Genipin Induces Autophagy and Suppresses Cell Growth of Oral Squamous Cell Carcinoma via PI3K/AKT/MTOR Pathway

Background: Oral squamous cell carcinoma (OSCC) is a common malignant tumor of the head and neck, and it accounts for more than 90% of oral cancer. Due to high mortality, limitations of traditional treatment and many complications, new treatment methods are urgently needed. This study aimed to look into the effect of new potential anti-tumor drug, genipin, on OSCC treatment.

Methods: In vitro, CCK-8, colony formation, and flow cytometry were used to detect the effect of genipin on SCC-9 and SCC-15 cell lines. Immunofluorescence, real-time PCR, and Western blotting were used to investigate its mechanism. Xenograft tumor model was used to explore the role of genipin in vivo.

Results: We found that genipin suppressed cell growth and induced apoptosis in vitro. In addition, the expression of p62 was down-regulated while Beclin1 and LC3II were up-regulated in SCC-25 and SCC-9 cells. 3-methyladenine (3-MA) significantly decreased LC3 (LC3II)+ puncta, but genipin rescued this reduction. Furthermore, genipin also reduced the expression of p-PI3K, p-AKT, and p-mTOR. In vivo experiment showed that genipin significantly curbed the tumor size and weight. The positive expression of Ki67 protein and number of apoptotic cells were increased.

Conclusion: Conclusively, this study implicated that genipin suppresses cell proliferation and stimulated apoptosis, and is the first exploration showing that genipin induces OSCC cell autophagy via PI3K/AKT/mTOR pathway inhibition.

Keywords: genipin, oral squamous cell carcinoma, autophagy, PI3K/AKT/mTOR pathway

Introduction

Oral cancer is an increasing threat worldwide, with around 640,000 new cases being detected annually globally.1,2 In addition, there are 145,000 deaths per year in oral cancer,3 and five-year survival rates of patients with advanced oral cancer are 30–40%.4 Owing to the lack of early symptoms and appropriate screening for precancerous lesions, most oral cancer cases are not detected until later stages. The incidence of oral cancer is mainly associated with the consumption of areca nut, tobacco, and alcohol.5 Notably, oral squamous cell carcinomas (OSCC) account for more than 90% of oral cancer and is more likely to occur in men, the risk increases with age.6 Tongue is the most common site of OSCC, which has little effect on other parts of the mouth. So far, the mechanism of oral squamous cell carcinoma has not been fully elucidated, which seriously restricts the prognosis and treatment strategy.
Genipin, an aglycone of geniposide, is a major component of the fruit of *Gardenia jasminoides,* and it is an iridoid glycoside hydrolyzed from intestinal microbiota. Genipin has been used in the pharmacological treatment of cardiovascular disorders, such as xenograft scaffolds of heart valves. Moreover, Genipin is also considered to be a therapeutic drug for liver fibrosis. Previous studies show that Genipin has a wide range of anticancer activities in different types of human cancer. Parallelly, several reports have suggested that Genipin can inhibit cell invasion and migration, and induce apoptosis of colon cancer cells. Previous studies show that Genipin has a wide range of anticancer activities in different types of human cancer. Parallelly, several reports have suggested that Genipin can inhibit cell invasion and migration, and induce apoptosis of colon cancer cells. However, the protective effect of Genipin on oral carcinoma is still ambiguous.

PI3K/AKT/mTOR signaling pathway plays hub roles in cell proliferation, angiogenesis, transcription, and translation. Substantially, PI3K is a phosphoinositide kinase that catalyzes PIP2 phosphorylation when activated by G protein conjugated and tyrosine kinase receptors. Consequently, phosphorylated PIP2 then further phosphorylates AKT and stimulate downstream mTOR, eventually activating protein translation and promote cell growth or proliferation. Besides, abnormal PI3K/AKT/mTOR pathway causes many human cancers, including human oral cancer. Therefore, PI3K/AKT/mTOR pathway may be a crucial target against cancer.

In this study, we explored the effect of Genipin on OSCC and tried to provide theoretical basis for drug treatment of cancer.

