E2F transcription factor 8 promotes cell proliferation via CCND1/p21 in esophageal squamous cell carcinoma

Huiwen Chang
Jianxiang Song
Jixiang Wu
Yajun Zhang

Department of Cardiothoracic Surgery, Yancheng Third People’s Hospital, The Affiliated Yancheng Hospital of Southeast University, Yancheng, Jiangsu 224001, PR China

Purpose: E2F transcription factor 8 (E2F8) is a novel member of the E2F family, but its function in esophageal squamous cell carcinoma (ESCC) remains unclear. This study aimed to research the function of E2F8 in ESCC.

Materials and methods: We used quantitative real-time PCR and Western blot analyses to detect the expression pattern of E2F8 in ESCC. The effects of E2F8 on proliferation were investigated by Cell Counting Kit-8, 5-ethynyl-2′-deoxyuridine, and colony formation assays. We also confirmed the function of E2F8 in vivo.

Results: E2F8 expression was upregulated in ESCC, and promoted cell proliferation and influenced the expression of CCND1/p21. Downregulation of E2F8 expression inhibited cell proliferation in vivo.

Conclusion: E2F8 was identified as a new potential oncogene in ESCC.

Keywords: E2F8, cell cycle, ESCC, cell proliferation

Introduction

Esophageal cancer is a common kind of malignant tumor with a high incidence and mortality. Most patients are diagnosed at an advanced stage, but even at the early stage of diagnosis, the prognosis is still dismal. Almost 90% of esophageal cancers in Asia are esophageal squamous cell carcinoma (ESCC).1 Great developments have recently been made in treatments for ESCC, and surgical techniques and perioperative treatments have advanced; however, the prognosis of ESCC remains poor.2 The main reasons for the late diagnosis and poor prognosis may be due to the mechanism of ESCC still remaining unknown and the lack of efficient biomarkers for diagnosis and therapy. Therefore, it is urgent to find the key markers and basic mechanisms that can affect the response of ESCC to chemotherapy.

E2F family members can function as transcription factors that bind to target promoters to affect their expression.3 As far as we know, eight members have been identified as in this family (E2F1–8). E2F1–3 were proven to be transcriptional activators, while E2F4–7 were shown to inhibit the transcriptional expression of downstream target genes.4,6 However, the functions of E2F8, a novel member, still remain unclear. It was reported that the combination of E2F8 and E2F7 is essential for embryonic development in mice,7,8 angiogenesis,9 and lymphangiogenesis10 in zebrafish. Moreover, previous studies demonstrated that E2F8 exhibited crucial functions in different kinds of cancers. Sun et al discovered that E2F8 influenced the cell cycle and further promoted papillary thyroid cancer development.11 Deng et al
found that E2F8 may work as a potential therapeutic target in hepatocellular carcinoma. All these studies indicated that E2F8 could act as a potential oncogene in several cancers. However, the features of E2F8 in ESCC have not been investigated until now. In the present study, we identified that E2F8 was upregulated in ESCC tissue and promoted the proliferation of ESCC cells by regulating CCND1/p21.

In conclusion, our findings demonstrated that E2F8 exhibited important functions affecting ESCC proliferation and regulating the cell cycle, indicating that E2F8 could be a potential novel therapeutic target in ESCC.

Materials and methods
Tissue specimens of 60 paired tumors and their adjacent normal tissue were obtained from patients who were diagnosed with ESCC and underwent resection surgery between 2014 and 2015 at the Department of Cardiothoracic Surgery, Yancheng Third People’s Hospital, The Affiliated Yancheng Hospital of Southeast University, Jiangsu, PR China. The pathological diagnosis was made by the Department of Pathology in Yancheng Third People’s Hospital, Jiangsu, PR China. Informed consent was obtained from the patients. If the patients were not able to answer or sign, consent was obtained from their relatives before the collection of specimens. All the consents were documented with written informed consent forms and were conducted in accordance with the Declaration of Helsinki. The study was approved with the Declaration of Helsinki. The study was approved by the Medical Ethical Committee of the Medical School of Southeast University. Samples were stored at −70°C after resection until RNA extraction.

ESCC cell lines and cell culture
ESCC cell lines, including TE-1, KYSE-150, and TE-10, were purchased from American Type Culture Collection (ATCC, Manassas, MA, USA). Normal human esophageal epithelial cells (HEEC) and ESCC cells, including the EC-1 and EC-109 cell lines, were obtained from the Cell Bank of the Chinese Academy of Sciences (Shanghai, PR China). The culture conditions were as follows: Dulbecco’s Modified Eagle’s Medium (Wisent Bioproducts, St-Bruno, QC, Canada) supplemented with 10% FBS (Wisent Bioproducts), 100 U/mL penicillin, and 100 µg/mL streptomycin in a moist incubator (stabilized at 5% CO₂ and 37°C).

