Assessment of insulin-like growth factor 1 receptor as an oncogene in esophageal squamous cell carcinoma and its potential implication in chemotherapy

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Abstract. Insulin-like growth factor-1 receptor (IGF-1R) is a tyrosine kinase receptor implicated in the pathogenesis of multiple cancers. After ligand binding, IGF-1R can initiate the activation of the PI3K/AKT/mTOR and Ras/Raf/MEK/MAPK pathways to modulate cell proliferation, survival, differentiation, motility, invasion and angiogenesis. IGF-1R is a prerequisite for tumor progression and is one of the most attractive targets for therapeutic interventions in several types of cancer. In the present study, we determined the expression of IGF-1R in an esophageal squamous cell carcinoma (ESCC) cohort, investigated the detailed function of IGF-1R and screened the potential application of IGF-1R in the clinic. We verified the higher expression of IGF-1R in ESCC tumor tissues as compared to adjacent normal tissues. We also found that high expression of IGF-1R was associated with advanced tumor progression. We used ESCC cell lines and a mouse xenograft model to detect the function of IGF-1R in vitro and in vivo. Our results suggest the oncogenic function of IGF-1R in regulating cell proliferation, clonogenesis, the cell cycle and apoptosis. In addition, we found that IGF-1R was associated with advanced tumor progression. We used ESCC cell lines and a mouse xenograft model to detect the function of IGF-1R in vitro and in vivo. Our results suggest the oncogenic function of IGF-1R in regulating cell proliferation, clonogenesis, the cell cycle and apoptosis. In addition, we found that IGF-1R was associated with the response to standard chemotherapy drugs 5-FU and cisplatin in an ESCC cell line. More importantly, we confirmed that the serum concentration of IGF-1/IGFBP3 can be used for predicting response to chemotherapy, and increased serum levels of IGF-1 and IGFBP-3 are associated with significantly higher rates of tumor response. In the present study, we demonstrated that IGF-1R is an important oncogene in ESCC and can be used to detect the chemotherapeutic response.

Introduction

Esophageal cancer is the eighth most common cancer worldwide with distinct geographical and ethnic characteristics (1). In China, it ranks as the fourth leading cause of cancer related mortality (2). Esophageal squamous cell carcinoma (ESCC) is the dominant type of esophageal malignancy in China. ESCC carcinogenesis is a multistage process characterized by morphological changes from normal esophagus to basal cell hyperplasia, dysplasia, carcinoma in situ and squamous cell carcinoma (SCC). It involves complex interactions between the environment and genetic factors (3). Although early intervention, as well as surgical, medical and radiotherapy techniques have undergone great achievements to date, the 5-year survival rate of ESCC patients remains low (4,5). Since most patients are at an advanced stage at initial diagnosis, the chance of surgery is lost (6). Chemotherapy-based comprehensive treatment is the main treatment for advanced patients. Yet, chemotherapeutic outcomes are not optimistic due to the poor selection and serious adverse reactions of standard chemotherapy drugs. Therefore, targeted therapy drugs that can avoid the chemotherapy-induced adverse reactions and elevate the treatment effect are critical to reduce the morbidity and mortality of ESCC.

Insulin-like growth factor-1 receptor (IGF-1R) is a tyrosine kinase receptor implicated in the pathogenesis of multiple cancers (7). Overexpression of IGF-1R has been reported in a range of human solid tumors such as breast, non-small cell lung and prostate cancer, sarcomas, hepatocellular carcinoma and pancreatic, ovarian and gastrointestinal cancers (8-11). After ligand binding, IGF-1R can initiate the activation of the PI3K/AKT/mTOR signaling and Ras/Raf/MEK/MAPK pathways resulting in the activation of multiple transcription factors such as ELK-1, CREB and AP-1 to modulate cell...
proliferation, survival, differentiation, motility, invasion and angiogenesis (12,13). Furthermore, more and more evidence indicates that IGF-1R is a prerequisite for tumor progression and is involved in the critical steps of the metastatic cascade (14,15). Therefore IGF-1R is one of the most attractive targets for cancer therapeutic interventions (16).