**Materials and Methods**

**Cell Culture and Genipin Treatment**

Normal human oral epithelial cells NHOK and human tongue squamous cancer cells SCC-25 and SCC-9 were obtained from Procell Life Science&Technology Co., Ltd (Wuhan, China), and were cultured in DMEM (Thermo Fisher Scientific, Waltham, USA) medium and supplemented with 10% fetal bovine serum (FBS, Gibco, Shanghai, China), penicillin (100 U/mL), and streptomycin (100 μg/mL) at 37°C with 5% CO₂. Then, cells were treated with Genipin (King Tiger, Chengdu, China), which was dissolved in 0.1% dimethyl sulfoxide (DMSO, Sigma-Aldrich, Shanghai, China). Cells treated with DMSO alone were set as control.

**Cell Viability Assay**

NHOK, SCC-25, and SCC-9 cells suspension were seeded in 96-well plate (100 μL/well), respectively. Then, the plates were placed in the incubator (37°C, 5% CO₂) for pre-culture. Next, 10 μL of CCK-8 solution (Beyotime, Shanghai, China) was added into each well according to the manufacturer’s instructions, and incubated for 48 hrs with different concentrations of Genipin (0 to 400 μM). The absorbance at 450 nm was detected by a microplate reader (Bio-Rad, Hercules, CA, USA) and the changes of cell viability were estimated.

**Clone Formation Assay**

Firstly, about 3 × 10³ SCC-25 and SCC-9 cells were treated by Genipin with three different concentrations (10, 30, 60 μM) for 72 hrs, respectively. Next, cells were seeded into 35 mm dishes and cultured for 14 days. After that, two types of cells were rinsed with PBS and fixed in 4% paraformaldehyde (Sigma, USA) for 15 mins at room temperature, respectively. Then, they were incubated with Wright-Giemsa dye for 5 mins with Wright-Giemsa stain (NIJCBIO, Nanjing, China) for 5 mins. Finally, the images of clone formation were captured and number of cell clones were calculated.

**Flow Cytometry**

Briefly, SCC-25 and SCC-9 cells were fixed for 24 hrs, respectively. Next, the fixed cells were stained with an Annexin V-FITC Apoptosis Detection Kit (Beyotime, Shanghai, China) according to the manufacturer’s instructions. After gently mixing, cells were incubated at room temperature for 20 mins in dark. Finally, red fluorescence was detected by flow cytometry (Thermo Fisher Scientific, Waltham, USA) at 488nm wavelength.

