REVIEW

The role of non-coding RNAs in drug resistance of oral squamous cell carcinoma and therapeutic potential

Xiang Meng1 | Qiu-Yue Lou2 | Wen-Ying Yang1 | Yue-Rong Wang1 | Ran Chen3 | Lu Wang1 | Tao Xu4,5 | Lei Zhang1,6

1 Key Lab. of Oral Diseases Research of Anhui Province, College & Hospital of Stomatology, Hefei, Anhui 230032, P. R. China
2 Department of Epidemiology and Biostatistics, School of Public Health, Anhui Medical University, Hefei, Anhui 230032, P. R. China
3 School of Stomatology, Anhui Medical University, Hefei, Anhui 230032, P. R. China
4 Inflammation and Immune Mediated Diseases Laboratory of Anhui Province, Hefei, Anhui 230032, P. R. China
5 School of Pharmacy, Anhui Key Lab. of Bioactivity of Natural Products, Anhui Medical University, Hefei, Anhui 230032, P. R. China
6 Department of Periodontology, Anhui Stomatological Hospital affiliated to Anhui Medical University, Hefei, Anhui 230032, P. R. China

Correspondence
Tao Xu, Inflammation and Immune Mediated Diseases Laboratory of Anhui Province, School of Pharmacy, Anhui Key Lab. of Bioactivity of Natural Products, Anhui Medical University, Hefei, Anhui, 230032, P. R. China.
Email: xutao@ahmu.edu.cn
Lei Zhang, Department of Periodontology, Anhui Stomatological Hospital affiliated to Anhui Medical University, Hefei, Anhui, 230032, P. R. China.
Email: zhanglei6551@126.com

Abstract
Oral squamous cell carcinoma (OSCC), the eighth most prevalent cancer in the world, arises from the interaction of multiple factors including tobacco, alcohol consumption, and betel quid. Chemotherapeutic agents such as cisplatin, 5-fluorouracil, and paclitaxel have now become the first-line options for OSCC patients. Nevertheless, most OSCC patients eventually acquire drug resistance, leading to poor prognosis. With the discovery and identification of non-coding RNAs (ncRNAs), the functions of dysregulated ncRNAs in OSCC development and drug resistance are gradually being widely recognized. The mechanisms of drug resistance of OSCC are intricate and involve drug efflux,
epithelial-mesenchymal transition, DNA damage repair, and autophagy. At present, strategies to explore the reversal of drug resistance of OSCC need to be urgently developed. Nano-delivery and self-cellular drug delivery platforms are considered as effective strategies to overcome drug resistance due to their tumor targeting, controlled release, and consistent pharmacokinetic profiles. In particular, the combined application of new technologies (including CRISPR systems) opened up new horizons for the treatment of drug resistance of OSCC. Hence, this review explored emerging regulatory functions of ncRNAs in drug resistance of OSCC, elucidated multiple ncRNA-mediated mechanisms of drug resistance of OSCC, and discussed the potential value of drug delivery platforms using nanoparticles and self-cells as carriers in drug resistance of OSCC.

KEYWORDS
drug delivery, drug resistance, long non-coding RNAs, microRNAs, non-coding RNA (ncRNA), oral squamous cell carcinoma

1 | INTRODUCTION

Oral squamous cell carcinoma (OSCC), accounting for about 40% of head and neck squamous cell carcinoma (HNSCC), is a heterogeneous neoplasm arising from the mucosal lining of the oral cavity, including the tongue, upper and lower gingiva, oral floor, palate, and buccal mucosa [1, 2]. Due to the low rate of early diagnosis, most patients are already at advanced stages by the time of diagnosis. At present, the treatment modalities for advanced OSCC mainly include surgery, chemotherapy, radiotherapy, or combinations of these modalities. Unfortunately, despite the application of various treatment modalities in the past few decades, the five-year overall survival rate of OSCC remains at 50% [3]. Chemotherapeutic agents, including platinum drugs, 5-fluorouracil (5-FU), paclitaxel (PTX), and doxorubicin (Dox), are the most common treatment options for OSCC. However, most patients would develop drug resistance. Currently, multidrug resistance (MDR) is one of the major hurdles of failed cancer chemotherapy and contributes to poor prognosis of patients [4]. The detailed mechanisms of MDR remain to be fully elucidated.