In addition to IGF-1R, the IGF signaling system consists of another two ligands, IGF-1 and IGF-II; another two cell membrane receptors, IGF-IIR and insulin receptor; and six IGF binding proteins (IGFBP 1-6) (17). High circulating levels of IGF-I and a decreased level of IGFBP3 have been linked to increased cancer risk and have been identified as prognostic markers for various cancers such as breast, prostate, colon and lung cancer (18). Almost 90% of the circulating IGFs bind to IGFBP3. IGFBP-3 inhibits IGF action at the cellular level by competitively binding IGFs and thereby preventing their binding to the IGF-1R. The molar ratio of IGF-I to IGFBP-3 has been regarded as an important index of IGF-I bioavailability. IGF-1R is associated with the sensitivity to chemotherapy in several types of cancers including lung, colon and head and neck cancer (19-21). The serum concentrations of IGF1 and IGFBP-3 may also play a role in the response to the chemotherapies.

IGF-1R has been reported as a marker for prognosis and as a therapeutic target in human ESCC (22). IGF-1 autocrine system could stimulate tumor growth and chemoresistance in human esophageal carcinoma cells (23). However, the detailed biological function of IGF-1R in ESCC still requires investigation, and the implications of IGF-1R in treatment are still unclear. In this study, we determined the expression of IGF-1R to IGF1 and IGFBP-3 may also play a role in the response to the chemotherapies.

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Materials and methods

Ethics statement. This study was approved by the Institutional Review Boards of the First Affiliated Hospital of Zhengzhou University. Written informed consent was obtained from all participants for the use of their blood or tissue samples. Full consent was obtained from the Institutional Review Boards of the First Affiliated hospital of Zhengzhou University. Animal care and experiments were approved by the Institute Animal Care and Use Committee of the Cancer Institute and Hospital, Chinese Academy of Medical Sciences and Peking Union Medical College.

Human ESCC cell lines and ESCC patient samples. ESCC cell lines EC9706, EC109 and NEC were gifts from the Department of Etiology and Carcinogenesis, Cancer Institute and Hospital, Chinese Academy of Medical Sciences and Peking Union Medical College. All the cells were cultured in RPMI-1640 medium with 10% fetal bovine serum (FBS), in 5% CO₂ at 37°C.

Eighty human ESCC tissues were fixed with 4% paraformaldehyde in PBS buffer and embedded in paraffin. The tissue samples were cut into 3-μm sections. The sections were incubated in 3% H₂O₂ for 15 minutes and washed three times with PBS buffer. The sections were stained with different antibodies, including IGF-1R, IGFBP-3, and other markers for the analysis of IGF-1R expression.

Bioinformatic analysis. The microarray data GSE 33426 and GSE 23400 were downloaded from the GEO database (http://www.ncbi.nlm.nih.gov/gds/). The differentially expressed genes were identified using GEO2R tools on GEO. The heatmap was generated using Genesis 1.0 (Graz, Austria). The biofunction and pathway analysis was performed with IPA® Software (http://www.ingenuity.com).

Immunohistochemistry. Fresh tumor tissues were fixed with formalin, embedded and finally cut into 4-μm sections. The sections were deparaffinized by dimethylbenzene and graded ethanol. Then the sections were incubated with 3% H₂O₂, goat serum, anti-IGF-1R antibody, biotin-labeled secondary antibody, peroxidase-conjugated streptavidin in order before DAB staining. All the reagents were purchased from ZSGB-Bio (Beijing, China). The staining intensity of IGF-1R was quantified into 3 groups according to the percentage of stained cells: negative, 0-5% cells were stained; weak positive, >5-50% cells were stained; strong positive, >50-100% cells were stained.

Western blot analysis. Protein was extracted from the cells using lysis buffer (50 mM Tris-HCl pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% Triton X-100). The protein concentration was measured using the Bradford method. The protein was separated by SDS-PAGE and transferred to a polyvinylidene fluoride (PVDF) membrane (Millipore, Billerica, MA, USA). After incubation with 10% milk, the primary IGF-1R antibody, and the secondary antibody, the blot signals were detected by chemiluminescence using the ECL kit (Applygen, Beijing, China). All the antibodies were purchased from Santa Cruz (Dallas, TX, USA).
Stable IGF-1R-knockdown cell line. pGCsilencer™ U6/Neo/GFP/RNAi-IGF-1R plasmid used for the knockdown of IGF-1R and the pGCsilencer™U6/Neo/GFP negative control were obtained from Genechem Co. (Shanghai, China). Both plasmids were transfected into EC9706 cells using Lipofectamine™ 2000 (Invitrogen, Carlsbad, CA, USA). The cells were digested and seeded into a new 24-well plate, diluted 5 times, 24 h after the transfection. Then, the regular medium was changed to medium containing 600 µg/ml G418 for stable transfection colony screening. After all of the cells without transfection died 4 days later, the remained cells were sustained by medium containing 200 µg/ml G418 and accepted as the stable transfection cell lines: IGF-1R-knockdown cell line and blank transfection control cell line. Western blotting and RT-PCR verified the knockdown of IGF-1R in the cell lines.

