Knockdown of LRP5 Promotes Proliferation and Invasion of Tongue Squamous Cell Carcinoma through Compensatory Activation of Akt Signaling

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Received: 2023.12.25; Accepted: 2024.04.03; Published: 2024.04.15

Abstract

The role of LRP5, a critical receptor in the Wnt signaling pathway, remains unexplored in tongue squamous cell carcinoma (TSCC). This study investigates the impact of LRP5 knockdown on the biological behaviors of TSCC cell lines both in vitro and in vivo. Our findings indicate that LRP5 knockdown significantly enhances cell proliferation, migration, and invasion in CAL27 and SCC25 cell lines. RNA-seq analysis reveals compensatory activation of the Akt pathway, with 119 genes significantly upregulated post-LRP5 knockdown. MMP1 expression suggests its potential involvement in TSCC progression. Western blot analysis demonstrates increased Akt phosphorylation, upregulated proliferation-related PCNA, and downregulated apoptosis-related caspase-3 after LRP5 knockdown. Down-regulation of E-cadherin and β-Catenin, proteins associated with cell adhesion and invasion, further elucidates the molecular mechanism underlying increased cell migration and invasion. Our study concludes that compensatory Akt pathway activation is essential for the LRP5 knockdown-induced migration and proliferation of CAL27 and SCC25 cells. These results highlight LRP5 as a potential therapeutic target for TSCC. Simultaneous inhibition of Wnt and Akt signaling emerges as a promising approach for TSCC treatment.

Keywords: LRP5; Akt signaling pathway; Tongue squamous cell carcinoma; Invasion; Wnt pathway

Introduction

Tongue squamous cell carcinoma is one of the most common cancers in the neck and head [1], which often results in short survival and poor prognosis, even after surgery and chemotherapy. Cell surface molecules can be used as good targets for anti-tumor drugs, such as epidermal growth factor receptor (EGFR), PD-L1, and insulin-like growth factor 1, all of which have been successfully used in the clinical treatment of tumors. It is much harder for resistant cells to develop resistance against surface receptor molecules than drug molecules. LRP5 (low-density lipoprotein receptor-related protein 5) is a transmembrane protein, as a co-receptor of the Wnt signaling pathway. With the co-receptor’s LRP5/6, the family of Frizzled (FZD) receptors interacts with WNT ligands to activate the canonical WNT/β-catenin signaling cascade. Deregulation of this pathway is involved in the progression, drug resistance, and immunity escape of several cancers [2–4]. Therefore, LRP5 may be an ideal cell surface molecular target to regulate the Wnt signaling pathway in the treatment of head and neck squamous cell carcinoma.

The WNT signaling pathway is crucial in both head and neck cancer, including TSCC, where its dysregulation contributes to tumor initiation,
progression, and metastasis. Specifically in TSCC, abnormal activation of WNT signaling fosters tumor cell proliferation, invasion, and epithelial-mesenchymal transition (EMT), resulting in aggressive tumor behavior and unfavorable prognosis [5,6]. The interaction between Wnt and Frizzled (Fz) activates the noncanonical pathway, while the activation of the canonical pathway requires the involvement of Wnt co-receptors LRP5/6. The canonical Wnt1 gene was initially discovered as a preferred integration site for the mouse mammary tumor virus in virally induced breast tumors. Subsequently, LRP5/6 were found to play a crucial role in Wnt1-dependent tumor development in transgenic mouse models [7-9]. In preliminary experiments, we found that elevated expression of LRP5 in TSCC may be associated with patient prognosis. Thus, components of the canonical pathway, including LRP5/6, are commonly implicated in cancer progression as oncogenes. LRP5/6's ectodomain comprises four β-propeller/epidermal growth factor (EGF) repeats (E1-4) and three LDL repeats (LDLR), with E1-4 serving as the binding domain for canonical Wnt ligands and the canonical pathway inhibitor Dkk1[10-13]. However, the proteins binding to LDLR remain unexplored. Activation of the canonical pathway requires the close proximity of LRP5/6 and Fz, facilitated by canonical Wnt ligand binding to E1-4 of LRP5/6 and the amino-terminal cysteine-rich domain (CRD) of Fz[14,15]. Conversely, Dkk1 induces LRP5/6 internalization via its receptor Kremen, preventing Wnt reception and inhibiting the canonical pathway. Key components like LRP5 act as co-receptors, transmitting WNT signals and facilitating oncogenic processes across various cancers [16-18]. Understanding the complex role of WNT signaling in head and neck cancer highlights its significance as a potential therapeutic target against this formidable disease.

