Overexpression of *SEZ6L2* predicts poor prognosis in patients with cholangiocarcinoma

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**Background:** With the feature of destructive and biliary malignancy, intrahepatic cholangiocarcinoma (ICC), presents unclear molecular mechanisms which contributes to typically poor prognosis for patients. *Seizure-related 6 homolog-like 2* (*SEZ6L2*) is a gene that encodes for a seizure-associated protein localized on the cell surface. Thus far, the function of *SEZ6L2* in ICC has not been reported.

**Methods:** We used data from The Cancer Genome Atlas and the Gene Expression Omnibus to analyze dynamics behind and levels of expression of *SEZ6L2* in ICC. Then we used qRT-PCR and Immunohistochemical staining to detect levels of expression of *SEZ6L2* and thereby determined the potential clinical significance of this protein in ICC.

**Results:** According to qRT-PCR and immunohistochemical analysis results, *SEZ6L2* was overexpressed in ICC. Kaplan-Meier and Cox proportional hazard analyses indicated that patients afflicted by ICC with high levels of relative expression of *SEZ6L2* have a poorer prognosis and that *SEZ6L2* may be an independent prognostic factor which enables to the accurate prediction of overall survival (OS) and disease-free survivals’ (DFS) expected rates. Subcutaneous xenograft models used to explore the role of *SEZ6L2* in tumor formation in vivo. The dynamics of the *SEZ6L2* gene being promote angiogenesis in cholangiocarcinoma are related to increasing expressive growth factors which include *EGF*, *VEGF*, *PDGF* and the activation of the *P38-MAPK* pathway.

**Conclusions:** Our findings suggest that *SEZ6L2* can serve as an advanced biomarker that can be used to accurately predict a patient prognosis and be used as a target for ICC treatment.

**Keywords:** Intrahepatic cholangiocarcinoma (ICC); seizure-related 6 homolog-like 2 (*SEZ6L2*); prognostic factor; *P38-MAPK* pathway

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**Introduction**

Intrahepatic cholangiocarcinoma (ICC) is an aggressive malignancy with poor prognosis for afflicted patients. The incidence of ICC has increased over recent years, ICC accounts for 15–20% of all primary hepatobiliary malignancies (1-3). According to some studies, several proteins, e.g., carbohydrate antigen (CA) 19-9, can serve as biomarkers that can be used in the process of accurate prognosis prediction (4,5). Thus, further research to elucidate mechanisms related to the ICC progression is still
needed, which can contribute to the development of new therapeutic strategies for afflicted patients. We have been screening for genes encoding up-regulated/down-regulated molecules in cholangiocarcinoma using microarrays derived from tumor cells purified by laser capture microdissection. To verify the clinicopathological and biological significance of the respective gene products, we performed tissue microarray analysis of clinical cholangiocarcinoma specimens. This systematic approach combines bioinformatics tools to search for cell surface and/or secreted proteins and finds that SEZ6L2 is frequently transcribed in a large number of patients with cholangiocarcinoma.

Seizure-related 6 homolog like 2 (SEZ6L2) is a gene that encodes a seizure-associated protein localized on the cell surface, which plays a role when it comes to the regulation of a variety of biological functions. SEZ6L2 is a type 1 membrane protein containing the C1r/C1s, Uegf, and Bmp1 (CUB) complement domains as well as complement control protein (CCP) domains (6). Proteins which contain these domains help to regulate the localization or function of neurotransmitter receptors (7,8), and serve as a helper subunit for GluR (9). Research indicated that the overexpression of SEZ6L2 in lung cancer makes it a potential prognostic marker for this type of tumor (10), however the dynamics and function of SEZ6L2 in cholangiocarcinoma still remains unclear.

In our study, we sought to investigate the levels of expression of SEZ6L2 in ICC by assessing several ICC specimen types with paired para-cancerous tissues. The relationships between the level of expression of SEZ6L2 and clinical pathological features, overall survival (OS), and disease-free survival (DFS) of patients with ICC were also analyzed. We present the following article in accordance with the ARRIVE reporting checklist (available at http://dx.doi.org/10.21037/tcr-20-1527).