With the development of sequencing technology, approximately 98% of the human genome is transcribed into RNA without protein-coding potential and hence is termed non-coding RNA (ncRNA) [5]. The nucleotide sequences transcribed from DNA constitute the primary structure of ncRNAs which regulate the transcriptional translation of target genes either directly or indirectly by binding to them through base-complementary pairing [6]. Furthermore, ncRNAs perform biological functions by folding to form more stable secondary or tertiary structures. For example, IncRNA maternally expressed gene 3 (MEG3), a human IncRNA, exerts its cancer suppressive effect by stimulating the p53 signaling pathway [7]. Uroda et al. [8] found that in an evolutionarily conserved region of MEG3, two distal motifs interacted to form alternative, mutually exclusive pseudoknot structures ("kissing loops") through base complementarity. The destruction of these interactions impeded the MEG3 folding, disrupted the MEG3-dependent p53 signaling, and ultimately inhibited its cancer suppressive effect. Therefore, the anticancer effect of lncRNA MEG3 can be maintained by stabilizing "kissing loops". According to the transcript size, ncRNAs can be roughly classified into small ncRNAs (18~200 nt; sncRNAs) and long ncRNAs (>200 nt; lncRNAs). There are different kinds of sncRNAs, including microRNAs (miRNAs) and small nucleolar RNAs [9]. It has been confirmed that ncRNAs exert their roles in gene regulation, mRNA maturation, and protein synthesis. Moreover, ncRNA-related signaling pathways can drive specific cell biological responses by recognizing extensive molecular targets [10]. For example, zinc finger protein 750 (ZNF750), a tumor repressor, was important for improving the prognosis of SCC patients [11]. Meanwhile, IncRNA terminal differentiation-induced noncoding RNA (TINCR) could function as its downstream target for cancer suppression. Further results revealed that ZNF750 upregulated the expression level of IncRNA TINCR and suppressed the malignant phenotype of SCC [12]. In addition, ChIRP analysis results revealed that IncRNA colon cancer-associated transcript-1 (CCAT1) exacerbated the in vitro and in vivo SCC malignant phenotype by forming a complex with master transcription factors (TFs, including TP63 and SOX2) and activating the
TP63/SOX2-CCAT1-EGFR cascade [13]. All of the above results suggested that ncRNAs can act as upstream or downstream regulators to drive SCC progression. Emerging evidence suggests that ncRNAs play roles not only in the development but also in the treatment of cancers [14, 15]. Thus, ncRNAs serve as imperative regulators in disease progression, prognosis, and treatment. In OSCC, Shi et al. [16] tested 260 OSCC serum samples and identified two miRNAs, namely miR-626 and miR-5100, which were closely and independently associated with OSCC prognosis. Similarly, the enhancer of zeste homolog 2 (EZH2) gene has been known to be a key oncogenic driver that repressed transcription [17]. After in vitro IncRNA ring finger and CCCH-type domains 2 (RC3H2) silencing, the expression level of EZH2 was downregulated, and proliferation, migration, and invasion of OSCC cells were inhibited, whereas in vivo IncRNA RC3H2 overexpression increased the expression level of EZH2 and significantly promoted the growth and invasion of OSCC cells by sponging miR-101-3p [18]. In another study, the knockdown of IncRNA urothelial cancer-associated 1 (UCA1) significantly intensified CDDP-induced apoptosis and chemosensitivity of tongue SCC (TSCC) cells, suggesting that UCA1 silencing could be used as a new strategy to improve the sensitivity of TSCC to CDDP [19]. In parallel, this provides new insights into the functions of ncRNAs in chemotherapy of OSCC. It can be seen that multiple ncRNAs can be therapeutic targets for commonly used chemotherapeutic agents against OSCC. However, while focusing on the synergistic treatment of OSCC with ncRNAs and chemotherapeutic agents, another problem on the horizon is gradually coming to light: ncRNAs-mediated drug resistance of OSCC.