For the role of LRP5 in tumor cells, there are two types of research results in the past. The first type is to knockdown the expression of LRP5 through siRNA or shRNA lentivirus, and it is found that knockdown the expression of LRP5 can inhibit the growth of tumor cells [19]. Or by overexpressing the dominant-negative, soluble LRP5 (sLRP5) can reverse epithelial-mesenchymal transition of tumor cells by blocking Wnt signaling [20-22]. Activation of LRP5 gene promotes CSCs-like phenotypes, including tumorigenicity and drug resistance in colorectal cancer cells [23]. Interestingly, some studies have shown that LRP5 knockdown does not slow the proliferation and invasion of certain cells, and the mechanism is not very clear. Ren [24] has shown that in breast cancer, knockdown of LRP5 or LRP6 suppresses the Wnt signaling, but it increases lung metastasis in breast cancer cells in nude mice. Maria and her colleagues [25] find that LRP5 overexpression inhibits cell proliferation and induces apoptosis in HL60 cells. Some scholars [26] chose to examine the effects of LRP5 mRNA expression on patient survival in six different malignancies, including colorectal adenocarcinoma, and failed to demonstrate a significant effect. In this study, we wanted to study the effects of LRP5 knockdown on the biological behavior of tongue cancer cell lines and their possible mechanisms.

According to previous studies, the activation of canonical Wnt/β-catenin signaling promotes invasion, proliferation, and anti-apoptosis in TSCC cells [27-30]. Targeted inhibition of the Wnt signaling pathway is considered promising in the treatment of TSCC and even head and neck cancer. Certain inhibitors of the β-catenin complex or neutralizing antibodies to FZD/LRP receptors have been gradually applied in some clinical trials. ICG-001 and the second-generation compound PRI-724 that selectively inhibit the CBP/β-catenin complex are used in clinical trials in patients with certain solid tumors [31]. Vantictumab [32], an anti-Frizzled-1/2/5/7/8 antibody, is also being used in clinical trials for certain solid tumors.

Our study found that after LRP5 knockdown in different tongue cancer cell lines, the growth of tumor cells was accelerated. Transcriptome sequencing found that after CAL27 and SCC25 were transfected with siLRP5 lentivirus, the Akt signaling pathway was abnormally activated. Compensatory activation of the Akt signaling pathway resulted in faster cell growth. We speculate that on the basis of inhibiting the Wnt signaling pathway, combined with inhibiting the activation of the Akt signaling pathway, the dual inhibition may limit the growth and progression of tongue cancer cell lines in vitro and in vivo.

Materials and methods

Antibodies and reagents

The LRP5 (#5731), p-Akt (#4060), pan-Akt (#4685), and E-cadherin (#14472) antibodies are all manufactured by Cell Signaling Technologies (Beverly, MA, USA). The p-mTOR (AP0115), mTOR (A2445), β-Actin (AC026), and PCNA (A12427) were purchased from Abclonal (China). The β-Catenin (51067-2-AP) and HRP-conjugated GAPDH (HRP-60004) are from Proteintech. The cleaved caspase-3 (ab32351) antibody is bought from Abcam (Cambridge, UK). The Akt inhibitors included GSK690693, AZD5363 and LY294002 and crystal

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violet are all manufactured by Beyotime (China). The alamarBlue is from thermofisher (Cleveland, OH, USA).

