FOXQ1 promotes cancer metastasis by PI3K/AKT signaling regulation in colorectal carcinoma

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Abstract: Colorectal cancer is one of the major health problems, with invade surrounding tissues, and migrate to distant organs being the most critical concern, thus identified metastasis associated hallmarks and more efficacious treatment are urgently needed. It found that forkhead box q1 (FOXQ1) is aberrant expression in variety of human cancers and FOXQ1 is involved in oncogenic pathways. However, the role of FOXQ1 has been unexplored in colorectal cancer metastasis to date. Here, expression of FOXQ1 was higher in colorectal cancer tissue samples and cancer cell lines than in normal colorectal tissue and cell lines. Further research suggested that FOXQ1 positively regulated cell proliferation in colorectal cancer and down-regulation of CDK6, extracellular regulated protein kinases 1/2 (ERK1/2) and mammalian target of rapamycin (mTOR). In corresponding to this result, over-expression of FOXQ1 significantly promoted colorectal cancer growth in vivo. Moreover, down regulation of FOXQ1 expression in colorectal carcinoma cell HCT116 and LOVO strikingly inhibits tumor growth in vivo. Finally, FOXQ1-dependent inhibition of colorectal cancer cell migration and invasion and down-regulation of focal adhesion kinase (FAK), phosphatidylinositol 3-kinase (PI3K) phosphorylation, AKT (v-akt murine thymoma viral oncogene) phosphorylation and matrix metalloproteinase-2/9 (MMP-2/9) expression. These integrated efforts have identified FOXQ1 as a tumor promoter and might provide promising approaches for colorectal cancer metastasis treatment.

Keywords: FOXQ1, colorectal carcinoma, migration, invasion

Introduction

Colorectal cancer represents one of the most common solid gastrointestinal cancers worldwide [1]. The main curative treatments for colorectal cancer are the surgical removal of lesions, radiotherapy, and chemotherapy. Although outcomes of clinical treatment for colon carcinoma have improved in recent decades, the overall 5-year survival rate for patients is only 50% because new diagnostic and treatment methods are lacking [2]. Therefore, it is of great clinical value to explore the molecular mechanisms underlying the development of colon carcinoma and to identify effective treatment strategies to improve the survival rate of patients with colorectal carcinoma [3]. Accumulating evidence suggests that the poor therapeutic effects and dismal survival rate with colorectal carcinoma are associated with aberrantly activated signaling pathways, including epidermal growth factor receptor (EGFR) signaling [4]. In late-stage colorectal cancer, the most commonly used targeted therapies are the monoclonal antibodies cetuximab and panitumumab, which prevent EGFR activation [5]. EGFR activation has been reported to play key roles in the development and progression of colorectal carcinoma and to contributes to the malignant phenotype of human colon cancer cells. For instance, activated EGFR signaling results in colorectal carcinoma cell resistance to chemotherapy and promotes cell survival, whereas inhibition of EGFR signaling dramatically reduces the proliferation of colorectal carcinoma cells [6]. EGFR signaling is involved in the EGF-induced epithelial-mesenchymal transition (EMT) and is positively associated with lymph node metastasis of colorectal cancer. Metastasis, which precipitates death in the vast majority of colorectal cancer patients, is a multi-step process by which tumor cells disseminate from their primary site and form secondary tumors at a distant site [7]. These stud-
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ies highlight molecular mechanisms that contribute to colorectal carcinoma metastasis and indicate the great need for new colorectal cancer interventions.

Recent studies have identified alterations in forkhead box q1 (FOXQ1) and other genes as likely mechanisms of secondary resistance to anti-EGFR antibody therapy that participate in colorectal cancer metastasis [8]. As a member of the forkhead box protein family, FOXQ1 is a transcription factor that regulates cell differentiation. FOXQ1 was recently found to be expressed in carcinomas of the breast, ovaries, and endometrium, where activation of the receptor by its ligand, produced either by the tumor cells or by stromal elements, stimulated tumor cell invasion and metastasis [9]. Furthermore, the activated or phosphorylated form of FOXQ1 has been associated with enhanced invasive and metastatic potential in advanced epithelial ovarian carcinoma. Phosphorylation of FOXQ1 on specific tyrosine residues enables it to bind directly to cytoplasmic effector proteins, which in turn relays receptor-induced signals through multiple signal transduction pathways [10]. Phosphorylation of FOXQ1 creates binding sites for a variety of cytoplasmic proteins that activate signal transduction pathways, including that of ERK1/2 and PI3K [11]. Several modes of regulating FOXQ1 expression have been demonstrated in normal and tumor cells, such as microRNA and the Wnt signaling pathway. Based on the hypothesis that blockade of FOXQ1 would be beneficial in cancer treatment, specific FOXQ1 inhibitors were developed to target malignant tumors. Collectively, these results identify FOXQ1 as a promising therapeutic target for solid tumors [12]. However, further studies are needed to fully elucidate how the multifunctional properties of FOXQ1 contribute to colorectal carcinoma invasion and metastasis.