**Real-Time PCR**

Total RNA of SCC-25 and SCC-9 cells was isolated using TRizol reagent (Invitrogen, Waltham, USA) and cDNA was synthesized by a PrimeScript RT Reagent kit (TaKaRa, Kusatsu, Japan), according to manufacturer’s instructions. Next, Real-time PCR was performed using a SYBR Green Master mix (TaKaRa, Kusatsu, Japan), including 1 μL primers (10 μmol/L) and 1μL cDNA (100 ng/μL). β-actin was used as endogenous control and relative mRNA levels normalized by β-actin were calculated employing 2^-ΔΔCt_ mean. The primers sequences are listed below: Ki67-F: 5’-GCAGGACTTCAC TTGCTTCC-3’, Ki67-R: 5’-TCAATTTGCGTTTGTTTCACG -3’; Survivin-F: 5’-GACCACCGATCTACACCT-3’, Survivin-R: 5’-TCTTCCACCTGCTTCTTGA$T$-3’; β-actin-F: 5’-CTGTCTGGCCGACCACCAT-3’, β-actin-R: 5’-GCAACT AAGTCATAGTCCGC –3’.
Western Blotting
Total protein was extracted from SCC-25 and SCC-9 cells using Radioimmunoprecipitation (RIPA) Assay Buffer (Thermo Fisher Scientific, Waltham, USA). And the protein lysates were separated on SDS-PAGE gel and blotted onto polyvinylidene difluoride (PVDF, Millipore, Boston, USA) membranes. Next, the primary antibody was incubated after blocking in 5% non-fat milk. All antibodies were obtained from the Abcam (Cambridge, UK) and shown below: Ki67 (ab16667, 1:1000), survivin (ab76424, 1:1000), cleaved PARP1 (ab32561, 1:1000), cleaved caspase-3 (ab2302, 1:1000), P62 (ab109012, 1:1000), Beclin1 (ab207612, 1:1000), LC3I (ab192890, 1:1000), LC3II (ab52628, 1:1000), PI3K (ab70912, 1:1000), p-PI3K (ab18265, 1:1000), AKT (ab179463, 1:1000), p-AKT (ab38449, 1:1000), mTOR (ab2732, 1:1000), p-mTOR (ab109268, 1:1000), and β-actin (ab179467, 1:1000). Finally, the strips were exposed with ECL reagent (Sangon Biotech, Shanghai, China) for 2 min at room temperature. Afterwards, the ECL solution was added and incubated in darkness at 37°C for 60 mins. The protein bands were detected by a One-Step TUNEL Apoptosis Assay Kit (Beyotime, Shanghai, China) and shown below: Ki67 (ab179463, 1:1000), p-mTOR (ab109268, 1:1000), P62 (ab109012, 1:1000), Beclin1 (ab207612, 1:1000), LC3I (ab192890, 1:1000), LC3II (ab52628, 1:1000), PI3K (ab70912, 1:1000), p-PI3K (ab18265, 1:1000), AKT (ab179463, 1:1000), p-AKT (ab38449, 1:1000), mTOR (ab2732, 1:1000), p-mTOR (ab109268, 1:1000), and β-actin (ab179467, 1:1000). The paraffin sections were sliced to 4 µm thick. According to the manufacturer’s instructions, the positive expression of Ki67 was detected using a DAB Substrate kit (ZSGB-BIO Co., Ltd., Beijing, China) and the number of apoptosis was detected by a One-Step TUNEL Assay Kit (Beyotime, Shanghai, China). Antibodies were obtained from the Abcam (Cambridge, UK).

Immunofluorescence Staining
SCC-25 and SCC-9 cells were grown on glass coverslip and cultured with or without Genipin (10, 30, and 60 µM), respectively. After washing twice in PBS, the cells were fixed with 4% paraformaldehyde for 20 mins. Then, cells were permeabilized for 10 mins in PBS containing 0.1% Triton X-100 (Sigma-Aldrich, Shanghai, China). Next, they were blocked in 4% goat serum blocking solution for 30 mins at room temperature. Afterwards, the cells were incubated with LC3 antibody at 4°C overnight. On the following day, the fluorescein-labeled secondary antibody was added and incubated in darkness at 37°C for 60 mins. Finally, the images were observed and photographed under a fluorescence microscope (Carl Zeiss, Jena, Germany).

Animal Experiments
Male BALB/c nude mice (4–6 weeks old, 20 ± 2 g) were obtained from the Animal Center of the Wuhan University Zhongnan Hospital (Wuhan, China). Mice were housed 6 per cage at 25°C and 80% humidity in a 12 hrs light/dark cycle for free access to food and water. OSCC xenograft model was established in mice by subcutaneous injection of SCC-25. When the tumor volume reached 100 mm³, 30 mg/kg Genipin was injected intraperitoneal for a week. Meanwhile, the control (n=6) was injected with 0.9% NaCl. At 30 days, the tumors were weighed and volume were measured. All animal experiments and operations were implemented in accordance with the guidelines of the National Institute of Laboratory Animal Care and Use (8th edition, National Academy of Sciences Press, 2011), and approved by the ethics committee of Wuhan University Zhongnan Hospital.

Histopathological Analysis
Histopathological assays were performed using fresh mouse tissue. In brief, after the mice were sacrificed, tumor and normal tissues were fixed in formalin and paraffin embedded. The paraffin sections were sliced to 4 µm thick. According to the manufacturer’s instructions, the positive expression of Ki67 was detected using a DAB Substrate kit (ZSGB-BIO Co., Ltd., Beijing, China) and the number of apoptosis was detected by a One-Step TUNEL Assay Kit (Beyotime, Shanghai, China). Antibodies were obtained from the Abcam (Cambridge, UK).