Drug resistance against OSCC poses a serious threat to the patients’ survival. Thus, strategies to mitigate the drug resistance processes are urgently needed. On one hand, nanoparticles are most commonly employed as delivery vehicles for tumor drugs and are also widely applied to counteract the drug resistance of OSCC [20]. On the other hand, engineering self-cells could evade tumor cell defenses and reverse the processes of tumor drug resistance [21]. Notably, the combined use of other tools (including CRISPR systems) also opens new windows for the treatment of tumor drug resistance. Herein, this review detailed the emerging regulatory functions of ncRNAs in drug resistance of OSCC and the mechanisms of ncRNA-related OSCC chemoresistance, thereby, providing insights to alleviate drug resistance of OSCC. Importantly, the application of nano-delivery and self-cellular drug delivery platforms may provide directions for the development of ncRNAs-based approaches to mitigate the drug resistance of OSCC.

## 2 | OVERVIEW ON OSCC AND CHEMoresistance

OSCC presents with pathological changes in the oral mucosa [1]. Depending on the site of occurrence, OSCC can be subdivided into three subtypes: buccal mucosal SCC, TSCC, and lip SCC. The probability of occurring in the tongue area is about 35.3%, followed by the floor of the mouth (22.8%) and the gingiva (12.6%) [22]. In the past ten years, the difference in prevalence rates between men and women has been narrowing, approaching a ratio of 1:1 [23]. However, the total number of cases has not decreased, suggesting a gradual increase in the number of female patients. Additionally, OSCC had approximately 350,000 new cases and 170,000 deaths globally in 2018, mainly in South and East Asian countries such as India, Sri Lanka, and China [24]. Owing to its high morbidity, mortality, and histological specificity, OSCC now constitutes a global public health concern imposing an enormous burden on both individuals and society.

Multiple factors have been shown to be jointly involved in OSCC progression. Tobacco was identified as a group 1 carcinogen that contributes to OSCC and currently remains one of the most dominant risk factors for OSCC. In 2017, cigarette smoking accounted for a large share in Asia, including Indonesia, China, and Mongolia [25]. Remarkably, due to the large population base, there were more OSCC patients in China than in other countries [24] and caused a heavier burden on the country. The result of a meta-analysis that included 254 studies involving more than seven countries demonstrated a relative risk of 3.43 for OSCC in smokers compared to non-smokers [26]. 4-(Methylamino)-1-(3-pyridyl)-1-butanone (NNK), one of the major components of cigarette smoke, is well known to be a potent carcinogen. Peng et al. [27] found that NNK upregulated the expression level of miR-944 in OSCC cells. Furthermore, miR-944 elicited pro-inflammation cytokines secretion, migration, and invasion, ultimately promoting the OSCC process. Also, alcohol is established as an independent risk factor for OSCC development. The risk of OSCC in Asians is associated with alcohol abuse, with adjusted ORs varying from 4.1 to 8.8. Moreover, smoking and alcohol consumption possess synergistic effects. In individuals who overused both, the relative risk for HNSCC was ≥15 [28]. A meta-analysis showed that HNSCC had a meta-relative risk of about 7.74 for betel quid containing tobacco and 2.56 for betel quid without tobacco in the Indian subcontinent (including India and Sri Lanka) [29]. Betel quid is a blend of areca nut, slaked lime, and betel leaf, which can be combined with tobacco, sweeteners, and/or spices. Particularly in India, betel quid serves as a predominant carrier of smokeless tobacco [30]. Human
papillomavirus (HPV) is also a risk factor for the OSCC process. Notably, HPV-16 is one of the most common sub-types of HPV causing OSCC. Approximately 14.9% of OSCC patients have DNA positive for HPV-16 [31]. Therefore, the complexity of etiologies gradually becomes a major obstacle in the treatment of OSCC.