In the present study, we found that FOXQ1 expression was higher in colorectal cancer tissues and cancer cell lines than in normal colorectal tissue and a normal cell line. FOXQ1 positively regulated cell proliferation in colorectal cancer cells lines and promoted metastasis of colorectal cancer cells. Downregulation of FOXQ1 hindered cellular migration and invasion of colorectal cancer cells, whereas silencing endogenous FOXQ1 resulted in the opposite outcome. Our findings suggest that FOXQ1 plays a critical oncogenic role in colorectal cancer progression and highlights its potential as a target for colorectal cancer therapies.

Materials and methods

microRNA database analysis

To examine the FOXQ1 expressed gene, the GENE-E tool (http://www.broadinstitute.org/cancer/software/GENE-E/download.html) was utilized to conduct the expression analysis of the microarray dataset. The dataset (GDS4382) were selected for the heterogeneity analysis of paired colorectal cancer tumors and adjacent non-cancerous tissues [13]. P values lower than 0.05 were considered significantly regulated and the top 10% of genes within the dataset were selected.

Cell culture

The human colorectal cancer cell lines (LOVO, HT-29, HCT116, Caco-2) and non-tumorigenic human colonic epithelial cells FHC used as control were obtained from the Chinese Academy of Sciences Cell Bank of Type Culture Collection (CBTCCCAS, Shanghai, China). The colorectal cancer cell lines were cultured as monolayers in ATCC-formulated DMEM: F12 media supplemented with 10% fetal bovine serum (FBS), penicillin (100 μg/mL), and streptomycin (100 μg/mL) and FHC was cultured in serum-free LHC-8 medium (Invitrogen, Carlsbad, CA) maintained in an incubator with a humidified atmosphere of 5% CO₂ at 37°C.

Stable cell lines

A siRNA plasmid targeting human FOXQ1 (sc-60660; Santa Cruz Biotechnology) or a non-silencing control siRNA plasmid (sc-37007; Santa Cruz Biotechnology) was transfected into HCT116 and LOVO, using Lipofectamine™ 2000 transfection reagent (Life Technologies). A FOXQ1 expression plasmid was generated by PCR sub-cloning full-length human FOXQ1 coding sequence into the retroviral transfer plasmid pQCXIP (Clontech, CA) to generate plasmid FOXQ1. Retroviral production and infection were performed as manufacturer’s instructions. Stable cell lines expressing FOXQ1 were selected for with 0.5 g/ml puromycin 10 days after infection [14].
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Cell proliferation

The cell proliferation was determined by 3-(4, 5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide (MTT) assay. Cells (1 × 10^4) seeded in 96-well tissue culture plates were incubated for 24, 48 and 72 hours, and at the end of the culture 10 μL MTT solution (5 mg/mL; Sigma) was added to each well with 100 μL of medium for 4 hours at 37°C. The medium was replaced with 200 μL imethyl sulfoxide (DMSO), followed by incubation for 10 minutes at room temperature. The absorbance was then measured at a wavelength of 490 nm using Synergy™ HT Multi-Mode Microplate Reader (Bio-Tek, Winooski, VT, USA). The effect of siRNA FOXQ1 or FOXQ1 over-expression on colorectal cancer cell viability was assessed as the percent of cell viability compared with vehicle-treated control cells, which were arbitrarily assigned 100% viability.

Colony formation

LOVO and HCT116 cells (1 × 10^2) were seeded in 25 mm petri dishes and were allowed to grow. Subsequently, growth medium was added every week for 21 days until colonies were easily distinguishable. Then, the cell clones were fixed with methanol and stained with 0.1% crystal violet for number counting [15].