**Cell Culture**

The CAL27 (CVCL_1107) and SCC25 (CVCL_1682) cells were cultured in high glucose Dulbecco's modified Eagle's medium (DMEM; Gibco, Invitrogen) with 10% fetal bovine serum (FBS; GEMINI) and 1% penicillin–streptomycin. Cells were purchased from Shanghai Zishi Biological Company and identified by STR. All experiments were performed using mycoplasma-free cells. Cells were incubated in a humidified atmosphere at 37°C with 95% air and 5% CO2. The siLRP5 and siNC lentiviruses were synthesized by Gcbio (Shanghai, China) and stably transfected after puromycin (1ug/ml) was added for 7 days. The alamerBlue assay procedure was conducted. First, cells were seeded into a 96-well plate at 1000 cells per well. After allowing the cells to attach and proliferate for a specified time, 10 μl alamerBlue was added to each well. Next, the plate was incubated for an hour. The results were then quantified by measuring the absorbance of the sample at 570 nm.

**Wound healing assay**

Cells were cultured to about 80% confluence before FBS starvation for 24 h. A 200 μl conventional pipette tip was used to create a scratch wound across the cell layer, followed by rinsing with PBS twice. Thereafter, culture was conducted in 1% FBS-containing DMEM. The wound width was evaluated via Image-Pro Plus 6.0 software after 20h.

**Protein extraction and western blot analysis**

Cells were washed twice with cold PBS and lysed with RIPA buffer containing PMSF at 4°C for 30 min. Then, the cells were collected and centrifuged at 4°C and 12,000 rpm for 15 min. For western blot assay, 20 μg of protein was isolated on 10% SDS-polyacrylamide gels, and transferred onto polyvinylidene difluoride (PVDF) membranes (Millipore, Bedford, MA, USA). The membranes were then blocked with TBST containing 5% fat-free milk, and incubated overnight at 4°C with primary antibodies against p-Akt (Ser473; 1:1000), pan-Akt (1:1000), LRP5 (1:1000), p-mTOR (52448; 1:1000), mTOR (1:1000), caspase-3 (1:500), E-caderin (1:1000), β-Catenin (1:1000), PCNA (1:1000), Beta-Actin (1:1000), and GAPDH (1:1000). The membranes were then washed with TBST, and incubated with HRP-conjugated secondary antibodies (1:4,000) for 1 h at room temperature. After being washed three times with TBST, the membrane was subjected to ECL detection using Bio-Rad chemiluminescence system (Bio-Rad, USA). Finally, the band densities were analyzed using ImageJ software (National Institutes of Health, Bethesda, MD, USA).

**Colony formation assay**

For colony formation assay, Cells were plated at 500 cells per well in six-well plates in DMEM medium plus 10% FBS. Cells were replaced with fresh medium every two to three days. After 8 days, colonies were fixed with paraformaldehyde, stained with crystal violet, imaged and counted under an inverted microscope. Each colony contains more than 100 cells. The data represents means ± standard error of three independent wells.

**Animal model**

All animal protocols were approved by the Animal Care and Use Committee at the Zhejiang University School of Medicine and followed the NIH Guidelines for Care and Use of Animals in Research (NIH Publication No. 85-23, revised 1996). Female BALB/c nude mice at 4 weeks old were purchased from Zhejiang Academy of Medical Sciences and fed with standard rodent chow under specific pathogen-free (SPF) conditions. Mice were given a subcutaneous injection of a mixture containing 2 × 10⁶ CAL27 or 1 × 10⁶ SCC25 cells, with each cell resuspended in 150 μl of PBS. Mice were anaesthetized 30 days later and sacrificed to harvest all tumors. The tumor weight was measured. For the mouse model of lung metastasis, female BALB/c nude mice at 4 weeks old were used. About 1 × 10⁶ cells in 100 ul PBS were injected into mice through the tail vein. One month later, the left and right lungs of each animal were fixed in 4 % paraformaldehyde and sections were stained with haematoxylin and eosin (H&E) for histology examination.

**Immunofluorescence assay**

For immunofluorescence assay, cells were seeded on glass coverslips in a 6-well plate and incubated for 48 hours. Then, cells were fixed with paraformaldehyde for 15 minutes at room temperature and washed with PBS. After a blocking step with normal goat serum for 60 min at room temperature, cells were incubated with associated antibodies (E-cad, 1:100; LR5, 1:200; p-Akt, 1:200) overnight at 4°C. Subsequently, slides were washed with PBS three times and incubated with DyLight 488 or DyLight 594 second antibody for 1 hour at room temperature. Following a double PBS wash, slides were stained with DAPI (4',6-Diamidino-2-Phenylindole) and examined with a fluorescence microscope.