Methods

Patients and tissue samples

We obtained all samples from patients with ICC at the First Affiliated Hospital of Sun Yat-Sen University (Guangzhou, China) between January, 2004 and December, 2015. All samples were from patients who underwent radical surgical resectioning without preoperative chemotherapy or radiotherapy. All patients were followed up with until at least through December 2017, and all initial and ongoing diagnoses were confirmed by using standard methodologies for pathological analysis. Complete clinical pathological and follow-up data for our study are available for all 101 ICC samples we collected. It conformed to the provisions of the Declaration of Helsinki (as revised in 2013). The study was approved by Institutional Research Ethics Committee of the First Affiliated Hospital of Sun Yat-sen University (2019-P-220) and informed consent was taken from all the patients. All fresh tumor tissue specimens were rapidly frozen in liquid nitrogen immediately after resection and then stored at −80℃.

Cell culture

Human cholangiocarcinoma cell line RBE was purchased from the cell bank of Shanghai Academy of Chinese Sciences. The frozen cells were removed from liquid nitrogen before the experiment and thawed rapidly in a constant temperature water bath at 37℃. During the experiment, the cells were shaken continuously to accelerate the thawing. Cell suspension was quickly transferred to a centrifugal tube containing 10 mL RPMI1640 medium containing 10% FBS in a super-clean workbench for mixing.

Cells were transferred to culture flask with a transpirator, and the algebra and date of cells were marked on the flask. The cells were placed in a 37℃, 5% CO2 incubator for further culture.

Quantitative real-time polymerase chain reaction (qRT-PCR)

The fresh frozen tissue was lysed in TRIzol solution (Invitrogen, Carlsbad, CA, USA) to obtain purified RNA. 1 mg of total RNA was transcribed reversely into cDNA with Verso cDNA Kit (Thermo Fisher Scientific). The relative gene expression level was monitored via qPCR by use of SYBR Premix Ex Taq II (Takara, Japan) and Terpal Cycler Dice TP850 (Takara) as per the standard methods provided by supplier. The 2-ΔΔCT method was applied for determination of target gene expression in experimental group and control group. qRT-PCR was conducted in triplicate for each sample, and detected gene expression was normalized to that of GAPDH.

Plasmid, small interfering RNA and transfection

The coding region of SEZ6L2 gene was cloned into
PUC57 vector to construct the overexpression vector pBPLV-SEZ6L2. Three SEZ6L2 interfering plasmids were constructed, and the sequence of interfering plasmids was as follows.

shRNA1, CCGGCCAGCCCAGTTCTGTGTAATAC TCGAGTTCAATATCTCAACTTGAGCTTTTTG; shRNA2, CCGGGCTCCAAGTTGAGATATTGAAC TCGAGTTCAATATCTCAACTTGAGCTTTTTG; shRNA3, CCGGCCGTAATAGTTGAGATATTGAAC TCGAGTTCAATATCTCAACTTGAGCTTTTTG. Then the interfering vector was transfected into RBE cells. After 96 hours, RNA was extracted from RBE cells for qPCR verification, and a plasmid with the highest efficiency was screened for subsequent experiments.

Western blot analysis

Western blot was used to detect the expression of SEZ6L2 protein and the expression of angiogenesis related factors CD36, EGF, PDGF, VEGF and VEGFR-1 after gene interference. The cells were collected and centrifuged, and the corresponding lysate was added. The cells were lysed at 4 °C for 30 min and centrifuged at 10,000 rpm for 10 min. The supernatant was carefully absorbed to obtain the total protein. Protein concentration was determined by BCA kit. After boiling, sampling and electrophoresis for 1 to 2 hours, the protein was transferred to PVDF membrane. The membrane was incubated overnight with a solution of the first antibody at 4 °C, and then the second antibody was incubated at room temperature for 1 to 2 hours. ECL exposure solution was dripped onto the membrane and exposed. Quantity one software was used to analyze the gray value of each antibody band.

Immunohistochemical staining

The de-paraffin-rehydrated sections were heated in sodium citrate buffer (pH 6.0) for 10 min for antigen retrieval. After inactivation of endogenous peroxidase by 3% hydrogen peroxide, the antibody against SEZ6L2 was incubated for 1 h at 37 °C. An IHC kit (Dako) was used to analyze the protein expression level as standard manual. Two more experienced pathologists scored the staining results. In this study, we subjected a total of 101 pairs of formalin-fixed paraffin-embedded ICC tissues to immunohistochemical staining.