Cell migration

The scratch wound healing assay was adapted to evaluate the ability of colorectal cancer cells migration. Cells were cultured in 6-well plates until reach to 90% confluence. The cell were monolayers scratched manually with a 200 μl pipette tip, washed with PBS for twice and cultured in serum-free medium for 48 hours. Then, cells were fixed with methanol, and the scratch area was photographed [16].

Cell invasion

Cell invasiveness was determined, using a Matrigel (BD Biosciences)-coated Transwell chamber (pore size of 8-um, Corning). Briefly, LOVO or HCT116 cells (5 × 10^4 cells per well) in 100 μL of serum-free medium were seeded into the upper chamber of the Transwell, whereas 600 μL of medium containing 10% FBS was applied to the lower chamber. After incubation for 24 hours at 37°C in a CO₂ incubator, invasive cells were fixed with 4% formaldehyde, stained with 0.1% crystal violet, and counted in five random microscopic fields under a light microscope [17].

Flow cytometry

FHC, LOVO and HCT116 cells were detached by 2 mM PBS/EDTA and fixed with 4% paraformaldehyde, washed twice with PBS and re-suspended for antibody staining at 1 × 10^6 cells/100 ul in 1% PBS/BSA. Gates (in the live cell population) were drawn in order to exclude > 99% of background staining (based on the isotype-stained samples). For FOXQ1 staining, anti-FOXQ1-APC conjugated and its highly adsorbed isotype matched control (Biolegend) were used [18].

Western blotting

Cell protein extraction were prepared with RIPA lysis buffer containing both protease and phosphatase inhibitors. Equal amounts of cell lysates (30 μg) were loaded on 10% sodium dodecyl sulfate (SDS)-polyacrylamide gel electrophoresis (PAGE) and transferred onto polyvinylidene difluoride (PVDF) membranes. After membranes were blocked, they were incubated with monoclonal antibody against FAK (1:500, Signalway Antibody) and phosphor-FAK Tyr407 (1:10000, Epitomics), AKT (1:5000, Epitomics) and phosphor-AKT Ser473 (1:1000, Cell Signaling Technology), PI3K and phosphor-PI3K Tyr458 (1:1000, Cell Signaling Technology), MMP-2 (1:1000, Cell Signaling Technology), MMP-9 (1:1000, Cell Signaling Technology), CD44 (1:1000, Cell Signaling Technology), PCNA (1:1000, Cell Signaling Technology), mTOR (1:1000, Cell Signaling Technology), ERK1/2 (1:1000, Cell Signaling Technology), FOXQ1 (1:1000, Cell Signaling Technology) and GPADH (1:5000, BioWorld Biotechnology). Target proteins were detected by the ECL system (Millipore, Braunschweig, Germany) and visualized with the ChemiDoc XRS system (Bio-Rad, Hercules, CA, USA) [19].

Quantitative real-time PCR

Total RNA (0.5 μg) was isolated from cultured cells using TRIzol reagent (Invitrogen, USA). First-strand cDNA was synthesized with 1 μg
total RNA using a PrimeScript RT reagent kit (TakaraBio, Japan). Quantitative real-time PCR was performed using iQ™ SYBR® Green Supermix and the iQ5 real-time detection system (Bio-Rad Laboratories, CA). The comparative cycle threshold (Ct) method was applied to quantify the expression levels through calculating the $2^{-\Delta\Delta Ct}$ method. The primer pairs used for PCR were as follows (sense and antisense, respectively): GAPDH (forward, 5'-GGAGCGAGATCCCTCCAAAAT-3'; reverse, 5'-GGCTGTTGCTATACTTCTCATGG-3') was used as an internal control; FOXQ1 (forward, 5'-CACGCAGCATATACG-3'; reverse, 5'-CGTTGAGCGAAAGGT-3').
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TGTGG-3'). cDNAs amplification and relative expression values were obtained from three independent experiments.

