DDX5 Silencing Suppresses the Migration of Basal cell Carcinoma Cells by Downregulating JAK2/STAT3 Pathway

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Abstract
Basal cell carcinoma is driven by the aberrant activation of hedgehog signaling. DEAD (Asp-Glu-Ala-Asp) box protein 5 is frequently overexpressed in human cancer cells and associated with the tumor growth and invasion. The purpose of this study was to investigate the role of DEAD (Asp-Glu-Ala-Asp) box protein 5 in the growth, migration, and invasion of basal cell carcinoma. The role of DEAD (Asp-Glu-Ala-Asp) box protein 5 was detected by quantitative real-time polymerase chain reaction, Western blot, and terminal deoxynucleotidyl transferase-mediated deoxyuridine triphosphate nick-end labeling assay in basal cell carcinoma cells. The associations between JAK2/STAT3 pathway and DEAD (Asp-Glu-Ala-Asp) box protein 5 were analyzed in basal cell carcinoma cells. Results showed that DEAD (Asp-Glu-Ala-Asp) box protein 5 is overexpressed in basal cell carcinoma cells. DEAD (Asp-Glu-Ala-Asp) box protein 5 knockdown inhibited the migration and invasion of basal cell carcinoma cells. DEAD (Asp-Glu-Ala-Asp) box protein 5 knockdown increased the apoptosis of basal cell carcinoma cells induced by tunicamycin. Results found that DEAD (Asp-Glu-Ala-Asp) box protein 5 knockdown increased JAK2 and STAT3 expression in basal cell carcinoma cells. JAK2 inhibitor decreased STAT3 expression and abolished the inhibitory effects of DEAD (Asp-Glu-Ala-Asp) box protein 5 silencing on migration and invasion in basal cell carcinoma cells. In conclusion, these results indicate that DEAD (Asp-Glu-Ala-Asp) box protein 5 is a potential target for inhibiting basal cell carcinoma cells growth, migration, and invasion by downregulating JAK2/STAT3 pathway.

Keywords
DDX5, BCC, migration, invasion, JAK2, STAT3

Abbreviations
BCC, basal cell carcinoma; cDNA, complementary DNA; DDX5, DEAD (Asp-Glu-Ala-Asp) box protein 5; DMEM, Dulbecco modified Eagle medium; HRP, horseradish peroxidase; IgG, immunoglobulin G; JAK2IR, JAK2 inhibitor; mRNA, messenger RNA; mAb, monoclonal antibodies; NBC, normal basal cell line; PBS, phosphate-buffered saline; qRT-PCR, quantitative real-time polymerase chain reaction; siRNA, small interfering RNA; STAT3IR, STAT3 inhibitor; TUNEL, terminal deoxynucleotidyl transferase-mediated deoxyuridine triphosphate nick-end labeling

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Introduction
Basal cell carcinoma (BCC) is one of the most common type of human skin cancers.1 Studies have found that BCC is characterized by BRCA2, BRCA1-associated protein-1 gene and P53 gene mutations, and aberrant expression of hedgehog signaling pathway.2-4 Previous study reports that the pathologic differential diagnosis of BCC includes cutaneous mixed sweat gland
tumor of the skin, myoepithelioma, and myoepithelial carcinoma. Basal cell carcinoma is associated with exposure to radiation in various populations, such as atomic bomb survivors, radiologists, and interventional cardiologists. Metastatic BCC is exceedingly uncommon, with a poorly defined natural history, and its incidence, risk factors, patterns of spread, prognosis, and potential treatment options are not well understood.

DNA replication plays an important process in BCC cells growth and invasion that can be exploited in cancer therapy. DEAD (Asp-Glu-Ala-Asp) box protein 5 (DDX5) is a prototypical member of the DEAD/H-box protein family and involves in ontogenesis and spermatogenesis in several human malignancies. DEAD (Asp-Glu-Ala-Asp) box protein 5 is also an ATP-dependent RNA helicase, which is first identified as a potential target for the treatment of breast cancer. Study has showed that DDX5 locus is frequently amplified in breast cancer and it can regulate DNA replication, which is required for cell proliferation in a subset of breast cancer cells. In addition, findings show that DDX5 plays an important role in the proliferation and tumorigenesis of non-small-cell lung cancer cells by activating the β-catenin signaling pathway, suggesting DDX5 may serve as a novel prognostic marker and potential therapeutic target in the treatment of non-small-cell lung cancer. Furthermore, knockdown of DDX5 inhibits the proliferation and tumorigenesis in esophageal cancer, and it may be a novel potential therapeutic target for the prevention and treatment of esophageal cancer. However, the expression and biological role of DDX5 in BCC remains largely unknown.