For immunohistochemistry assay, the tumor tissues from the mice subcutaneous xenograft model
were fixed with paraformaldehyde for 24 hours, embedded in paraffin and then sectioned (4 μm) using a RM2135 rotary microtome (Leica Geosystems). The sections were subjected to a series of preparatory treatments: roasted at 60°C for 2 hours, dewaxed with dimethyl benzene, dehydrated with alcohol, and antigen retrieval was performed with a solution. The sections were then incubated with anti-p-Akt primary antibody (diluted 1:200) overnight at 4°C following 1 × TBST washing. After 1 × TBST washing, the sections were further incubated with a biotinylated secondary antibody for 1 h before being washed and counterstained with hematoxylin (Sigma-Aldrich). Imaging was conducted using an inverted fluorescence microscope.

Bioinformatics analysis

After mRNA was extracted from CAL27 and SCC25 cells stably transfected with siLRP5 and siNC lentiviruses, followed by reverse transcription to generate cDNA, and finally the sequencing of the cDNA was obtained. After quality control and filtering of the raw sequencing data by Fastp software, the data was compared to the reference genome using Hisat2. Then the gene reads data is converted into a standardized TPM. The probe sets without corresponding gene symbols or the genes with multiple probe sets were removed or averaged respectively. The DESeq R package was used to obtain DEGs with MARS algorithm [33]. Log2FC > 1 or Log2FC < -1, p-value < 0.05 were considered to indicate statistical significance. At last, the common up-regulated genes and down-regulated genes between different groups were selected. We performed gene ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) analysis of common DEGs (119 up-regulated genes) between the siNC and siLRP5 groups. GO and KEGG enrichment analyses were performed via an R package called clusterProfiler[34]. The analyses were done by enrichGO and enrichKEGG functions, respectively [35]. The Pathview library of the bioconductor was used to generate the Akt signaling pathway. The fold values of significantly changed genes were mapped by colors on native KEGG, Akt signaling pathway, where green represents down-regulated expression and red represents up-regulated expression levels in relation to the control group. A p value < 0.05 was considered significant. The data correlating MMP1 expression levels with the survival prognosis of head and neck squamous cell carcinoma patients was obtained from GEPIA [36]. The expression data of AKT1 was obtained from proteinatlas.org and analyzed using R language, utilizing the survminer and survival packages for survival analysis curve plotting. The AKT1 expression data is included in the supplementary files.

Statistical analysis

All statistical analyses were performed using SPSS version 19.0 (SPSS, Chicago, IL). Student’s t-test or Analysis of Variance was used to compare group distributions. All results were expressed as mean ± standard deviation (S. D.). A value of P less than 0.05 was considered statistically significant.

Results

Knockdown of LRP5 promotes tongue squamous cell carcinoma CAL27 and SCC25 proliferation and migration in vitro

To study the effect of LRP5 knockdown on tongue squamous cell carcinoma, LRP5 siRNA lentivirus was used to transfect CAL27 and SCC25. We evaluated the effect of LRP5 knockdown on biological behaviors of tongue squamous cell carcinoma cell line CAL27 and SCC25. The efficiency of knockdown of LRP5 was evaluated by Western blot assay. Results of Western blot assay showed that LRP5 expression was significantly down-regulated (Fig. 1A). Following knockdown of LRP5, increased cell proliferation was observed in Alamarblue and colony formation assays. (Fig. 1B-D) Moreover, cell migration significantly elevated in the wound healing assay (Fig. 1E, F).