Mouse xenograft model

Six-week-old athymic male BALB/c nude mice (Shanghai Slack Laboratory Animal co., Ltd) were acclimatized under controlled standard conditions for a week prior to the experiment according to Guide for the Care and Use of Laboratory Animals of Nanjing University of Chinese Medicine. LOVO or HCT116 (5 × 10⁶) cells re-suspended in 100 μl of PBS and were subcutaneously injected into the back flank area of nude mice. The diameter of the tumor was measured externally with a caliper ruler every 3 days, and the volume of the tumor (mm³) was estimated by the following formula: (short diameter) 2 × (longest diameter) × 0.5 [20]. For immunohistochemical analysis, tumor tissues were removed, formalin-fixed and paraffin-embedded. Serial 4-μm tumor sections were deparaffinized in xylene, rehydrated through descending grades of ethanol, and subjected to immunohistochemical analysis using antibodies against CD44, extracellular regulated protein kinases 1/2 (ERK1/2) and mTOR.

Statistical analysis

The Fisher exact test or the χ² test was used to evaluate the associations between categorized variables. The numerical data were compared by Student's t-test or Mann-Whitney U test. Survival rates in patients were calculated by the Kaplan-Meier method and comparison was made by log-rank test. P values less than 0.05 were considered to be statistically significant.

Results

FOXQ1 expression is elevated in colorectal cancer

We tested five representative colorectal cancer cell lines and non-tumorigenic human colonic epithelial cell line FHC for the expression of FOXQ1. Quantitative PCR revealed that all colorectal cancer cell lines expressed detectable amounts of FOXQ1 mRNA (Figure 1A). Western blotting with anti-FOXQ1 antibody confirmed higher detectable amounts of FOXQ1 in all five cell lines (Figure 1B). To determine changes in FOXQ1 in clinical specimens of colorectal cancer tissue, the level of FOXQ1 in colorectal carcinoma tissues was examined using a microarray expression database (DataSet Record GDS4382). Based on gene summary views (neoplastic vs. normal tissue), FOXQ1 was significantly upregulated in colorectal cancer tissue (Figure 1C). The top 10 differentially regulated in the dataset were selected for further analysis. The FOXQ1 gene was identified as significantly upregulated in colorectal cancer tissue based on their presence in non-cancerous tissue (Figure 1C). Fluorescence-activated cell sorting analysis after staining with anti-FOXQ1 antibody revealed the existence of HCT116 and LOVO cell subpopulations that expressed FOXQ1 (range 28.3-35.9%, Figure 1D), whereas no FOXQ1-expressing FHC cells were identified (2.9%).

FOXQ1 regulates cell proliferation and colony formation in colorectal cancer cells

To investigate the role of FOXQ1 in colorectal cells proliferation, we transfected HCT116 and LOVO cells with FOXQ1 siRNA or a wild-type FOXQ1 expression plasmid (FOXQ1) to silence or overexpress FOXQ1, respectively. Lower levels of FOXQ1 expression were observed in the cells transfected with FOXQ1 siRNA, and higher expression levels were observed in cells transfected with the FOXQ1 plasmid (Figure 2A). As shown in Figure 2B, downregulation of FOXQ1 with specific siRNA inhibited proliferation of both HCT116 and LOVO cells. A growth curve analysis showed that growth of HCT116 and LOVO cells with downregulated FOXQ1 decreased relative to that of control cells. However, overexpression of FOXQ1 in HCT116 and LOVO cells significantly promoted cell proliferation. The soft agar colony formation assay is a method used to confirm cellular anchorage-independent growth in vitro. As expected, inhibiting FOXQ1 expression in HCT116 and LOVO cells markedly decreased the efficiency of cell colony formation, whereas HCT116 and LOVO cells transfected with FOXQ1 showed an enhanced ability to form colonies (Figure 2C). To study the molecular mechanisms responsible for FOXQ1-regulated colon cancer cellular proliferation, we evaluated, by quantitative PCR, whether the effect of FOXQ1 on HCT116 and LOVO cell proliferation was accompanied by modulation of proteins that are involved in
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Tumor cell growth and survival. This revealed significant downregulation of CD44, extracellular regulated protein kinases 1/2 (ERK1/2), and mammalian target of rapamycin (mTOR) in the FOXQ1 siRNA-treated HCT116 and LOVO cells (Supplementary Figure 1). Additionally, we observed that FOXQ1 transfection increased the levels of CD44, ERK1/2, and mTOR (Supplementary Figure 1). We then assessed the impact of FOXQ1 on the protein levels of these targets by overexpressing FOXQ1 and knocking down FOXQ1 in HCT116 and LOVO cells. Consistently, western blots of these putative targets (Figure 2D) showed that high levels of FOXQ1 expression increased levels of CD44, ERK1/2, and mTOR, and that FOXQ1 knockdown resulted in significantly decreased levels of these proteins (Figure 2D). Overall, these results indicated that FOXQ1 positively regulated cell proliferation and clone formation ability in colorectal cancer cells in vitro.