In this study, we examined the role of DDX5 in regulating BCC cell growth and invasion and explored its possible molecular mechanism. Report has showed that inhibition of JAK/STAT3 pathway involves in angiogenic activity in BCC cell and a study indicates that DDX5 expression is associated with tumor cells growth. In addition, overexpression of STAT3 inhibited BCC's tumor progression in the inflammatory microenvironment. Furthermore, activation of phosphorylated STAT3 increases viability of BCC cells and promote BCC development. Therefore, we assumed that DDX5 regulated the growth and invasion of BCC cells through JAK/STAT3 signaling pathway.

Materials and Methods

Ethical Statement

This study was approved by the Ethical Committee of Tongren Hospital, Shanghai Jiao Tong University School of Medicine (Shanghai, China). All patients were required to write informed consent.

Clinical Tissues

Twenty patients with micronodular BCC were recruited in this study. A total of 20 micronodular BCC tissues and matched adjacent nontumor tissues (10 cm far away from BCC tissues) were collected from Shanghai Jiao Tong University School of Medicine (Shanghai, China) during August 2014 to July 2018.
Table 1. Primers for qRT-PCR.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward (5'-3')</th>
<th>Reverse (5'-3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>Caspase-3</td>
<td>CATGATTAGCAAGTTACAGTGATGC</td>
<td>CACAGTCTTAAAGTGCGGGGGA</td>
</tr>
<tr>
<td>Caspase-9</td>
<td>CGTTCTATGTTACCGACATGACG</td>
<td>GTTCCATACTGACATGACATG</td>
</tr>
<tr>
<td>Bcl-xl</td>
<td>GCTGGTGTTGACTTTTCTTCC</td>
<td>GGCCTAGTCGTCGTCGTCGTCGTCGTCG</td>
</tr>
<tr>
<td>Bcl-2</td>
<td>GATGAAATCTCATCATTATAAGCTGTCACA</td>
<td>'GGCGTACGCCCTGTCCACCTGTGTCGTCAC</td>
</tr>
<tr>
<td>DDX5</td>
<td>GCCGGGACCGAGGGTTTGTTTGGT</td>
<td>CTTGTCGTCGGGCTCATCAGCCA</td>
</tr>
<tr>
<td>JAK2</td>
<td>TTATCCTACTTCTCGTCTCCACAGTC</td>
<td>TTACATCTCTCGTCTCCACAGTA</td>
</tr>
<tr>
<td>STAT3</td>
<td>AGCTGACGTGGTGTGACAGT</td>
<td>ACCCATGGGATTACACTTTG</td>
</tr>
<tr>
<td>β-Actin</td>
<td>CGGAGTCAACGGATTTGTC</td>
<td>AGCCTTCTCCATGGTGTGA</td>
</tr>
</tbody>
</table>

Abbreviations: DDX5, DEAD (Asp-Glu-Ala-Asp) box protein 5; qRT-PCR, quantitative realtime polymerase chain reaction.

Table 1. Polymerase chain reaction thermocycling conditions were 45 amplification cycles, denaturation at 95°C, primer annealing at 64°C with touchdown to 56°C, and applicant extension at 72°C. Relative levels of mRNA expression were calculated using the $2^{-\Delta\Delta C_{t}}$ method and results were expressed as a fold change relative to β-actin control.