Knockdown and overexpression of E2F8
To knockdown E2F8, siRNAs targeting the coding region of E2F8 were obtained from GenePharma (Shanghai, PR China), and siRNA transfection reagent (Thermo Fisher Scientific, Waltham, MA, USA) was utilized according to the manufacturer’s protocols. An E2F8 inhibitor lentivirus (shE2F8) was designed based on the sequence of siRNA for E2F8. Nontargeting control siRNA was used as a negative control. The sequences used were as follows: siRNA for E2F8: 5′-GGGCAAGACUGUAUCACCTG-3′ (sense), 5′-GUGUAUCAGUCUUUGGCCC-3′ (antisense); and nontargeting control siRNA (NC): 5′-UUCCUC CGAACGUGUCACCTT-3′ (sense), 5′-ACGUGAC ACGUUCGGGAAATTT-3′ (antisense). To overexpress E2F8, ESCC cell lines were transfected with the E2F8-overexpressing recombinant vector pCDNA3.1-E2F8 (GenePharma); the empty plasmid was utilized as the negative control. The primer to overexpress E2F8 was designed as follows: 5′-CGGGATCCGAATTTACAGAATGG AGAAC-3′ (forward); 5′-CCCGTCTAGATATTGGAC ATCCTCAGGTGAGACTT-3′ (reverse).

Quantitative real-time PCR (qRT-PCR)
Following the manufacturer’s instructions, total RNA was extracted from ESCC tissue and cells with TRIzol reagent (Thermo Fisher Scientific). For qRT-PCR of the mRNA content, total RNA was reverse transcribed into cDNA with a PrimeScript RT reagent kit (Takara, Shiga, Japan). The reactions were performed using a 7,500 Real-time PCR System (Thermo Fisher Scientific) with a SYBR Premix Ex Taq kit (Takara). The specific primers for the target mRNA and internal control were designed as follows: E2F8 forward: 5′-CCAACCTGTGCTGAATA-3′ and E2F8 reverse: 5′-TTCGTGGCTTACATACCT-3′; and β-actin forward: 5′-GCATCGTCCACACTGGGC-3′ and β-actin reverse: 5′-ACCTGGCCGTCAGGCAGCT-3′. The expression pattern was normalized to the internal control (β-actin), and the results are shown in the form of relative expression and were calculated by the 2^−ΔΔCt method.

Colony formation assay
Under the culture conditions described earlier, 500 cells/well were seeded into six-well plates. After 1 week, each well underwent the following procedure: washed with PBS for three times at room temperature, fixed in ethyl alcohol for 30 seconds, and finally stained with crystal violet. Colonies were counted with the naked eye after rinsing with PBS.

Cell proliferation assay
A Cell Counting Kit-8 assay (CCK-8; Dojindo, Tokyo, Japan) was used to detect cell proliferation according to the manufacturer’s instructions. A total of 2,000 cells/well
were seeded into 96-well plates and cultured as previously described. Every 24 hours, 10 µL of CCK-8 solution was added to the serum-free medium. After 2 hours of incubation, the absorbance was detected by a microplate reader at a test wavelength of 450 nm and a reference wavelength of 630 nm.

5-Ethynyl-2′-deoxyuridine (EdU) assay
An EdU assay kit (RiboBio, Guangzhou, PR China) was used to measure cell proliferation. The cells were inoculated into 24-well plates (2×10⁴ cells/hole) and cultured for 24 hours with DMEM (10% FBS) before adding EdU (50 µM). According to the protocols, cells were incubated at 37°C for 2 hours, fixed in 4% formaldehyde for 30 minutes, and permeabilized at room temperature for 10 minutes with 0.5% Triton X-100. After washing with PBS, 1× ApolloR reaction cocktail (400 µL) was added to the EdU mixture for 30 minutes. Subsequently, Hoechst33342 (400 µL) was added for 30 minutes to visualize the nuclei. Cell images were obtained under a Nikon microscope (Nikon Corporation, Tokyo, Japan). Proliferation was analyzed using the average number of cells in three fields of vision for each sample.

Cell cycle analysis
First, the transfected cells were digested with trypsin and then centrifuged for 5 minutes at 1,200 rpm. After washing with PBS twice, the cells were immobilized overnight in 75% ethanol at −20°C. The cells were washed with PBS twice before detection, fixed with 500 µL of propidium iodide staining solution, and incubated at room temperature for 15 minutes. A BD FACSCanto II (BD Biosciences, San Jose, CA, USA) flow cytometer and FlowJo software were used to analyze the cell cycle.