**Downregulation of FOXQ1 blocks cellular migration and invasion of colorectal cancer cells**

Increased mobility is a crucial property of invasive cancer cells. To determine the effect of FOXQ1 on cellular migration, HCT116 and LOVO
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Cells were transfected with a FoxQ1 siRNA plasmid and subjected to wound-healing migration and Transwell invasion assays. Non-specific siRNA (control siRNA) was used as a negative control. As shown in Figure 3A, the cell migration rate was significantly decreased in FoxQ1-downregulated cells compared with the rate in control cells. The invasion ability of colorectal cancer cells was examined in HCT116 and LOVO cells transfected with the FoxQ1 siRNA plasmid cultured in a 24-well Transwell plate with an 8-μm pore polyester insert. As shown in Figure 3B, significantly fewer FoxQ1-downregulated HCT116 and LOVO cells showed invasion as compared to control cells. To further investigate the molecular mechanism by which FoxQ1 contributes to cellular migration and invasion in colon carcinoma, the expression of several metastasis-related proteins were determined. An immunoblotting assay was performed to assess changes in the phosphorylation state of important metastasis-associated kinases, including focal adhesion kinase (FAK), protein kinase B (AKT), and phosphatidylinositol 3-kinase (PI3K). Figure 3C shows that FAK, AKT, and PI3K phosphorylation was effectively inhibited in FoxQ1 siRNA-transfected HCT116 and LOVO cells, but not in control cells. Matrix metalloproteinases (MMPs), which are regarded as capable of degrading extracellular matrix, play a critical role in the processes of tumor invasion, migration, and metastasis. To determine whether FoxQ1 downregulation affects MMP-2/9 levels, we applied a western blot to FoxQ1 siRNA-transfected cells to examine the expression of MMP-2/9. As expected, we observed a significant reduction in MMP-2/9 expression after RNAi-mediated FoxQ1 knockdown (Figure 3D). Consistent with this result, the activity of MMP-2/9 in HCT116 and LOVO cells was also markedly decreased by siRNA mediated knockdown.
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To determine the functional role of FOXQ1 in cellular migration and invasion, HCT116 and LOVO cells that stably expressed FOXQ1 were established and subjected to wound-healing migration and Transwell invasion assays. As shown in Figure 4A, ectopic expression of FOXQ1 in colorectal cancer cells markedly increased the cell migration rate compared with the rate of cells transfected with the control vector. In addition, cell invasion potential was examined using Matrigel-coated Transwell plates. As shown in Figure 4B, the invasive abilities of HCT116 and LOVO cells were significantly increased in cells that overexpressed FOXQ1. Furthermore, phosphorylation of numerous well-characterized metastasis-related kinases was shown to be increased in FOXQ1 overexpressing cells (Figure 4C). Additionally, a western blot assay showed that overexpression of FOXQ1 remarkably upregulated MMP-2/9 expression (Figure 4D) and activity (Supplementary Figure 3). Collectively, these results suggested that FOXQ1 upregulation promotes the aggressiveness of colorectal cancer cells in vitro.

**Downregulation of FOXQ1 significantly inhibited cellular growth in vivo**

To investigate the effect of FOXQ1 expression on colorectal cancer cell growth in vivo, HCT116 and LOVO cells with downregulated FOXQ1 were subcutaneously injected into the posterior flanks of nude mice. Xenograft tumor volumes were measured every 3 d when they were pal-
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Figure 5. Down-expression of FOXQ1 significantly inhibits colorectal cancer cells growth in vivo. A. Control siRNA or siRNA against FOXQ1 were transfected into HCT116 cells and then cells was subcutaneously injected into back flank of nude mice (n = 6 each). Representative pictures of tumor in control siRNA or FOXQ1 siRNA HCT116 cell-inoculated nude mice. B. Tumor growth kinetics (mean ± SD) of FOXQ1 siRNA cells versus control HCT116 cells in nude mice (n = 6 each). The tumor volumes were measured at the indicated number of days after mice were transplanted with indicated cells. For indicated comparison, **P < 0.01. C. 21 days after xenograft implantation, mice were sacrificed. HCT116 colorectal cancer cells xenograft tumor sections were subjected to immunohistochemical analysis for CD44 (upper panel), ERK1/2 (middle panel) and mTOR (lower panel).