**Western Blotting**

Basal cell carcinoma or normal basal cells were homogenized in lysate buffer containing protease inhibitor (Sigma-Aldrich, St. Gallen, Switzerland) and centrifuged at 8000 × g at 4°C for 10 minutes. Protein concentration was measured by a bicinechonic acid protein assay kit (Thermo Scientific, Pittsburgh, Pennsylvania). Subsequently, protein samples (40 μg) were loaded and separated using 15% sodium dodecyl sulfate-polyacrylamide gel electrophoresis, as described previously. Subsequently, proteins were subsequently blotted on a nitrocellulose membrane and hybridized using rabbit antihuman primary antibodies: DDX5 (1:2000, ab21696, Abcam, Cambridge, UK), Claudin3 (1:1000, ab15102, Abcam), MTA3 (1:1000, ab87275, Abcam), Caspase-3 (1:1000, ab238440, Abcam, Cambridge, UK), Caspase-9 (1:1000, ab32539, Abcam), Bcl-2 (1:1000, ab32124, Abcam, Cambridge, UK), Bcl-xl (1:1000, ab32370, Abcam, Cambridge, UK), and β-actin (1:1000, ab8226, Abcam, Cambridge, UK) after blocking in 5% bovine serum albumin (Sigma-Aldrich) for 1 hour at 37°C. Membranes were then incubated with horseradish peroxidase (HRP)-conjugated goat antirabbit immunoglobulin G (IgG) monoclonal antibody (mAb; PV-6001, ZSGB-BIO, Beijing, China) secondary antibodies for 24 hours at 4°C. The membrane was also washed with TBST.
for 3 times and protein bands were detected by an enhanced chemiluminescence detection system, and the band intensities were analyzed by ImageJ software 1.2.

Cell Migration and Invasion analysis
Basal cell carcinoma cells were transfected with siR-DDX5 and/or treated with JAK2IR (1 mg/mL, 420099, Sigma-Aldrich, St. Gallen, Switzerland) or STAT3IR (1 mg/mL, 573125, Sigma-Aldrich, St. Gallen, Switzerland); 1 × 10^5/well concentration of the BCC cells with 150 µL serum free DMEM were added into the upper chamber with the noncoated membrane. Matrigel-unicoated and -coated migration inserts (8 µm pore size; Millipore, Bedford, MA, USA) were used to evaluate cell migration and invasion. After 24 hours incubation, BCC cells were fixed in 4% paraformaldehyde for 10 minutes at 37°C. Cells were washed with PBS 3 times and stained with 0.1% crystal violet dye (Sigma-Aldrich, St. Gallen, Switzerland) for 15 minutes at 37°C. The cells were removed with a cotton swab and counted at 3 randomly selected views using a light microscope (Olympus BX51, Olympus; Tokyo, Japan).

Immunohistochemistry Analysis
Basal cell carcinoma tissues and matched adjacent nontumor tissues were fixed in 4% paraformaldehyde overnight and then embedded in paraffin wax; 4 µm BCC tissue sections were deparaffinized in xylene, rehydrated through graded ethanols, followed by blocking of endogenous peroxidase activity in 3% hydrogen peroxide for 10 minutes at room temperature and analyzed for DDX-5 expression. Tumor sections were incubated with specific primary antibodies for DDX5 (1:2000, ab21696, Abcam) for 12 hours at 4°C. Tumor tissues were then incubated with HRP-conjugated goat anti-rabbit IgG mAb (1:5000, dilution, PV-6001, ZSGB-BIO). A Ventana Benchmark automated staining system was used for purpose protein expression in tumor tissues (Olympus BX51, Olympus). The staining results were semiquantitatively evaluated by the multiplicity of staining intensity and the percentage of positive staining cells (magnifications: ×400).

Terminal Deoxynucleotidyl Transferase-Mediated Deoxuryridine Triphosphate Nick-End Labeling Assay
The treated BCC cells (1 × 10^6) were treated with tunicamycin (1 µg/mL) for 4 hours at 37°C and fixed with 10% paraformaldehyde for 10 minutes at room temperature. Cells were washed with PBS and apoptosis of BCC cells was analyzed using terminal deoxynucleotidyl transferase-mediated deoxuryridine triphosphate nick-end labeling (TUNEL) assay kit (DeadEnd Colorimetric Tunel System, Promega, Madison, Wisconsin) according to the manufacturer’s instructions. Cells were immersed in 50 µL TUNEL reaction fluid in a humid environment at 37°C for 1 hour. After washing with PBS 3 times, cells were incubated with 4′,6-diamidino-2-phenylindole at 37°C for 30 minutes. Finally, samples were washed with PBS 3 times and then captured with a ZEISS LSM 510 confocal microscope at 488 nm. The apoptosis rate was calculated by using the software of Developer XD 1.2 (Definiens AG, Munich, Germany).