Statistical Analysis
Data are presented as the mean ± standard deviation (SD), and IBM SPSS Statistics 25.0 was used for statistical analysis (IBM, Armonk, USA). One-way ANOVA analysis of variance was conducted to analyze the significance of differences. Multiple comparisons using one-way analysis of variance and Newman–Keuls test. P<0.05 was considered to indicate a statistically significant difference.

Results
Genipin Inhibited Viability of Oral Carcinoma Cells
Base on the toxic effect of Genipin in colon cancer, CCK8 assay was conducted to measure the cell activity using a concentration gradient of Genipin (0 to 400 µM). As shown in Figure 1B, we found that Genipin inhibited SCC-25 and SCC-9 cells in a concentration-dependent manner, but when the concentration was below 100 µM, there was no significant effect on normal oral epithelial cells (NHOK). In addition, when the concentration were ≥60 µM, Genipin obviously suppressed the viability of SCC-25 and SCC-9 cells. Given this, we selected three concentrations of 10, 30, and 60 µM for subsequent in vitro experiment.

Genipin Suppresses OSCC Cell Growth
According to the clone formation experiment, Genipin concentration-dependent inhibited the growth of OSCC cells (Figure 2A). When Genipin exceeded 30 µM, the cell colony
formation rate was significantly reduced compared with the control group. In addition, we also detected the expression of ki67 (proliferation maker) and Survivin (apoptosis maker) by qPCR and Western blotting, respectively. As shown in Figure 2B, the mRNA and protein levels of Ki67 and Survivin were decreased by the treatment of Genipin in SCC-25. Parallelly, the result in SCC-9 cells was similar to SCC-25 (Figure 2B). Hence, the above data suggested that Genipin suppressed OSCC cell proliferation.

Genipin Induced OSCC Cell Apoptosis
In order to further explored the effect of Genipin on cell proliferation, we detected cell apoptosis using flow cytometry. We found that the apoptosis rate of SCC-25 and SCC-9 cells significantly increased after Genipin treatment (10, 30, and 60 μM), compared with control (Figure 3A and B). Moreover, the marked protein (caspase-3 and PARP) of apoptosis were detected by Western blotting, and the data showed that the protein levels of cleaved-caspase-3 and cleaved-PARP were obviously elevated after Genipin treatment (Figure 3C and D). These results confirmed that Genipin promoted cell apoptosis in OSCC.

Genipin Promoted OSCC Cell Autophagy
In order to study the effect of Genipin on autophagy, we detected the related proteins expression of autophagy in SCC-25 and SCC-9 cells through Western blotting. The results showed that Genipin significantly down-regulated the expression of p62 and upregulate the expression of Beclin 1 and LC3II/LC3I in a concentration-dependent manner (Figure 4A and B). In addition, immunofluorescence assay showed that intracellular LC3 (LC3II)+ puncta increased significantly after Genipin treatment (Figure 4C). Besides, autophagy process was further confirmed by autophagy inhibitor 3-methyladenine (3-MA, 5 mM), we found that 3-MA significantly inhibited LC3+ puncta, and Genipin weakened this trend after combined treatment with Genipin and 3-MA (Figure 4D).

PI3K/AKT/mTOR Pathway Was Involved in the Anticancer Effect of Genipin on OSCC Cell
Since the abnormal regulation of PI3K/AKT/mTOR signaling pathway is very common in human cancer, we have explored whether the Genipin regulated this pathway in human OSCC. As shown in Figure 5A and C, the phosphorylation levels of PI3K, AKT, and mTOR were down-regulation after treated with Genipin. Then, we added PI3K activator 740Y-P (20 μM). As shown in Figure 5B and D, 740Y-P increased the expression of p-PI3K, p-AKT, and p-mTOR, as well as the protein levels of cleaved-caspase-3 and LC3II. But, after co-treatment with Genipin and 740Y-P, Genipin weakened the activation of the pathway by 740Y-P.