Animal studies

Human cholangiocarcinoma cell line RBE (0.2 mL PBS containing 5×10^5 cells) transfected with indicated plasmids was subcutaneously inoculated into the back of 5-week-old female Balb/c nude mice (8 mice in each group). Tumor diameter was measured every 5 days in 20 days. The tumor volume (mm^3) was calculated based on the longest and shortest diameter. The formula was as follows: volume = (shortest diameter)2 × (longest diameter) ×0.5. After 30 days of inoculation, the mice were killed and the xenografts were dissected for further analysis. Approval for the animal experiments was obtained from Sun Yat-sen University Animal Care and Use Committee, Institute of Clinical and Translational Research, Sun Yat-sen University (2019-P-220). And the study was conducted in compliance with guidelines for the care and use of Laboratory Animals (Ministry of Health, People's Republic of China, 1998).

Bioinformatic analysis

Clinical information and data for levels of expression related to ICC were obtained from The Cancer Genome Atlas (TCGA; http://gdc.cancer.gov/) and from the Gene Expression Omnibus (GEO). We used R and GraphPad Prism 7 software to apply a log-transformation to the data and analyze the results. Negative binomial distributions, empirically based Bayesian estimations, exact tests, generalized linear models (GLMs) and quasi-likelihood tests in the edgeR and limma packages were also used in this study to assess important patterns in the data. We used the metric of a calculated logFC (fold-change) ≥1.0 or log FC ≤−1.0, and we defined an associated P value of <0.05 as a statistically significant gene.

Statistical analysis

We used SPSS version 23.0 (IBM) for statistical analysis. We used a Student’s t-test to compare differences between treatment groups and explore the dynamics of the relationships between categorical variables with the χ² test. The relative impact of various prognostic factors on the OS and DFS rates of patients with ICC was assessed using the Cox regression and Kaplan-Meier analysis. Two-tailed P values <0.05 were used to indicate statistically significant results at which the null hypothesis of no differences between treatment groups would be rejected.
Results

The overexpression of SEZ6L2 in comparison with para-cancerous tissues

We examined SEZ6L2 expression using mRNA sequencing or microarray datasets from TCGA and GEO to investigate levels of expression of SEZ6L2 in patients with ICC. Based on the results from analysis of TCGA, we compared 36 samples of ICC afflicted tissues with 9 samples of normal tissues, and the comparative results indicated that levels of expression of SEZ6L2 were significantly higher in tumor afflicted tissues than were respective levels in para-cancerous tissues (Figure 1A). In addition, the results from analysis of GSE26566 (P<0.0001) and GSE32225 (P=0.0155) indicated that the level of expression of SEZ6L2 was significantly elevated in ICC afflicted tissue sample compared with para-cancerous tissues (Figure 1B,C). Moreover, we used qRT-PCR and 12 pairs of fresh clinical samples which verified upregulation of the levels of expression of SEZ6L2 in tumor tissues (Figure 1D). The results of immunostaining analyses reflect that levels of expression of SEZ6L2 were high in 51 of the 101 total ICC cases in our study (Figure 2A; Table 1). Taken together, these results indicated that SEZ6L2 is overexpressed in ICC afflicted tissues comparing with the respective levels of expression in paired samples of para-cancerous tissues.

Correlations between SEZ6L2 expression and clinical parameters in ICC

We next analyzed the immunostaining results and corresponding clinical data for the 101 pairs of ICC afflicted and non-diseased tissues to assess relationships between the levels of expression of SEZ6L2 and standardized ICC clinicopathological parameters. These clinical features are presented in Table 1. Results indicated that the level of expression of SEZ6L2 was strongly positively related to tumor size (P=0.002), lymphatic metastasis (P=0.016) and
TNM stage (P=0.018). However, there we observed no significant difference with regard to the level of expression of \textit{SEZ6L2} and patient measures of sex, age, liver cirrhosis, or perineural and vascular invasion.

\textbf{SEZ6L2 is an independent prognostic marker in ICC patients}

To determine whether \textit{SEZ6L2} would be a good candidate to be used as an independent prognostic factor for treatment of patients with ICC, we utilized Cox regression analysis was utilized to examine OS and DFS. Univariate analysis indicated the stage of TNM, the levels of expression of \textit{SEZ6L2}, and the degree of lymphatic metastasis was associated with OS. Furthermore, results from multivariate analysis indicated that the stage of TNM, medical complications, and levels of expression of \textit{SEZ6L2} were independent prognostic factors for OS (Table 2). As shown in Table 3, univariate analysis indicated that the levels of expression of \textit{SEZ6L2} and the stage of TNM are related to DFS. Multivariate analysis also indicated that the level of expression of \textit{SEZ6L2} was an independent prognostic factor for patients with ICC (Table 3). Taken together, the results of this aspect of our study indicate that \textit{SEZ6L2} can serve as an independent prognostic factor for patients with ICC.