Furthermore, the pathological development of OSCC is characterized by multiple stages in the context of complex etiologies. OSCC is usually preceded by oral potentially malignant disorders (OPMD) which increases the risk of becoming or already harboring invasive carcinoma. Leukoplakia and erythroplakia are the two most common and malignant types of OPMDs [32]. Leukoplakia has no characteristic histopathology. Microscopically, leukoplakia shows hyperkeratosis, epithelial hyperplasia, and/or epithelial thickening with an annual malignant transformation rate of 2.6% [33]. Unlike leukoplakia, erythroplakia often exhibits characteristic histopathological changes, including high-grade dysplasia, carcinoma in situ, or invasive SCC. Clinically, the lesions usually appear as well-demarcated and red velvety patches. However, some lesions appear rough and granular on the surface [34]. Further result has revealed that molecular-level changes (especially in ncRNAs, including miRNAs) can identify lesions that progress to cancers. For example, combination of miR-150-5p/miR-222-3p and miR-150-5p/miR-423-5p can distinguish normal healthy individuals from leukoplakia and OSCC patients, respectively [35]. Moreover, nuclear transport was established as a key program for regulating the progression of SCC. ΔNp63α, an oncogenic transcription factor, could maintain the undifferentiated state of SCC cells by controlling their nuclear transport. Meanwhile, karyopherin-β1 (KPNB1), a nuclear transport receptor, was shown to assist ΔNp63α in performing nuclear transport [36]. Hazawa et al. [37] demonstrated that importazole (an inhibitor of KPNB1) increased the expression level of p53-upregulated modulator of apoptosis (PUMA) and decreased the expression level of nucleoparin 62 (NUP62) by attenuating ΔNp63α nuclear import, which ultimately enhanced apoptosis in SCC cells. However, it was not clarified whether ΔNp63α-mediated nuclear transport is involved in the regulation of ncRNAs in the OSCC process. Perhaps it will be a direction to explore drug resistance of OSCC in the future. Additionally, the mechanisms of clinical action of molecular alterations in OSCC, including early diagnosis and treatment, remain unclear.

In the face of complex etiologies and clinical manifestations of OSCC, a variety of chemotherapeutic agents are used in clinical treatment. Cisplatin (CDDP), also known as cisplatinum, is a first-line and cellular non-specific chemotherapeutic drug prescribed for the treatment of solid cancers such as breast cancer [38], head and neck carcinoma [39], and testicular cancer [40]. CDDP can enter cells through multiple accesses and bind with genomic DNA or mitochondrial DNA to form DNA-platinum adducts, which then arrest DNA replication, inhibit cell mitosis, and induce apoptosis [41]. Given the positive tumor cytotoxicity of CDDP, it is also used in the treatment of OSCC patients [42]. Similar to CDDP, 5-FU mainly disrupts DNA replication and inhibits thymidylate synthase [43]. In OSCC, 5-FU induces apoptosis through intrinsic mitochondrial-mediated signaling pathways [44]. PTX is considered to be an antitumor drug that stabilizes microtubules and blocks mitosis [45]. In the treatment of OSCC, PTX reduces the expression level of vascular endothelial growth factor but has no significant inhibitory effect on tumor growth [46]. Therefore, PTX is often used as an adjunct drug to other anticancer drugs in the treatment of OSCC. For example, PTX enhanced the toxicity of cetuximab to OSCC cells and induced apoptosis of OSCC cells [47]. However, the use of chemotherapeutic agents in OSCC is limited by the development of intrinsic or acquired drug resistance in patients. For example, the mechanisms responsible for CDDP resistance are multifactorial, such as enhancement of DNA repair, elevation of cellular detoxification, inhibition of apoptosis, and regulation of ncRNAs [48]. Besides CDDP, other chemotherapeutic agents, such as PTX and 5-FU, have shown varying degrees of drug resistance. As it can be seen, MDR emergence undoubtedly poses a severe test for the treatment of OSCC patients. It was shown that ncRNAs could be involved in a variety of biological processes to promote tumorigenesis, including signaling protein interaction, modulation of translation, miRNA sponge, stabilizes protein complex [49]. Yan et al. [50] performed an integrated analysis and identified that in OSCC pathogenesis, elevated expression levels of hsa-miR-21, hsa-miR-31, and hsa-miR-338 were associated with cellular protein metabolic process, macromolecule metabolic process. Decreased expression levels of hsa-miR-125b, hsa-miR-133a, hsa-miR-133b, and hsa-miR-139 were associated with negative regulation of macromolecule biosynthetic process and gene expression. All of the above aberrantly expressed miRNAs were involved in OSCC pathogenesis. Furthermore, in a meta-analysis that included 15 studies with 1200 OSCC samples, nine upregulated miRNAs (miR-21, miR-455-5p, miR-155-5p, miR-372, miR-29b, miR-1246, miR-196a, and miR-181) and seven down-regulated miRNAs (miR-204, miR-101, miR-32, miR-20a, miR-16, miR-17, and miR-125b) were identified to be associated with poor prognosis of OSCC [51]. Specifically, the pooled hazard ratio values (95% confidence interval) related to different miRNA expression for overall survival and disease-free survival were 2.65 (2.07-3.39) and 1.95 (1.28-2.98), respectively [51]. These results suggest that ncRNAs not only promoted OSCC occurrence but were also involved in poor
prognosis. Recently, ncRNAs have gradually been found to play regulatory roles in the drug resistance of the OSCC process. Therefore, elucidation of ncRNAs regulating the drug resistance of the OSCC process is urgently needed to mitigate or even avoid drug resistance.