Genipin Alleviated OSCC Tumor in vivo
Combined with the results of in vitro experiments, we established a SCC-25 xenograft tumor model. The data showed that compared with the control group, the tumor weight and the tumor volume in the treatment group were significantly reduced (Figure 6A–C). Immunohistochemistry assay showed that the expression of Ki67 was down-regulation (Figure 6D), and TUNEL assay manifested that the number of apoptosis was increased in tumor tissue (Figure 6D).
addition, we also found that the protein levels of cleaved-caspase-3 and LC3II were increased in the Genipin treatment group, compared to the control (Figure 6E).

Discussion
At present, OSCC remains a major threat to increased morbidity and poor 5-year survival. Although OSCC has advantages of timely treatment due to relatively easier access for early diagnosis, it is also an aggressive disease with local recurrence and cervical lymph node metastasis. The basic method of therapy includes surgical resection leaving a margin of healthy tissue, and neck dissection during node metastases. However, conventional therapy often leads to impaired oral functions. Moreover, radiotherapy often leads to several of complications such as xerostomia, radiation caries, taste disorders, and mucositis. This study provided insight into the
protective function of Genipin for OSCC, our findings showed that Genipin inhibited the growth of OSCC cells, induced apoptosis, and improved the autophagy level, which provided a new strategy for the oral cancer treatment.

Genipin is derived from geniposide and is the major component of Gardenia jasminoides Ellis fruit. Li et al.26 implicated that Genipin inhibited the cell growth in human bladder cancer. Jiang et al.27 claimed that Genipin induced apoptosis in OSCC cells. Figure 3 shows the effect of Genipin on cell apoptosis in OSCC. SCC-25 and SCC-9 cells were treated with three concentrations of Genipin (10, 30, and 60 μM). (A and B) SCC-25 and SCC-9 cells apoptosis were measured by flow cytometry. (C) The protein levels of cleaved-caspase-3 and Cleaved-PARP in SCC-25 were detected Western blotting. (D) The protein levels of cleaved-caspase-3 and Cleaved-PARP in SCC-9 were detected Western blotting. *p<0.05, **p<0.01 compared with the control group. Data are presented as mean ± SD, and all the experimental in triplicate.
HO-1 expression/activity and subsequently decreased vascular smooth muscle cell (VSMC) proliferation and migration. In our study, we found that Genipin inhibited the clone formation of SCC-25 and SCC-9 cells, and the expression of Ki67 and Survivin were down-regulation. Our animal experiments further confirmed the inhibitory effect of Genipin on cell autophagy in OSCC. SCC-25 and SCC-9 cells were treated with three concentrations of Genipin (10, 30, and 60 μM). (A) The autophagy relevant protein (p62, Beclin1, LC3I/II) levels in SCC-25 were detected by Western blotting. (B) The autophagy relevant protein (p62, Beclin1, LC3I/II) levels in SCC-9 were detected by Western blotting. (C) LC3 (LC3II) puncta in SCC-25 and SCC-9 cells were measured by immunofluorescence assay, respectively. (D) After added the autophagy inhibitor 3-MA, LC3 (LC3II) puncta in SCC-25 and SCC-9 cells were measured by immunofluorescence assay, respectively. *p<0.05, **p<0.01. Data are presented as mean ± SD, and all the experimental in triplicate.
Genipin on tumor growth, the data showed that Genipin (30 mg/kg) treatment straightly reduced the size and weight of xenograft tumors, accompanied by a decrease in the expression of Ki67. Besides, the previous study demonstrated that Genipin treatment in human gastric cancer cell line induced apoptosis in a dose-dependent manner via p53-independent Egr1/p21 signaling pathway. In addition, the evidence also suggested that Genipin existed the anti-tumor activity of inducing apoptosis and inhibiting invasion in breast cancer.