\textbf{Poor survival prediction of strong SEZ6L2 expression in patients with ICC}

We next evaluated the prognostic value of the levels of expression of \textit{SEZ6L2} in patients with ICC using the Kaplan-Meier analysis. The data of 101 patients including survival data were sampled and analyzed, the results indicated that patients with a high level of expression of \textit{SEZ6L2} had correspondingly lower rates of OS and DFS than those with a low level of expression of \textit{SEZ6L2} (P=0.005 and P=0.024 respectively; Figure 2B,C). Thus, a high level
of expression of SEZ6L2 appeared to successfully predict poor survival rates in patients with ICC.

**SEZ6L2 promoting ICC tumor growth in vivo**

The expression of GFP green fluorescent protein was the strongest when lentivirus SEZ6L2 was transfected into cholangiocarcinoma cell line RBE and assessed under inverted fluorescence microscopy. The efficiency of transfection of lentivirus SEZ6L2 knockdown vector was observed to be 75% (Figure 3A). Eighteen female nude mice aged 5–9 weeks were divided into an experimental group (transfected sh-SEZ6L2) and a negative control group (no-load lentivirus, N=8 mice each group). Results indicated that there were no significant differences in growth rates of transplanted tumors between the two treatment groups within 5 days after transplantations (P>0.05). After 10 days of transplantation, the growth rate of transplanted tumors in the experimental group was slower and smaller than those in the negative control group (P<0.05, Figure 3B,C,D). Immunohistochemical results showed positive expression of SEZ6L2 and vascular endothelial growth factor (VEGF) as observed through the appearance of obvious yellow and brown granules in the cytoplasm and as some cell membranes were stained (data not shown). Accordingly, the levels of expression of SEZ6L2 and CD34 in the experimental group were significantly lower than those respective levels in the negative control groups (Figure 3E,F).

**SEZ6L2 expression promoting angiogenesis of cholangiocarcinoma**

The results from qRT-PCR analysis indicated that the levels of expression of CD36, EGF, P38, PDGF, SEZ6L2, VEGF, and VEGFR-1 in the expression vector group were significantly higher than those respective levels in the control group and the empty vector treatment group (P<0.01, Figure 4A). In contrast, the effect of the interference vector group was opposite, and the levels of expression of related genes were inhibited successfully and found to be significantly different among the treatment groups (P<0.01, Figure 4B). Results from western blotting showed that the levels of expression of CD36, EGF, P38, PDGF, SEZ6L2, VEGF, and VEGFR-1 in the expression vector group increased significantly comparing with the control and the empty vector groups (P<0.01, Figure 4C). The interference vector group had the opposite effect, which successfully inhibited expression of CD36, EGF, P38, PDGF, SEZ6L2, VEGF, and VEGFR-1. The scientific significance (P<0.01, Figure 4D) was consistent with the results of qRT-PCR.

**Discussion**

Typically poor prognosis for patients with ICC is characterized with the feature of a median survival rate for patients who do not undergo surgery within 6 months, and characterized by a 5-year survival rate for patients following complete resection of only 20% to 40% (11-13). According
to previous research, the poor prognosis for patients with ICC is largely resultant due to the high frequency of recurrence and metastasis after surgical resection, as well as the resistance to systemic courses of chemotherapy-based treatments (14). Therefore, the identification of heretofore mostly unknown potential biomarkers that could be used for aiding early diagnosis of cases of ICC is urgently needed. By referring to the literature, it was reported that in non-small cell lung cancers (NSCLC), patients whose tumours revealed a higher level of SEZ6L2 expression suffered shorter tumour-specific survival compared to those with no SEZ6L2 expression (15). In addition, SEZ6L2 was significantly up-regulated in tumour tissues of patients with colorectal cancer compared with adjacent normal tissues. Up-regulation of SEZ6L2 was correlated with a poor prognosis in patients with colorectal cancer. Meanwhile, SEZ6L2 overexpression was closely associated with poor OS and DFS in hepatocellular carcinoma patients. Moreover, SEZ6L2 is an independent prognostic predictor for the survival of HCC patients (16). Until now, the clinical significance of SEZ6L2 in ICC remains uncertain and unclear. However, in our study, it is found that the levels of expression of SEZ6L2 were higher in ICC afflicted tissues comparing with the ANL T s samples. Taking all these