LRP5 knockdown promotes CAL27 and SCC25 proliferation and invasion in vivo

The lung metastasis nude mice model and subcutaneous tumor model were established, the proliferation and invasion ability of tumor cells in vivo was evaluated after LRP5 knockdown. The results showed that knockdown of LRP5 increased the metastatic nodules of lungs (Fig. 2A, B). The hematoxylin and eosin staining of lung slides also suggested a marked increase in the area of lung metastasis in the siLRP5 group (Fig. 2A). Compared to those of control xenografts, LRP5 knockdown accelerated the tumor growth in nude mice by weight over a period of 30 days in the subcutaneous tumor model. (Fig. 2C, D).

Akt pathway is activated by LRP5 knockdown

To further investigate the mechanisms of LRP5 knockdown, RNA-seq was performed. The differentially expressed genes (DEGs) after LRP5 knockdown were obtained (logFC = 1, p < 0.05) (Fig. 3A, B). Then DEGs of the two cell lines CAL27 and SCC25 were crossed, and 119 up-regulated DEGs (Table S1) were obtained after LRP5 knockdown (Fig. 3C, D).
Using functional enrichment analysis, it was obtained that the Akt signaling pathway was compensatory activated compared to the control group in both cell lines (Fig. S1). Using bioinformatic tools, we found 25 genes that were significantly upregulated in both cell lines. The heatmap displays 15 representative commonly upregulated genes (Fig. 3D). It has been reported in the literature that the expression of MMP1 is increased in tongue squamous cell carcinoma and is associated with the survival and prognosis of patients [37–39]. They [37] found that knocking down MMP1 can reduce the proliferation and invasion of the tongue squamous cell carcinoma cell line FADU, and also lead to decreased phosphorylated Akt. Additionally, we have analyzed the correlation between MMP1 expression levels and the prognosis of head and neck squamous cell carcinoma patients through database analysis (with tongue squamous cell carcinoma comprising the majority, approximately 25.2%, from GDAC Firehose). We found that high expression of MMP1 and AKT1 is associated with lower survival chances in patients with head and neck squamous cell carcinoma compared to low expression (Fig. 3E, F). PCR verified that MMP1 was indeed elevated after LRP5 knockdown (Fig. 4A). The results of western blot assay showed that the phosphorylation of Akt was increased, the expression of proliferation-related PCNA was increased, and the apoptosis-related protein caspase3 was decreased (Fig. 4B-E). We also found that the protein E-cadherin and β-Catenin related to cell adhesion and invasion were down-regulated, which further proved the molecular mechanism of increased cell migration and invasion (Fig. C, E). Furthermore, in immunofluorescence staining assay, we observed that the fluorescence intensity of p-Akt increased after LRP5 knockdown, while that of E-cadherin decreased (Fig. 4F, G). In the in vivo subcutaneous tumor xenografts model, the intensity of p-Akt immunohistochemical staining was significantly greater in the siLRP5 group than in the control group (Fig. 5A-D).

![Figure 1](https://www.jcancer.org)

**Figure 1.** LRP5 knockdown promoted the proliferation and migration of CAL27 and SCC25 cells. (A) Western blot assay showed the expression of LRP5 was down-regulated after cells were transfected with lentivirus and selected with 1 µg/ml puromycin for 7 days. (B and C) Cell proliferation was measured by alamarBlue and colony formation assays. (D) The relative value of colony formation compared to the siNC group was indicated. (E and F) Cell migration was measured by wound healing assay. Cell-based experiments were repeated three times with triplicate wells each. (*P <0.05, **P <0.001, *** P <0.0001)
Compensatory Akt activation is required for LRP5 knockdown-induced cell proliferation and migration