Exhilaratingly, in this study, the results showed that Genipin promoted the apoptosis of SCC-25 and SCC-9 cells in vitro and significantly raised the protein level of cleaved-caspase-3 and Cleaved-PARP. Similarly, we also found that Genipin promoted the expression of cleaved-caspase-3 in xenograft mouse model. Therefore, in combination with previous studies, we notarized that Genipin induced apoptosis in OSCC.

In cell biology, autophagy is a catabolic process of its own components by a lysosomal machine. Autophagy acts a key role in cell survival, which is a key pathway for homeostasis, development, and other pathophysiological processes. Moreover, autophagy showed an increase in the expression of autophagy-related proteins such as LC3-II, Beclin-1, and ATG5, with a decrease in the expression of p62. Kinarivala et al reported that the activated Beclin-1 could induce autophagy. In this study, we investigated the effect of Genipin on autophagy in OSCC for the first time. In vitro, we found that Genipin treatment up-regulated the protein levels of Beclin1 and LC3II, while down-regulated the protein level of P62. After co-incubation with autophagy inhibitor 3-MA, the autophagy process was alleviated, compared with Genipin alone. Ulteriorly, animal experiments confirmed that Genipin induced an increase in the expression of LC3II protein.

PI3K/AKT/mTOR signaling pathway has been widely known to be involved in the progression and tumorigenesis of many types of cancers including Oral Squamous cancer. There is sufficient evidence that PI3K/Akt/mTOR axis played an important role in the occurrence of oral cancer. For instance, Rizzo et al found that PI3K mutation was associated with the occurrence of oral squamous cell carcinoma. Prodromidis et al found that the up-regulation of Akt and

Figure 5 Regulating effect of Genipin on PI3K/AKT/mTOR pathway in OSCC. SCC-25 and SCC-9 cells were treated with three concentrations of Genipin (10, 30, and 60 μM). (A) The expression of PI3K, p-PI3K, AKT, p-AKT, mTOR, and p-mTOR in SCC-25 were measured by Western blotting. (B) After added the PI3K activator 740Y-P, the expression of PI3K, p-PI3K, AKT, p-AKT, mTOR, p-mTOR, cleaved-caspase-3, and LC3II in SCC-25 were measured by Western blotting. (C) The expression of PI3K, p-PI3K, AKT, p-AKT, mTOR, and p-mTOR in SCC-9 were measured by Western blotting. (D) After added the PI3K activator 740Y-P, the expression of PI3K, p-PI3K, AKT, p-AKT, mTOR, p-mTOR, cleaved-caspase-3, and LC3II in SCC-25 were measured by Western blotting. *p<0.05, **p<0.01. Data are presented as mean ± SD, and all the experimental in triplicate.
mTOR expression in OSCC was more common than in oral lichen planus (OLP), which was thought to be the result of PI3K activation. Here, we found that Genipin inhibited the phosphorylation of PI3K, AKT, and mTOR in a concentration-dependent manner, indicating that the PI3K/Akt/mTOR signaling pathway was inactivated. In order to further verify the regulatory effect of Genipin in PI3K/Akt/mTOR, we added PI3K activator 740Y-P. As we predicted, 740Y-P increased the expression of p-PI3K, p-AKT, and p-mTOR, as well as the protein levels of cleaved-caspase-3 and LC3II. However, after co-treatment with Genipin and 740Y-P, Genipin weakened the activation of the pathway by 740Y-P. Therefore, it is believed that Genipin inhibits OSCC cells growth and improves the autophagy through suppressing PI3K/Akt/mTOR signaling pathway, and PI3K/Akt/mTOR pathway is an attractive target for the treatment of OSCC.39

Conclusively, this study showed that Genipin inhibited the cells' viability, induced apoptosis, and promoted autophagy in OSCC, and its mechanism might be related to the inhibition of PI3K/Akt/mTOR signaling pathway. It provided a new candidate for OSCC drug treatment. Although we found that OSCC inhibited tumor growth in animal models, its protective mechanism in vivo needed further study.

Disclosure
The authors report no conflicts of interest in this work.

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