| Table 2 Cox proportional-hazard regression analysis for overall survival |
|---------------------------------------------------------------|
| **Variables** | **Univariate analysis** | **Multivariate analysis** |
| | HR | 95% CI | P value | HR | 95% CI | P value |
| Sex | 1.124 | 0.651–1.941 | 0.675 |
| Age [60] | 0.850 | 0.490–1.475 | 0.563 |
| Differentiation | 0.919 | 0.364–2.319 | 0.858 |
| Lymphatic metastasis | 1.956 | 1.124–3.435 | 0.018* |
| TNM stage | 2.271 | 1.310–3.938 | 0.003* |
| HBsAg | 1.008 | 0.564–1.803 | 0.978 |
| Liver cirrhosis | 0.917 | 0.469–1.793 | 0.799 |
| Size | 1.429 | 0.824–2.480 | 0.204 |
| SEZ6L2 | 2.176 | 1.242–3.814 | 0.007* |

Lymphatic metastasis, TNM stage, SEZ6L2 are included in the Multivariate analysis. *, significant difference.

| Table 3 Cox proportional-hazard regression analysis for disease-free survival |
|---------------------------------------------------------------|
| **Variables** | **Univariate analysis** | **Multivariate analysis** |
| | HR | 95% CI | P value | HR | 95% CI | P value |
| Sex | 1.056 | 0.613–1.921 | 0.843 |
| Age [60] | 0.765 | 0.440–1.330 | 0.343 |
| Differentiation | 0.840 | 0.333–2.122 | 0.713 |
| Lymphatic metastasis | 1.689 | 0.971–2.937 | 0.063* |
| TNM stage | 1.988 | 1.151–3.438 | 0.014* |
| HBsAg | 0.855 | 0.478–1.530 | 0.598 |
| Liver cirrhosis | 0.943 | 0.483–1.839 | 0.862 |
| Size | 1.545 | 0.890–2.680 | 0.122 |
| SEZ6L2 | 1.856 | 1.066–3.232 | 0.029* |

Lymphatic metastasis, TNM stage, SEZ6L2 are included in the multivariate analysis. *, significant difference.
results into consideration, survival data analysis indicated that SEZ6L2 overexpression plays a significant role in the corresponding observation of poor survival rates of patients with ICC. These findings lead to the conclusion that SEZ6L2 is highly expressed in ICC afflicted tissues and can serve as a potential biomarker for promoting outcomes and accurate prognoses for patients with ICC.

SEZ6L2 is a brain-specific receptor-like protein that is predominantly expressed in the soma and dendrites of neurons, which required for synaptic maturation (6). A previous study indicated that the promotion and regulation of SEZ6L2 methylation contributes to cell proliferation and metastasis through the regulation of dynamics in the pathways for STAT3, EGR1, and PAX4, and thus played an important role in the dynamics of pathogenesis of osteosarcoma (17). Both DNA methylation and histone modification are epigenetic modifications that can control gene expression and promote cell proliferation (18), cell differentiation, and cell adaptation to environmental conditions. Dysregulation of these processes may lead to various pathologies (19). So far, isoforms functions of SEZ6L2 have not been clarified in great detail. Therefore, further studies are still needed to confirm the results of our studies and hypothesize hypothesis related to the use of SEZ6L2 to improve the prognosis of patients with ICC.

SEZ6L2 encodes a 92.5-kDa protein with an N-terminal signal peptide, five SUSHI domains, three CUB domains, and a C-terminal transmembrane domain (20,21). These encoded domains and products are likely mainly involved in developmental processes, cell to cell interactions, and dynamics related to cell adhesion (22,23). It is known that cell to cell interaction and cell adhesion are strongly related to proliferation and metastasis, and may play additional potentially critical roles in tumor development and malignant transformations. In particular, the leading cause of death among cancer patients is metastasis (24).