Cell proliferation study. Cells were seeded in a 96-well plate at a density of 4,000 cells/well. The next day, 10 µl of 5mg/ml MTT was added to the medium for a 3-h incubation, and at a density of 4,000 cells/well. The next day, 10 µl of 5mg/ml MTT was added to the medium for a 3-h incubation, and the time point was set as zero then detected every 24 h for 6 days. Then the medium was removed, and 150 µl DMSO was added. Finally the OD value was read at 490 nm on a Model 680 microplate reader (Bio-Rad, Hercules, CA, USA) after a 10-min incubation.

Flow cytometry. Cells were digested by trypsin, washed twice in cold PBS, fixed with ice cold 70% methanol, and incubated at 4°C overnight. Cells were then washed in PBS and incubated with 25 µg/ml propidium iodide containing 30 µg/ml ribonuclease for 30 min at room temperature. Cells were analyzed on the FACSCalibur Cell Sorting System (Becton Dickinson, Franklin Lakes, NJ, USA) using BD FACSuite™ Software in triplicate.

Cell colony formation study. Cells were seeded onto 6-well plates at a density of 300 cells/well. Following 12 days, visible colonies were fixed with methanol and stained with 0.4% crystal violet solution. Colonies were counted, and the number of visible colonies in each well was determined. Each assay was performed in triplicate.

Mouse ESCC xenograft study. Male BALB/c nude mice 5-6 weeks old were purchased from the Institute of Laboratory Animal Sciences, Chinese Academy of Medical Sciences and Peking Union Medical College. The mouse were divided into three groups and were inoculated with stable IGF-1R-knockdown cells, empty vector control cells or wild-type EC9706 cells, respectively (six mice in each group). A total of 2x10⁶ cells were inoculated subcutaneously into the right dorsal flank. The size of the tumor was measured twice per week using a digital vernier caliper. The tumor volume was determined from the orthogonal dimensions (d1, d2, d3) using the formula (d1 x d2 x d3) x π/6. Five weeks after inoculation, the mice were euthanized by cervical dislocation, and xenograft tumors were collected.

ELISA. Collected venous blood was centrifuged for 10 min at 2,000 rpm. The upper tube of serum was then taken for cryopreservation and stored at -40°C for ELISA assay. IGF-1 and IGFBF-3 ELISA kits were obtained from Senxiang Biotech (Shanghai, China). The standard curve was established according to the manufacturer's instructions. Serum samples (100 µl) were incubated in each well for 120 min, followed by 50 µl primary antibody buffer for 60 min, 100 µl secondary primary antibody buffer for 60 min, 100 µl substrate buffer for 10 min and 1 drop stop solution. Then the plate was read at 490 nm on the Model 680 microplate reader. The serum concentrations of IGF-1 and IGFBP3 were calculated by the standard curves.

Statistical analysis. All statistical analyses were performed using the SPSS 13.0 package. Pearson's χ² test was used to detect the expression difference in the cancer and normal tissues. All tests of statistical significance were two-sided, and P<0.05 was considered to indicate a statistically significant difference. All quantitative data are presented as means ± SD. Independent samples t-test and one-way ANOVA were adopted to compare the means.

Results

Bioinformatic analysis of IGF-1R in ESCC. In order to study the expression and potential function of IGF-1R in ESCC, we took advantage of publicly available gene expression data (GSE33426 and GSE23400). In the referenced studies, tumor cluster and normal cells separated by laser-capture microscope or tumor tissues and adjacent normal tissues were compared by cDNA microarray. IGF-1R was identified as a differentially expressed gene in both studies. Moreover, we found 85 and 50 genes, respectively, related with the IGF-1R signaling pathway (Fig. 1A and B). Twenty-three genes in the IGF-1R signaling pathway were identified in both studies (Fig. 1C and D). After IPA analysis, we found that these genes were mainly mapped to AKT and MEK cascade and were related to cell growth, proliferation and survival (Fig. 1E). The bioinformatic analysis supported our hypothesis that IGF-1R plays an important role in ESCC.