To determine the functional role of Akt activation in the LRP5 knockdown-induced migration and proliferation of tongue squamous cell carcinoma, cells were treated with Akt inhibitors LY294002, GSK690693 and AZD5363. As shown in Fig. 6A-E, the treatment of CAL27 and SCC25 with GSK690693 and AZD5363 inhibited cell proliferation and migration. Among them, the effect of GSK690693 was the most obvious. As shown in Fig. 6F-I, the expression of phosphorylated Akt was reduced under the action of LY294002. Interestingly, under the action of Akt inhibitors GSK690693 and AZD5363, the expression of p-Akt increased instead. This is consistent with literature reports. It is generally believed that GSK690693 and AZD5363 block the transmission of Akt signaling, making p-Akt accumulate in cells in compensation [40–44]. The phenomenon of increased levels of p-Akt upon addition of AKT inhibitors such as GSK690693 or AZD5363 may be due to a mechanism known as "feedback activation". In this scenario, AKT inhibitors suppress the activity of the AKT signaling pathway, leading cells to generate a self-regulatory response, attempting to restore the activity of the AKT signaling pathway by increasing p-Akt levels. This feedback mechanism may result in elevated p-Akt levels, despite the overall suppression of AKT signaling pathway activity. We detected p-mTOR by western blot and found that GSK690693, AZD5363, and LY294002 did inhibit the expression of p-mTOR, which is a downstream molecule of Akt signaling (Fig. 6F and 6H; Fig. S2). Therefore, it seems that the compensatory activation of Akt pathway is required for LRP5 knockdown-induced migration and proliferation of CAL27 and SCC25 cells (Graphical Abstract).
Figure 3. MMP1 and Akt pathway were induced in the siLRP5 group than the control group. CAL27 and SCC25 in LRP5 knockdown group and siNC group were analyzed by transcriptome sequencing separately. (A and B) The volcano map showed differentially expressed genes (DEGs) after knockdown of LRP5 in two kinds of cells. (C) The venn map showed that 1165 genes were upregulated in CAL27 and 989 genes were upregulated in SCC25 after knockdown of LRP5. There were 119 identical genes among them. (D) The heatmap of DEGs showed that MMP1 was elevated (Mark with a red triangle). (E) According to the GEPIA website, patients with high expression of MMP1 in head and neck squamous cell carcinoma have a lower likelihood of survival. (F) According to public data from the HPA website (supplementary files provide the raw data for analysis), R language analysis revealed that patients with high expression of AKT1 in head and neck squamous cell carcinoma have a worse prognosis relative to patients with low expression of AKT1.

Discussion

In this study, the researchers investigated the role of LRP5 in tongue squamous cell carcinoma (TSCC) and its potential as a therapeutic target. They found that knockdown of LRP5 in TSCC cell lines resulted in increased cell growth, which was attributed to the compensatory activation of the Akt signaling pathway. These findings suggest that inhibiting the Wnt signaling pathway alone may not be sufficient to limit the growth and progression of TSCC and that combined inhibition of both the Wnt and Akt pathways may be necessary.

In addition to its role in promoting cell survival and growth, Akt is also known to be involved in mediating resistance to various anti-tumor therapies [45]. Indeed, many studies have reported that treatment with chemotherapy, radiation, or targeted therapies can lead to compensatory activation of Akt in cancer cells, which in turn promotes their survival and resistance to treatment. For instance, one study showed that treatment with the chemotherapy drug cisplatin led to upregulation of Akt in ovarian cancer cells, which was associated with increased resistance to the drug [46]. Similarly, in melanoma cells, treatment with the BRAF inhibitor vemurafenib was found to induce compensatory activation of Akt, which was shown to be involved in resistance to the drug [47]. Other studies have reported similar findings in various types of cancer, including breast, prostate, and pancreatic cancer. For example, in breast cancer cells, treatment with the anti-estrogen drug tamoxifen was found to induce Akt activation, which was associated with increased survival and resistance.
to the drug [48]. In prostate cancer cells, treatment with the androgen receptor antagonist enzalutamide was found to upregulate Akt signaling, which was shown to be involved in the development of resistance to the drug [49]. Together, these findings suggest that Akt plays a critical role in mediating resistance to various anti-tumor therapies, and that targeting this pathway may represent a promising strategy to overcome treatment resistance and improve patient outcomes [50]. Further studies are needed to fully elucidate the molecular mechanisms underlying compensatory Akt activation in cancer cells and to identify new therapeutic targets for the treatment of cancer.

