved in the review and editing of the manuscript. Xingyu Mu was involved in the writing of the original draft. Yong Wang was involved in the writing of the original draft. Wanqing He was involved in the review and editing of the manuscript. Zhong Zheng, Xinyi Zheng, and Yiwen Zhu contributed equally to this work.

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Supplementary Materials

Table S1: monoclonal antibodies used for flow cytometric assay. Table S2: antibodies used for western blotting. Table S3: statistics of the RNA-seq data for human MDSCs. Table S4: statistics of the RNA-seq data for mice MDSCs. (Supplementary Materials)

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