Discussion

Colorectal cancer remains a major clinical challenge because of poor prognosis and limited treatment options that prevent recurrence [13]. The oncogenesis of colorectal cancer involves changes in multiple oncogenes and suppressor genes. Therefore, many molecular biomarkers can be utilized for viable approaches to improving cancer prognosis and treatment [14]. Our study showed the role of a specific tumor-promotion protein, FOXQ1, in colorectal cancer. Therefore, integrated studies on the contribution of FOXQ1, with its multifunctional properties, to tumorigenesis will be important. In the current study, we found that FOXQ1 might play an important role in malignant progression of colon cancer and regulation of the FAK/PI3K/AKT signaling pathway. Preparation of a microR-
NA database and western blot analysis revealed that FOXQ1 was significantly upregulated in colon cancer tissues and cancer cell lines. Overexpression of FOXQ1 augmented the anchorage-independent growth and invasive abilities of human colon cancer cells, promoting their growth in vivo. Downregulation of FOXQ1 hindered cellular migration and invasion of colon cancer cells, as well as tumor growth in vivo. These findings provide novel insights into the potential roles of FOXQ1 dysregulation in promoting carcinogenesis and progression of colorectal cancer.

FOXQ1 expression levels are significantly higher in primary cancers than in matched noncancerous tissues [15], which suggests that downregulation of FOXQ1 in colorectal cancer might regulate the initiation of colorectal cancer [16]. Next, we investigated the role of FOXQ1 in the proliferation of cancer cells. When FOXQ1 was silenced in HCT116 and LOVO cells, the proliferation of cells decreased. In nude mice xenograft experiments, we found that downregulation of FOXQ1 significantly inhibited cellular growth in vivo. CD44, ERK1/2, and mTOR are reported to play fundamental roles in many biological processes, including cell proliferation [17]. Herein, we found that silencing FOXQ1 significantly inhibited the expression of various target genes that specifically regulate proliferation of colorectal cancer cells. Additionally, overexpression of FOXQ1 augmented the anchorage-independent growth and proliferation abilities of colorectal cancer cells in vitro, suggesting a potential role for FOXQ1 dysregulation in carcinogenesis and colon carcinoma progression.

Clinical specimens from ovarian cancer metastases have shown strong FOXQ1 immunostaining, in contrast to non-invasive borderline tumors, which do not express FOXQ1, and benign ovarian tissue, which expresses little to no FOXQ1 [18]. However, blockade of FOXQ1 signaling, which does not affect primary tumor growth, can enhance spontaneous metastasis of breast carcinoma to both the lung and bone. Given the pro-metastasis and anti-metastasis potential of FOXQ1 in cancer progression, an in vitro model that allows the role of FOXQ1 in the migration and invasion of colorectal cancer cells to be characterized may provide insight into the etiologic role of FOXQ1 in colorectal carcinoma metastasis [19]. We found, in wound healing and migration assays, that metastasis of HCT116 and LOVO human colon cancer cells was significantly decreased when the FOXQ1 gene was disrupted by siRNA. Additionally, overexpression of FOXQ1 greatly enhanced migration and invasion of HCT116 and LOVO cells. The FAK/PI3K/AKT pathway is involved in many cellular processes, including proliferation, differentiation, apoptosis, cell cycle progression, cell motility, tumorigenesis, tumor growth, and angiogenesis. PI3K is a
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member of the intracellular lipid kinase family, which catalyzes the generation of phosphatidylinositol-3,4,5-trisphosphate (PIP3) from phosphatidylinositol-4,5-trisphosphate (PIP2) [20]. Activated AKT translocates to the nucleus and activates mTOR and downstream targets [21, 22]. Phosphorylated PI3K/AKT also acts as a scaffold, primarily regulating focal signaling related to cell adhesion to the extracellular matrix and MMP-mediated matrix degradation [23, 24]. Several studies have indicated that the PI3K/AKT signaling pathway is involved in regulation of MMP-2 and MMP-9 activity [25-27]. Our results showed that overexpression of FOXQ1 in HCT116 and LOVO cells upregulates phosphorylation of FAK/PI3K/AKT, and FOXQ1 silencing reduces FAK/PI3K/AKT phosphorylation. Our findings identify FOXQ1 as a potential enhancer of metastatic potential in colorectal cancer.