Figure 4. LRP5 knockdown induced compensatory activation of the Akt signaling pathway. (A) Quantitative PCR verified that MMP1 increased after LRP5 knockdown in both CAL27 and SCC25 cell lines. (B and D) After knockdown of LRP5, p-Akt expression significantly increased. In addition, the expression of apoptosis-related protein cleaved caspase-3 decreased. (C and E) Western blot showed that the expression of the cell cycle-related protein PCNA increased, which verified the accelerated cell proliferation. In addition, the expression of adhesion-related proteins E-cad and β-catenin decreased. (F and G) By immunofluorescence staining, in the siLRP5 group, the cells had less E-cad fluorescence, but stronger fluorescence expression of p-Akt compared to the siNC group. (scale bar = 10 μm)
In Ren’s paper [24], knocking down LRP5 or LRP6 promoted lung metastasis of breast cancer cells in a nude mouse model. They found that under conditions of reduced LRPS/LRP6, Wnt5a strongly activates the non-canonical pathway, leading to elevated p-c-jun and promoting tumor cell metastasis. Wnt3a can activate canonical Wnt signal transduction [51,52]. Therefore, different tumors may rely on the canonical or non-canonical Wnt signal pathway, which may lead to different roles of LRP5 in different tumors. For example, some literature reports that Wnt5a inhibits tumor growth [53,54]. Other studies have found that Wnt3a not only promotes tumor metastasis through the Wnt/β-catenin signal but also induces Rho kinase and PLC-dependent cell migration by activating PLC [52,55]. It can be seen that under different ligands and receptor combinations, the canonical or non-canonical Wnt signal pathway can act synergistically or alone on tumor cells.

The findings of this study are also consistent with previous research that has shown the complexity of signaling pathways in cancer and the potential for compensatory activation of other pathways when one is inhibited. This highlights the importance of understanding the mechanisms of action of potential therapeutic targets and the need for combination therapies that target multiple pathways. The researchers used siRNA knockdown of LRP5 to investigate its role in TSCC. While this is a common method of gene knockdown, it has limitations and can result in off-target effects. The researchers acknowledged this limitation and performed transcriptome sequencing to investigate the downstream effects of LRP5 knockdown. This approach helped to validate the findings of the study and provide insights into the mechanisms of action.
Overall, this study provides important insights into the role of LRP5 in TSCC and highlights the potential for combination therapies that target multiple pathways. However, further research is needed to validate these findings and investigate the potential for clinical translation.

Figure 6. Akt inhibitors inhibited cell proliferation and migration caused by knockdown of LRP5. (A-C) About 500 cells were plated in six-well plates. Akt inhibitors (LY294002 = 1 μM; GSK690693 = 1 μM; and AZD5363 = 1 μM) or DMSO were added to the siLRP5 group and cultured for 3 days, and then replaced with normal complete medium without inhibitors. 5 days later, the cells were fixed and stained with crystal violet. Akt inhibitors can significantly inhibited the colony formation of tumor cells.
after LRPs knockdown. Among them, GS690693 showed the most obvious inhibitory effect, and LY294002 had the worst effect. (D and E) Cells in different groups were pretreated with DMSO or Akt inhibitors (LY294002 = 1 μM, GS690693 = 1 μM; and AZD5363 = 1 μM) for 3 hours. Then, wound healing experiments were carried out. Akt inhibitors also inhibited the migration of tumor cells after LRPs knockdown by wound healing assay. (F-I) Western blot showed that Akt inhibitors increased the expression of cleaved caspase-3 and inhibited the expression of p-mTOR. Interestingly, GS690693 and AZD5363 increased p-Akt accumulation in cells. However, LY294002 decreased the expression of p-Akt. GS is the abbreviation of GS690693. AZD is the abbreviation of AZD5363. L729 is the abbreviation of LY294002. (P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001)

Supplementary Material

Acknowledgments
This work was supported by grants from the National Natural Science Foundation of China (81801026), the Natural Science Foundation of Zhejiang Province (Q16H140002).

Author contributions
All authors contributed to the study’s conception and design. Material preparation, data collection, and analysis were performed by Chengze Wang, Yongzheng Li, Xiaoyan Miao, Lingling Dong and Ying Wang. The first draft of the manuscript was written by Chengze Wang, and Guoli Yang, and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

Data availability
The data that supports the findings of this study are available in the supplementary material of this article.

Competing Interests
The authors have declared that no competing interest exists.

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