IGF-1R is highly expressed in human ESCC cancer tissues. Firstly, we used immunohistochemical (IHC) staining to detect the expression of IGF-1R in ESCC tumor tissues (Fig. 2A and C) and adjacent normal tissues (Fig. 2B and D). After IHC staining, IGF-1R showed strong staining in the cancer cell membrane. Of the 80 cancer specimens, 11 cases were negative, 28 cases were weakly positive and 41 cases were strongly positive. The total positive rate was 86.25% and the strong positive rate was 51.25%. Of the 18 normal tumor-adjacent tissues, 7 cases were negative, 9 cases were weakly positive and 41 cases were strongly positive. The total positive rate was 61.11% and the strong positive rate was 11.11%. Both the total positive rate and the strong positive rate were significantly higher in the ESCC tissues than in the normal tumor-adjacent esophageal tissues (total positive rate, χ²=4.630, P<0.05; strong positive rate, χ²=9.614, P<0.01). In addition, we found a relationship between the expression of IGF-1R and cancer patient characteristics (Table I). We found that high expression of IGF-1R was associated with more aggressive lymph node metastasis, lower histological grade and advanced clinical stage. Both positive and strong positive rates of IGF-1R expression were also higher in patients with
Figure 1. Bioinformatic analysis identifies the upregulation of the IGF-1R signaling pathway components in ESCC. (A) Heat map depicting 83 genes related to the IGF-1R pathway in GSE33426. (B) Heat map depicting 50 genes related to the IGF-1R pathway in GSE23400. (C) Venn diagram showing the common genes identified in GSE33426 and GSE23400. (D) Gene list for the 27 common genes identified in GSE33426 and GSE23400. (E) The core molecular events of the IGF-1R signaling pathway in ESCC.

Figure 2. Expression of IGF-1R in ESCC and normal tissues. Tissues in A-D were from the same patients. (A and C) Staining in tumor tissues. (B and D) Staining in adjacent normal tissues.
lymph node metastasis than those without lymph node metastasis ($P<0.01$), and were also higher in patients with advanced clinical stage (III-IV stage) than those with early clinical stage (I-II stage). The expression of IGF-1R was also increased from a low differentiation grade to a moderate differentiation stage and high differentiation grade ($P<0.05$ between each of the two groups). No significant difference was noted in relation to age or gender ($P>0.05$). These results showed that IGF-1R was highly expressed in ESCC tissues, and the high expression of IGF-1R in the ESCC tissues was related to tumor progression.

**IGF-1R acts as an oncogene in ESCC in vitro and in vivo.** We determined the expression of IGF-1R in EC9706, EC109 and NEC cells and found that IGF-1R was relatively highly expressed in these three ESCC cell lines. We established stable IGF-1R-knockdown cells by transfection of pGCsilencer™U6/Neo/GFP/RNAi IGF-1R in EC9706 cells. Western blotting and RT-PCR verified the successful knockdown of IGF-1R in these cells (Fig. 3A and B). Compared with the empty vector control, the cell proliferation was inhibited and the doubling time was extended significantly in the IGF-1R-knockdown cells (Fig. 3C). Significant cell cycle arrest occurred in the IGF-1R-knockdown cells with an increased percentage of G1 phase cells ($P<0.05$) and a decreased percentage of S phase cells ($P<0.01$) (Fig. 3D). In addition, a higher apoptosis rate ($P<0.05$) and weaker clonogenic ability ($P<0.05$) were found after IGF-1R-knockdown (Fig. 3E). All of these results showed that IGF-1R acts as an oncogene in ESCC in vitro. We also used a mouse ESCC xenograft model to detect the tumor growth ability of IGF-1R in vivo. Ten days after cell inoculation, subcutaneous nodules were noted and finally developed into tumors. Although all three cell lines induced visible tumors in the nude mouse, the size of the tumors induced by the stable IGF-1R-knockdown cells was significantly smaller than those formed by the empty control ($P<0.05$) and wild-type cells ($P<0.05$) (Fig. 4A and B). The animal assay showed that IGF-1R can induce tumor growth in vivo.

**High expression of IGF-1R is related to low sensitivity to chemotherapy in the ESCC cell lines.** We compared the sensitivity to 5-FU and cisplatin between the IGF-1R-knockdown cells and the empty vector control. We found that the growth inhibition rate of 5-FU and cisplatin in the pGCsilencer™U6-IGF-1R-transfected cells was higher than that in the empty vector-transfected and the non-transfected cells ($P<0.05$) (Fig. 5A). The high expression of IGF-1R facilitates cancer cell resistance to chemotherapy.