3  FUNCTIONS AND MECHANISMS OF NCRNAS IN DRUG RESISTANCE OF OSCC

3.1  Functions of ncRNAs in drug resistance of OSCC

OSCC cells progress to drug-resistant process through multiple functional roles, of which ncRNAs appear to act as mediators. Despite broad similarities in the functional progression of drug resistance, there are differences in the functional patterns mediated by different types of ncRNAs. Herein, we present the functions of ncRNAs in intensifying or weakening drug resistance of OSCC, and highlight the significance of ncRNAs as therapeutic targets for alleviating OSCC resistance (Table 1).

3.1.1  miRNAs and drug resistance of OSCC

A variety of anticancer drugs have been prescribed to treat OSCC. Platinum drugs are widely used in clinic because of their unique anticancer mechanisms and broad spectrums of anticancer activity. By using miRNA microarrays, Yu et al. [52] showed that expression levels of miR-214 and miR-23a were upregulated and accompanied by CDDP resistance, while the expression level of miR-21 was decreased with CDDP sensitivity in TSCC CDDP-resistant subline (Tca/CDDP) cells compared to CDDP-sensitive TSCC cells. miR-372 overexpression was detected in OSCC. Further, miR-372 was found to inhibit zinc finger and BTB domain-containing 7A protein (ZBTB7A), while ZBTB7A silencing increased the oncogenic potential and drug resistance of OSCC cells. Therefore, the sensitivity of OSCC to CDDP could be improved by miR-372 silencing [53]. Besides the aforementioned signaling pathways, exosomes exert synergistic roles in miRNA-mediated drug resistance of OSCC. For example, exosomes released from CDDP-resistant OSCC cells delivered miR-21 and ultimately enhanced CDDP resistance of OSCC cells [54].

For the other two chemotherapeutic agents (5-FU and Dox), miRNAs are also found to modulate the drug resistance of OSCC. Huang et al. [55] indicated that miR-365-3p enhanced OSCC chemosensitivity to 5-FU. The expression level of miR-221 was upregulated in Dox-treated OSCC cells compared to untreated OSCC cells. However, knockdown of miR-221 enhanced the sensitivity of OSCC cells to Dox [56]. Collectively, it can be concluded that miRNAs are involved in MDR regulation. Therefore, it is reasonable to target and regulate the expression levels of miRNAs to alleviate the drug-resistant process of OSCC.