Statistical Analysis
Data are expressed as the mean ± standard deviation and performed at least 3 independent replicates. All data were analyzed by SPSS software 19.0 (SPSS, Inc, Chicago, Illinois). Data were analyzed using Student t test or one-way analysis of variance followed by Tukey multiple comparison post hoc tests; *P < .05 and **P < .01 were considered statistically significant.

Results
Expression of DDX5 in BCC Tissues and BCC Cell Lines
Expression of DDX5 was investigated in BCC tissues and BCC cell lines. Results showed that DDX5 expression was approximately 2.08-fold in BCC tissue samples compared with normal basal tissue (Figure 1A, P < .01). As shown in Figure 1B and C, BCC cell line expressed higher level of DDX5 gene (2.39-fold, BCC vs normal basal cell line [NBC]; P < .01) and protein (1.55-fold, BCC vs NBC; P < .01) expression level than NBC. These results suggest that DDX5 expression is upregulated in BCC tissues and cell lines.

DDX5 Knockdown Inhibits BCC Cell Line Migration and Invasion
The effects of DDX5 on migration and invasion of BCC cells were analyzed in this study. DEAD (Asp-Glu-Ala-Asp) box protein 5 knockdown (siR-DDX5) inhibited the migration and invasion of BCC cells (Figure 2A and B, P < .01 vs siR vector). Results observed that knockdown of DDX5 inhibited the expression of Claudin3 and MTA3 in BCC cells (Figure 2C and D, P < .01). These results suggest that DDX5 involves in BCC cell line migration and invasion.

DDX5 Knockdown Promotes BCC Cells Apoptosis
The effects of DDX5 knockdown on BCC cells apoptosis were analyzed in this study. As shown in Figure 3A, DDX5 knockdown increased the apoptosis of BCC cells compared to control (P < .01 vs siR vector). Gene and protein expression of the active fragment for caspase-3 and caspase-9 was increased by DDX5 knockdown in BCC cells compared to the cells transfected by siR vector (Figure 3B and C, P < .01 vs siR vector). DEAD (Asp-Glu-Ala-Asp) box protein 5 knockdown decreased Bcl-2 and Bcl-xl expression in BCC cells (Figure 3D and E, P < .01 vs siR vector). These results suggest that DDX5 knockdown promotes BCC cells apoptosis.

DDX5 Knockdown Regulates BCC Cells Metastasis via JAK2/STAT3 Pathway
The potential mechanism mediated by DEAD (Asp-Glu-Ala-Asp) box protein 5 was analyzed in BCC cells. Results found
that DDX5 knockdown increased JAK2 and STAT3 expression in BCC cells (Figure 4A and B, \( P < .01 \) vs siR vector). However, JAK2IR decreased STAT3 expression and abolished the inhibitory effects of DDX5 knockdown (siR-DDX5 + JAK2IR) on migration and invasion in BCC cells (Figure 4C-F, \( P < .01 \) vs control or siR-DDX5 + JAK2IR). These results suggest that DDX5 knockdown downregulates JAK2/STAT3 pathway in BCC cells.

**DDX5 Knockdown Regulates BCC Cells Apoptosis via JAK2/STAT3 Pathway**

The potential mechanism of apoptosis mediated by DDX5 was analyzed in BCC cells. The results showed that JAK2IR increased apoptosis and abolished the proapoptotic effects of DDX5 knockdown in BCC cells (Figure 5A, \( P < .01 \) vs control or siR-DDX5 + JAK2IR). Also, STAT3IR abolished the proapoptotic effects of DDX5 knockdown in BCC cells (Figure 5B, \( P < .01 \), vs control or siR-DDX5 + STAT3IR). Results demonstrated that JAK2IR canceled DDX5 knockdown-regulated caspase-3, caspase-9, Bcl-2, and Bcl-xl expression in BCC cells (Figure 5C-D; \( P < .01 \) vs control or siR-DDX5 + JAK2IR, \( P < .01 \) vs control or siR-DDX5 + STAT3IR, respectively). These results suggest that DDX5 knockdown regulates BCC cells apoptosis via JAK2/STAT3 pathway.