**Serum concentrations of IGF-1/IGFBP-3 are associated with the chemotherapy response.** We determined the serum concentrations of IGF-1 and IGFBP-3 in 172 patients both before and after chemotherapy treatment (Fig. 5B). The mean serum concentrations of IGF-1 pre-therapy and post-therapy in all patients were 268.87±61.66 and 266.42±49.98 ng/ml, respectively, without significant difference. The mean serum concentrations of IGFBP-3 pre-therapy and post-therapy in all patients were 2523.2±469.83 and 2598.8±563.56 ng/ml, respectively, without significant difference. In contrast, the serum concentration of IGFBP-3 increased ($P<0.01$) and the IGF-1/IGFBP-3 ratio decreased ($P=0.01$) significantly after chemotherapy in the chemotherapy responsive group (CR and PR). The serum concentration of IGF-1 decreased after chemotherapy in the responsive groups but without significance ($P>0.05$). Accordingly, in the chemotherapy unresponsive group (SD and PD) the IGF-1/IGFBP-3 ratio increased.

### Table I. Association of IGF-1R with clinical characteristics.

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<th>Clinical characteristics</th>
<th>Negative n (%)</th>
<th>Weak positive n (%)</th>
<th>Strong positive n (%)</th>
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<th>P-value</th>
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<td>8 (10.7)</td>
<td>19 (32.2)</td>
<td>25 (57.2)</td>
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<td>3 (15.4)</td>
<td>9 (36.5)</td>
<td>16 (48.1)</td>
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<td>&lt;51</td>
<td>1 (7.7)</td>
<td>5 (38.5)</td>
<td>7 (53.8)</td>
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<td>51-65</td>
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<td>18 (38.3)</td>
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<tr>
<td>&gt;65</td>
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<td>5 (25.0)</td>
<td>10 (50.0)</td>
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<td>Lymph node metastasis</td>
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<td>3 (5.6)</td>
<td>11 (20.3)</td>
<td>40 (74.1)</td>
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<td>17 (65.3)</td>
<td>1 (3.9)</td>
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<td>High</td>
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<td>I-II</td>
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Figure 3. Biological function of IGF-1R in the ESCC cell lines. (A) IGF-1R protein expression in three ESCC cell lines. (B) Expression of IGF-1R in the EC9706 cells after pGCsilencer™U6/Neo/GFP/RNAi-IGF-1R transfection (siRNA) or without any treatment (Control) or pGCsilencer™U6/Neo/GFP empty plasmid (NC). (C) Growth curve of cells after different treatments. (D) Cell cycle and apoptosis analysis for cells after different treatment. (E) Colony formation abilities of cells after different treatments. *P<0.05, **P<0.01.

Figure 4. IGF-1R induces tumor growth in mouse xenografts. (A) The volume of tumors in the different groups. (B) The size of tumors in the different groups. *P<0.05
significantly (P<0.05) after chemotherapy; the concentration of IGFBP-3 decreased but without significance (P>0.05). There was no variation in IGF-1 (Table II). These results showed that the IGF-1/IGFBP-3 ratio can be used to predict the chemotherapeutic effect; a decreased IGF-1/IGFBP-3 molar ratio presents a better chemotherapeutic effect. We also evaluated the chemotherapy response rate in 4 groups divided according to the quartiles of plasma IGF-1 and IGFBP-3. Chemotherapy response rate increased from low concentrations of IGF-1 and IGFBP-3 to high concentrations of IGF-1 and IGFBP-3. These results showed that increased serum levels of IGF-1 and IGFBP-3 are associated with significantly higher rates of tumor response (P<0.05).

Discussion

In the present study, we determined the expression of IGF-1R in an ESCC cohort and verified the high expression of IGF-1R in ESCC tumor tissues when compared with that in adjacent normal tissues. High expression of IGF-1R was found to be associated with aggressive lymph node metastasis, lower histological grade and advanced clinical stage. After knockdown of IGF-1R in ESCC cell lines and the mouse xenograft assay, we verified the pro-proliferation, apoptosis inhibition and tumor growth function of IGF-1R in vitro and in vivo. We also found that IGF-1R was associated with the response to standard chemotherapy drugs 5-FU and cisplatin in the ESCC cell lines. More importantly, we found that the serum concentration of IGF-1/IGFBP3 can be used for the predicting chemotherapeutic effect. Increased serum levels of IGF-1 and IGFBP-3 were found to be associated with significantly higher rates of tumor response.