Collectively, our results show that FOXQ1 was markedly upregulated in colorectal carcinoma cells and clinical colorectal cancer samples, which might be significant in colorectal cancer diagnosis. Moreover, overexpression of FOXQ1 increased colorectal carcinoma aggressiveness, including cell proliferation, migration, and invasion in vitro and growth in vivo, and activated the FAK/PI3K/AKT signaling pathway. Understanding the biological function of FOXQ1 in colorectal carcinoma progression both advances our knowledge of the mechanisms that underlie colorectal carcinoma tumorigenesis and establishes FOXQ1 as a potential therapeutic target for clinical management of and therapies for human colorectal cancers [28].

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Disclosure of conflict of interest

None.

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References

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Supplementary materials

Quantitative real-time PCR

Total RNA (0.5 μg) was isolated from cultured cells using TRIzol reagent (Invitrogen, USA). First-strand cDNA was synthesized with 1 μg total RNA using a PrimeScript RT reagent kit (TakaraBio, Japan). Quantitative real-time PCR was performed using iQ™ SYBR® Green Supermix and the iQ5 real-time detection system (Bio-Rad Laboratories, CA). The comparative cycle threshold (Ct) method was applied to quantify the expression levels through calculating the 2^(-ΔΔCt) method. The primer pairs used for PCR were as follows (sense and antisense, respectively): GAPDH (forward, 5'-GGAGCGAGATCCCTCAAAT-3'; reverse, 5'-GGCTGTTGTCATCTTCATGG-3') was used as an internal control; CD44 (forward, 5'-CTGCCGCTTTGCAGGTGTA-3'; reverse, 5'-CATTGTGGCAAGGTGCTATT-3'), ERK1/2 (forward, 5'-TACACAACTCTCGTACATCG-3'; reverse, 5'-CATGTCGAAAGCGCAGTAGATT-3'), mTOR (forward, 5'-TCCGAGAGTGAAGTCAAGAGG-3'; reverse, 5'-CACCTCCACTCTATGAGGC-3'). cDNAs amplification and relative expression values were obtained from three independent experiments.

MMP-2/9 activity assay

The activity of MMP-2/9 was determined by QuickZyme Mouse MMP-2/9 activity assay (QuickZymeBio-Sciences, Netherlands) according to the manufacturer’s instructions. Briefly, after transfection for 48 h, cells were washed with fresh medium and replaced with serum-free medium. After additional 24 h, the medium was collected and centrifuged at 10000 g for 10 min. Respective supernatant was added to the 96-well strip coated with MMP-2/9 antibody and incubated at 4°C overnight. After washing with wash buffer for 4 times, 50 µl assay buffer was added into the well, followed by adding 50 µl detection reagent. After incubation at 37°C for 1 h, OD405 was measured with Tecan (Männedorf, Switzerland).

Supplementary Figure 1. qRT-PCR was performed to evaluate the mRNA levels of CDK4, ERK1/2 and mTOR in HCT116 and LOVO cells transfected with FOXQ1 siRNA (A) or FOXQ1 (B). Histograms reporting the levels of CD44, ERK1/2 and mTOR mRNA (normalized to the GAPDH mRNA) as assessed in indicated cancer cell lines. Mean ± SD of three independent experiments. For indicated comparisons, *P < 0.05 and **P < 0.01.
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**Supplementary Figure 2.** The activity of MMP-2/9 in control siRNA or FOXQ1 siRNA transfected cells was determined by MMP-2/9 activity assay. Data are from three independent experiments and as mean ± SD. For indicated comparisons, **P < 0.01.**

**Supplementary Figure 3.** The activity of MMP-2/9 in control vector or FOXQ1 transfected cells was determined by MMP-2/9 activity assay. Data are from three independent experiments and as mean ± SD. For indicated comparisons, **P < 0.01.**

**Supplementary Figure 4.** Mean tumor weights. The data shown were represented as the mean ± SD. For indicated comparisons, **P < 0.01.**
Supplementary Figure 5. Mean tumor weights. The data shown were represented as the mean ± SD. For indicated comparisons, **$P < 0.01$. 