Western blot analysis
According to the manufacturer’s instructions, a RIPA kit (Beyotime, Shanghai, PR China) was used to extract proteins from ESCC cells and tissue. Based on the molecular weight of each protein on the twelve alkyl SDS-PAGE and utilizing method of Electrophoresis (EMD Millipore, Bedford, MA, USA). We used 5% nonfat powdered milk in Tris-buffered saline to block the membrane for 2 hours. Then, the membranes were incubated overnight with a specific antibody at 4°C. After rinsing in TBST (Tris-buffered saline containing 0.05% Tween-20) for three times (10 minutes each time), membranes were incubated in secondary antibody solutions (rabbit or mouse) at room temperature for 2 hours and then washed again (three times, 10 minutes each time). Protein expression levels were detected by ECL Plus (EMD Millipore) with a Bio-Imaging System. The following primary antibodies were used: anti-E2F8, anti-CCND1, and anti-p21; anti-GAPDH was used as an internal control.

Statistical analysis
All experiments in this research were performed in triplicate independently. All analyses were performed using SPSS 19.0 software. The data are shown as the mean ± SD. Student’s unpaired t-test was used to determine any significant differences in the results. The following indicators were used for data that were considered statistically significant: * indicates P<0.05, ** indicates P<0.01, and *** indicates P<0.001.

Results
E2F8 was upregulated in ESCC tissue and ESCC cell lines
To identify the expression pattern of E2F8 in ESCC, 60 paired ESCC tissue and adjacent normal tissue samples were obtained. Two methods, qRT-PCR and Western blotting, were utilized to detect the expression of E2F8. The qRT-PCR results showed that E2F8 expression was upregulated in ESCC tumor tissue compared to adjacent normal tissue in the 60 paired ESCC tumor tissue and adjacent normal tissue samples. E2F8 protein expression was also upregulated in six paired ESCC tumor tissue and adjacent normal tissue samples assessed by Western blotting (Figure 1A). The expression levels of E2F8 in ESCC cell lines, including EC-1, TE-1, EC-109, KYSE-150, and TE-10, were further investigated. Compared to normal esophageal epithelial HEEC cells, ESCC cells lines showed upregulated E2F8 expression by both qRT-PCR and Western blotting (Figure 1B). In summary, E2F8 expression was upregulated in ESCC.
E2F8 promotes ESCC cell proliferation and arrests the cell cycle in S phase

Two cell lines were selected based on their E2F8 expression patterns. E2F8 was overexpressed in EC-1 cells, which had the lowest expression level among the five ESCC cell lines studied, and E2F8 was downregulated in TE-10 cells, which had the highest expression level among the five ESCC cell lines studied. Using qRT-PCR and Western blotting, the transfection efficiencies were confirmed. The results indicated that E2F8 expression was successfully upregulated in EC-1 cells and downregulated in TE-10 cells (Figure 2A). To investigate the effects of E2F8 on proliferation, three assays, CCK-8, colony formation, and EdU assays, were conducted. CCK-8 assays showed that downregulating E2F8 expression decreased the proliferation ability of TE-10 cells, while upregulating E2F8 expression increased the proliferation ability of EC-1 cells (Figure 2B). Images of representative plates with colonies (Figure 2C) similarly showed that the downregulation of E2F8 expression in TE-10 cells decreased their colony formation ability compared to that of the control group. Upregulated E2F8 expression in EC-1 cells increased their colony formation ability compared to that of the NC group (Figure 2C). The results of EdU assays showed that the rate at which TE-10 cells incorporated EdU in the downregulated E2F8 group distinctly decreased compared to the rate in the control group; however, the opposite results were detected in the EC-1 cell lines when comparing the rate at which cells incorporated EdU in the upregulated E2F8 group to that of the NC group (Figure 2D). Cells incorporating EdU were considered to be proliferating cells. Flow cytometry was performed to analyze the effects of dysregulated E2F8 expression on the cell cycle. The results showed that downregulating E2F8 expression in TE-10 cells produced a significant increase in the percentage of cells in the G0/G1 phase, while upregulating E2F8 in EC-1 cells resulted in cell cycle arrest in S phase (Figure 2E). These findings demonstrated that E2F8 promoted ESCC cell proliferation and arrested the cell cycle at S phase.

E2F8 influences CCND1/p21

Previous studies have revealed that E2F8 influences the cell cycle by interacting with key factors. We wondered whether E2F8 could affect these factors in ESCC. qRT-PCR and Western blotting were performed to detect the expression patterns of CCND1 and p21 in the two pairs of genetically engineered cell lines. The results demonstrated that downregulating E2F8 expression decreased the expression...
Figure 2 (Continued)
E2F8 promotes tumorigenicity in vivo

To reveal the effects of E2F8 on tumorigenicity in vivo, TE-10 cells treated as previously described were injected into the flanks of nude mice to generate tumors ectopically.
The results showed that compared to the TE-10 control cells, the cells transfected with shE2F8 generated smaller and lighter tumors (Figure 4).