Figure 3 Knockdown of SEZ6L2 impaired cholangiocarcinoma carcinogenesis in vivo. The expression of GFP green fluorescent protein and assessed under inverted fluorescence microscopy after lentivirus was transfected into cholangiocarcinoma cell line RBE, the efficiency of transfection of lentivirus SEZ6L2 knockdown vector was observed to >80% (A). The nude mice were sacrificed 28 days after the injection and tumors from respective groups were shown (B,C). Tumor growth curves after the injection of RBE cells. Tumor volume was calculated every 3 days (D). Relative expression was measured in IHC staining of SEZ6L2 in tumors with different treatments (E). Relative expression was measured in IHC staining of CD34 in tumors with different treatments.
Additionally, cell migration and the ability of cancer cells to disseminate from the primary site forming distant metastases, are the primary causes of cancer-related morbidity in patients with solid tumors (25). Until now, the impact of SEZ6L2 on ICC and its mechanisms were unreported in the literature, and although our novel results are interesting, further investigations will still be required. However, this study clearly demonstrated that SEZ6L2 has potential for clinical use as a reliable biomarker for prognosis and investigations of ICC.

In our study, functional and immunohistochemical analysis of candidate molecules on tissue microarray has revealed that SEZ6L2 is likely to be a potential target for development of novel tools for diagnosis of ICC. In addition, in vitro and in vivo results suggested that SEZ6L2 functioned as an oncogene in cholangiocarcinoma patients. The knockdown of SEZ6L2 impairs tumour growth by promoting angiogenesis, which indicates that SEZ6L2 could be a potential therapy target for ICC.

By designing small molecule inhibitors or gene therapy targeting SEZ6L2 (knockout of oncogene SEZ6L2), the therapeutic purpose of inhibiting tumorigenesis and
development can be achieved. In addition, the application of tumor-targeting nanocarrier system to encapsulate chemotherapeutic drugs and gene-related agents (such as SEZ6L2 siRNA) for co-delivery of drug agents to tumor cell targets to achieve synergistic cancer suppression by chemotherapy and gene therapy is also a highly desirable approach for cancer treatment.

Cytokines including fibroblast growth factor-2, vascular endothelial growth factor (VEGF), EGF, and TGF-beta can regulate the levels of expression of SEZ6L2 (26,27). VEGF is one of the strongest VEGFs. The combination of VEGF and its receptor, VEGFR, specifically stimulates mitosis of vascular endothelial cells and promotes angiogenesis. Furthermore, as a powerful tumor lymph angiogenesis factor, VEGF promotes proliferation of lymphatic system tumors, as well as induces changes in the levels of expression in tumor tissues and tumors. It has also been determined that there is a close relationship between the degree of vascularization and biological behaviors including the rates and degrees of infiltration and metastasis (28). PDGF is a receptor tyrosine kinase, which plays an important role in the growth, proliferation, and differentiation of cells including vascular endothelial cells by binding with receptors. PDGF also promotes angiogenesis of tumors, thus promoting the growth, invasion, and metastasis of tumors.

Therefore, we used qPCR and western blot to detect the changes of the expression levels of EGF, P38, PDGF, SEZ6L2, VEGF, and VEGF receptor-in from determinations of protein and mRNA levels after SEZ6L2 interference and induced overexpression. The analytical results indicated that the levels of expression of VEGF, VEGFR, and PDGF were significantly increased after the SEZ6L2 gene was applied with induced interference and overexpression. In some cancer cells, there is a positive feedback mechanism of regulation between TGF-beta and SEZ6L2. These results imply that SEZ6L2 may promote the expression of VEGF, VEGF receptor, and PDGF through such a positive feedback regulation, thus ultimately promoting the angiogenesis of cholangiocarcinoma.

The results of our study lead to the conclusion that SEZ6L2 overexpression can be used as a metric to accurately predict a poor prognosis for patients with ICC. Furthermore, observations indicate that SEZ6L2 can serve as an independent prognostic factor for predicting outcomes related to OS and DFS in patients with ICC. Nonetheless, further work is needed to elucidate the molecular mechanisms through how SEZ6L2 promotes the progression of ICC.

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Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). Approval and consent obtained for the use of human tissue were obtained from the Institutional Research Ethics Committee of the First Affiliated Hospital of Sun Yat-sen University (2019-P-220). Informed consent was taken from all the patients. Approval for the animal experiments was obtained from Sun Yat-sen University Animal Care and Use Committee, Institute of Clinical and Translational Research, Sun Yat-sen University (2019-P-220). The study was conducted in compliance with guidelines for the care and use of Laboratory Animals (Ministry of Health, People’s Republic of China, 1998).

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