High expression of IGF-1R has been reported in several cancers including ESCC. Imsumran et al used IHC to detect the expression of IGF-1R in ESCC tumor tissues and found that ~50% of tissues showed high expression of IGF-1R which
was associated with invasive depth, metastasis, advanced tumor stage and recurrence (22). In this study, we also detected high expression of IGF-1R in ESCC tissues compared with that in adjacent normal tissues. We found that high expression of IGF-1R was associated with more aggressive lymph node metastasis, lower histological grade and advanced clinical stage, similar to the results of previous research. We suggest that the expression of IGF-1R can be used as a biomarker for the diagnosis and tumor progression of ESCC.

IGF-1R plays an important role in carcinogenesis mainly based on the activation of the PI3K/Akt/mTOR signaling and the Ras/Raf/MEK/MAPK pathways. The activation of AKT and MAPK pathways help cancer cells acquire the ability for proliferation, evasion of apoptosis, insensitivity to anti-growth signals, unlimited replicative potential, metastasis and angiogenesis (24,25). Both of these pathways are reported to be activated in ESCC. After using siRNA to knock down the expression of IGF-1R, we found significant inhibition of cell growth, cell cycle arrest, reduced apoptosis and fewer colonies. These results are the same as previous findings. Various strategies such as anti-IGF-1R antibodies, IGF-1 mimetic peptides, antisense strategies, IGF-1R-specific peptide aptamers, targeted degradation of IGF-1R and expression of dominant-negative IGF-1R mutants have been explored to inhibit IGF-1R signaling (26). AVE1642, a humanized version of the murine monoclonal antibody can bind specifically and with high affinity to human IGF-1R preventing IGF-1R binding to its ligand resulting in receptor inactivation (27). It has been reported to delay the growth of tumor xenografts and to prolong the survival of tumor-bearing nude mice. An IGF-1R inhibitor NVP-AEW541 showed significant inhibition of pancreatic cell lines, and is being used for an in vivo study (28). Another small molecular IGF-1R inhibitor BMS-554417 showed antitumor activity in breast cancer in vitro and in vivo (29). The small molecular inhibitor for IGF-1R and IGF-1R antibodies both can be used for ESCC treatment (30). In further research, we will utilize the IGF-1R antibody or a small molecular inhibitor for IGF-1R to treat cells or animals to verify new strategies for the targeted therapy for IGF-1R.

Chemotherapy resistance is a serious obstacle to cancer therapy. Any cellular and molecular events associated with chemotherapeutic effects are possible targets for better outcomes from treatment. We found that high expression of IGF-1R facilitated ESCC cell resistance to chemotherapy. Liu et al reported that IGF-1 prevented the apoptosis in ESCC CE81T/VG cells induced by chemotherapeutic drugs, such as cisplatin, 5-FU and camptothecin (23). Thus, interruption of IGF-1R function may provide a strategy by which to retard tumor growth and increase the sensitivity of esophageal carcinoma to chemotherapy. Due to the important bioactivities of the ligand for IGF-1R, IGF1, we hypothesized that chemotherapy may influence the serum levels of IGF-1 and IGFBP3. In order to prove this hypothesis, we compared the serum concentrations of IGF-1 and IGFBP3 in ESCC patients before and after chemotherapy. Our results showed that the IGFBP3 ratio was significantly decrease after chemotherapy in the chemotherapy responsive group (CR and PR) and was increased in the chemotherapy unresponsive group (SD and PD). The IGF-1/IGFBP3 ratio can be used to predict the chemotherapeutic effect. The serum concentration of IGFBP3 was increased in the chemotherapy responsive group and had a tendency to decrease in the chemotherapy unresponsive group. We divided patients into 4 groups according to quartiles of serum IGFBP3 and then compared the chemotherapy response rate in these 4 groups. We found that the chemotherapy response rate increased from low concentrations of IGF-1 and IGFBP3 to high concentrations of IGF-1 and IGFBP3. These results showed that increasing serum IGF-1 and IGFBP3 are associated with significantly higher rates of tumor response (P<0.05). Since almost 90% of circulating IGFs bind with IGFBP3, we believe that the concentration of IGF-1/IGFBP3 is more effective for predicting the chemotherapeutic effect. We believe that the potential function of IGF-1/IGFBP3 to predict chemotherapeutic effect is also related to the function of IGF-1R.

In conclusion, in the present study we verified the oncogenic function of IGF-1R in vitro and in vivo. We proved that a high serum concentration of IGF-1R is associated with chemotherapy resistance. This study offers strong evidence for the application of IGF-1R as a new target for ESCC therapy.

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References