**Discussion**

Esophageal cancer is a malignant tumor with high morbidity and mortality due to recurrence, metastasis, and drug resistance. The main cause of these phenomena is the lack of knowledge about the disease mechanism; thus, it is imperative to reveal the potential mechanism of this disease. It may be much easier to diagnose and treat this kind of disease with identification of effective biomarkers or oncogenes. Among the various kinds of esophageal cancer, ESCC is the most common.\(^1\) Thus, in the present study, we focused on researching the mechanism and potential biomarkers of ESCC.

mRNA is a large class of RNA molecules that transfer genetic information from DNA to ribosomes; they specify the amino acid sequences of the protein products based on the expression of genes.\(^13\) Regarding cancer treatment, mRNAs have been widely investigated as biomarkers in multiple cancers. Zhang et al revealed that TRIM27 functions as an oncogene by activating epithelial–mesenchymal transition and p-AKT in colorectal cancer.\(^14\) Sun et al proved that TRIM59 facilitated the proliferation of colorectal cancer and promoted metastasis via the PI3K/AKT pathway.\(^15\)

The E2F transcription factor family is of paramount importance for coordinating cell cycle progression. Among the known E2F family members, eight members, E2F1–8, have been recognized. In general, E2F1–3 are considered as transcriptional activators, whereas E2F4–7 play inhibitory roles in the transcriptional expression of downstream target genes.\(^4\)–\(^6\) All these observations demonstrated that E2F transcription factors have significant effects on cell fate. Thus, it is urgent to identify the specific role for each E2F family member in specific malignancies. E2F1–7 were described in previous studies.\(^4\)–\(^6\) In the present study, we focused on E2F8, a new member of the E2F family. E2F8 has been previously researched and shown to regulate the cell cycle in several cancers. Zhang et al discovered that E2F8 expression is upregulated in colorectal cancer.\(^16\) Jin et al discovered that by suppressing E2F8, metformin can induce cell cycle arrest in the G1 phase in lung cancer cells.\(^17\) Sun et al found that E2F8 can promote papillary thyroid cancer progression by regulating the cell cycle.\(^11\) Ye et al indicated that upregulating E2F8...
expression modulates the G1/S phase transition and further promotes cell proliferation in breast cancer. However, the function of E2F8 still remains unknown in ESCC. In the present study, the expression pattern of E2F8 in ESCC was detected by qRT-PCR and Western blotting. The results showed that E2F8 expression was upregulated in ESCC tissue and cell lines. Transfected cell lines that upregulate or downregulate E2F8 expression were constructed. Using these cell lines, the effects of E2F8 on cell proliferation were investigated by CCK-8, colony formation, and EdU assays. Persistent proliferation is considered a hallmark of cancer.

Our studies demonstrated that E2F8 expression was upregulated in ESCC, and upregulated E2F8 expression promoted the proliferation of ESCC cells. In contrast, downregulated E2F8 expression presented the opposite effects by inhibiting ESCC cell proliferation. These findings indicated that E2F8 was dysregulated in ESCC and may play a significant role in ESCC by influencing cell proliferation. Cell proliferation is the main aspect that is controlled by the cell cycle. In general, the cell cycle is dysregulated in cancer cells, which destroys cell density homeostasis and leads to uncontrolled cell proliferation. We wondered whether E2F8 could affect the cell cycle. Using flow cytometry, upregulated E2F8 expression was discovered to arrest the cell cycle in S phase. Previous studies proved that the E2F8 gene, located on chromosome 11p15, encodes a family member of the transcription factors that regulate the genes required for progression through the cell cycle, and E2F8 could regulate progression from G1 to S phase by ensuring that the nucleus divides at the proper time.

Our results were consistent with previous findings, further indicating that E2F8 is an indispensable cell cycle regulatory factor in ESCC. We further detected the relationship between E2F8 and CCND1/p21, which were identified as key cell cycle factors. Through Western blotting, E2F8 was revealed to have the ability to influence p21, which can be a universal inhibitor of cyclin kinases and CCND1, which was identified as a key factor in regulating the cell cycle. It can be further deduced that E2F8 might be a transcriptional repressor of p21 and transcriptional activator of CCND1 in ESCC. CCND1 functions as an oncogene that is involved in many cancers by regulating the G1 to S phase transition of the cell cycle, while p21 acts as a master effector in multiple tumor suppressor pathways. Thus, E2F8 might play critical roles in regulating the cell cycle in ESCC by influencing CCND1/p21. However, the detailed mechanism of E2F8-mediated regulation of the cell cycle still needs further investigation. Taken together, the activities of E2F8, CCND1 and p21 could be activated or suppressed and the proliferative abilities of cells could be changed. Finally, E2F8 expression promoted ESCC cell proliferation in vivo.

In conclusion, our study identified E2F8 as an oncogene that promoted cell proliferation by CCND1/p21 in vitro and in vivo.

Author contributions

All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

References