**Discussion**

Previous reports have indicated that DDX5 expression is overexpressed in human cancer tissues compared with normal tissues including breast cancer, gastric cancer, and prostate cancer.\(^{13,23,24}\) Evidence indicates that inhibition of JAK2/STAT3 signaling pathway induces cancer cells
apoptosis and decreases migration of cancer cell, such as ovarian cancer, colorectal cancer, and gastric cancer. In this study, we analyzed DDX5 expression in BCC tissue and BCC cell line. This study also explored the possible mechanism of apoptosis and invasion mediated by DDX5 in BCC cells. Results in this study indicate that DDX5 is overexpressed in BCC tissues and BCC cell lines. Findings in this study suggest that DDX5 induces BCC cells apoptosis, inhibits migration and invasion through inhibition of the JAK2/STAT3 signal pathway.

Claudins are key cells adhesion molecules, which are associated with cancer development. Study has found that MTA3 can regulate malignant progression of human cancer through various signaling pathways. Here, we reported that DDX5 knockdown inhibited migration and invasion of BCC cells. However, we will further investigate the expression of DDX5 in the more aggressive BCC subtypes based on their morphologies in our future work.

Increasing apoptotic sensitivity plays a crucial role in the treatment of human cancer. In this study, we showed that DDX5 knockdown increased the apoptosis of BCC cells induced by tunicamycin. Study has found that inducing mitochondria-dependent cleavage of caspase-9 and -3 or decreasing in Bcl-2 and Bcl-xl can promote human BCC

Figure 3. Effects of DDX5 knockdown on BCC cells apoptosis. A, DDX5 knockdown increases the apoptosis of BCC cells. Effects of DDX5 knockdown on gene (B) and protein (C) expression of pro caspase-3 and cleaved caspase-3 and caspase-9 in BCC cells. Effects of DDX5 knockdown on gene (D) and protein (E) expression level of Bcl-2 and Bcl-xl in BCC cells. **P < .01 vs siR vector. BCC indicates basal cell carcinoma; DDX5, DEAD (Asp-Glu-Ala-Asp) box protein 5.
Our data demonstrated that DDX5 knockdown induced caspase-9 and -3 expression and decreased Bcl-2 and Bcl-xl through interfering with JAK2/STAT3 signaling pathway in BCC cells.

A study has found that under *in vitro* experimental condition, activation of JAK/STAT3 signaling pathway induces BCC cells invasion. Activating STAT3 signaling pathway in BCC contributes tumor growth and tumor progression in the...
Figure 5. DEAD (Asp-Glu-Ala-Asp) box protein 5 knockdown regulates BCC cells apoptosis via JAK2/STAT3 pathway. Effects of JAK2 inhibitor (A) or STAT3 inhibitor (B) DDX5 knockdown-increased apoptosis of BCC cells. Effects of JAK2 inhibitor on DDX5 knockdown-regulated caspase-3 and caspase-9 (C), Bcl-2 and Bcl-xL (D) expression in BCC cells. **P < .01 vs control or siR-DDX5 + JAK2IR. BCC indicates basal cell carcinoma; DDX5, DEAD (Asp-Glu-Ala-Asp) box protein 5; JAK2IR, JAK2 inhibitor.
inflammatory microenvironment for BCC cells.\textsuperscript{18} Findings in this study reported that JAK2IR decreased STAT3 expression and abolished the regulatory effects of DDX5 knockdown on apoptosis, migration, and invasion of BCC cells. Findings also found that JAK2 and STAT3IR abolished DDX5 knockdown-regulated caspase-3, caspase-9, Bcl-2, and Bcl-xl gene and protein expression in BCC cells. However, other regulatory factors involved in the progression of BCC cells growth and invasion need to be further studied in our future investigation.

In conclusion, we have shown that DDX5 knockdown increased tunicamycin-induced apoptosis in BCC cells and suppressed migration and invasion of BCC cells. Our findings suggest that the mechanism of DDX5 may involve modulation of the JAK2/STAT3 signaling pathway in BCC cells. However, antitumor mechanisms of DDX5 need to be clarified further, and the experimental cancer mice performed in the future.

**Authors’ Note**
The analyzed data sets generated during the study are available from the corresponding author on reasonable request. This study was approved by the Ethic Committee of Shanghai Jiao Tong University School of Medicine. The approval number given by the Ethics Committee is 2017-020-01.

**Declaration of Conflicting Interests**
The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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**Supplemental Material**
Supplemental material for this article is available online.

**References**