3.1.2  lncRNAs and drug resistance of OSCC

A number of lncRNAs have been identified to be abnormally expressed in OSCC cells and involved in chemoresistance via regulating different target genes and biologic processes. For example, the interactions of lncRNAs mediating drug resistance of OSCC were based on the ceRNA theory that lncRNAs could act as miRNA sponges to weaken regulations of miRNAs on mRNAs [60]. Specifically, lncRNA HOXA11-AS overexpression was detected in OSCC tissues and cells compared to adjacent normal tissues and human oral keratinocytes. Mechanistically, lncRNA HOXA11-AS sponged miR-98–5p, which in turn suppress OSCC proliferation [61]. Meanwhile, lncRNA HOXA11-AS could also inhibit the expression level of miR-214-3p, and promote drug resistance of OSCC [62]. Notably, lncRNAs could also regulate exosomal miRNAs. For example, the downregulation of lncRNA X inactive specific transcript (XIST) enhanced exosomal miRNA-503 secretion and promoted cancer metastasis [63]. However, this regulatory mechanism has not been identified in OSCC and could be further explored in the future. Several oncogenic lncRNAs, such as lncRNA UCA1, lncRNA ANRIL, as well as lncRNA HOX transcript antisense RNA (HOTAIR), have been shown to participate in the drug resistance of OSCC. LncRNA UCA1 was demonstrated to be upregulated in CDDP-resistant OSCC cells compared to CDDP-sensitive cells. Mechanistically, the expression level of miR-184 was inhibited by lncRNA UCA1. Moreover, in CDDP-resistant OSCC cells, miR-184 overexpression facilitated tumor suppression and chemosensitivity [64]. In addition to mediating the involvement of miRNAs in drug resistance of OSCC, lncRNAs may also recognize drug transporter proteins. For example, compared to normal and tumor adjacent tissues, lncRNA ANRIL was overexpressed in OSCC tissues. Meanwhile, lncRNA ANRIL silencing inhibited tumor cell proliferation, induced apoptosis, and increased CDDP cytotoxicity by impairing drug transporters MRP1 and ABCC2 [65]. Autophagy has also been shown to be involved in lncRNA-mediated CDDP resistance. For example, lncRNA HOTAIR was reported as an oncogene, which was overexpressed in OSCC cells compared to corresponding normal oral mucosa tissues and human oral keratinocytes [66]. Notably, lncRNA HOTAIR silencing inhibited cellular autophagy by downregulating expression levels of autophagy-related genes (ATG3 and ATG7), ultimately enhancing sensitivity to CDDP [67]. All of the above
| ncRNA     | Expressions | Sample(s)                      | Targets and signaling pathways | Clinical Responses | Drugs      | Reference |
|-----------|-------------|--------------------------------|--------------------------------|--------------------|------------|-----------|
| miR-214   | +           | Tca8113 and Tca/CDDP cells     | NM                             | Promotes survival  | CDDP       | [52]      |
| miR-23a   | +           | Tca8113 and Tca/CDDP cells     | TOP2B                          | Promotes survival  | CDDP       | [52]      |
| miR-21    | -           | Tca8113 and Tca/CDDP cells     | NM                             | Promotes survival  | CDDP       | [52]      |
| miR-372   | +           | SAS, OC-3, OECM-1, HSC-3 and FaDu cells | miR-372/ZBTB7A/T8 R2 axis | Anti-apoptosis     | CDDP; taxol | [53]      |
| miR-21    | +           | HSC-3-R and SCC-9-R cells      | PTEN and PDCD4                 | Promotes metastasis| CDDP       | [54]      |
| miR-365-3p| -           | OC-3, CGHNC-9, and C9-IV3 cells; clinical tumor tissues | miR-365-3p/EHF/KRT16/β5-integrin/c-Met signaling pathway | Promotes metastasis and stemness | 5-FU       | [55]      |
| miR-221   | +           | SCC-4 and SCC-9 OSCC cells     | miR-221/TIMP3 axis            | Anti-apoptosis     | Dox        | [56]      |
| miR-371   | +           | SAS cells                      | AKT, β-catenin, and Src       | Anti-apoptosis     | CDDP; taxol | [57]      |
| miR-373   | +           | SAS cells                      | AKT, β-catenin, and Src       | Anti-apoptosis     | CDDP; taxol | [57]      |
| miR-654-5p| +           | Tca-8113 and CAL-27 cells; primary fresh OSCC tissues | miR-654-5p/GRAP/Ras/E signaling pathway | Promotes proliferation | CDDP; 5-Fu | [58]      |
| miR-1246  | +           | SAS, GNM, OC-3 and Fadu cells  | miR-1246//CCNG2 axis           | Enhances stemness  | CDDP       | [59]      |
| IncRNA HOXA11-AS | + | TSCCA, CAL-27, SCC-9 and Tca8113 cells; OSCC tumor tissues | HOXA11-AS/miR-214-3p/PIM1 axis | Promotes proliferation | CDDP       | [62]      |
| IncRNA UCA1 | +       | Tca8113, TSCCA, CAL-27 and SCC-9 cells; OSCC tumor tissues | UCA1/miR-184/SF1 axis          | Promotes proliferation | CDDP       | [64]      |
| IncRNA ANRIL | +       | OSCC-3, SCC-4, HSC-3 and CAL-27 cells; OSCC tumor tissues | MK/ANRIL/Mβ and ABCC2/caspase-3/BCL-2 axis | Promotes proliferation; Anti-apoptosis | CDDP       | [65]      |
TABLE 1 (Continued)

| ncRNA      | Expressions | Sample(s)          | Targets and signaling pathways | Clinical Responses | Drugs | Reference |
|------------|-------------|--------------------|--------------------------------|--------------------|-------|-----------|
| lncRNA HOTAIR | +           | KB and CAL-27 cells | Autophagy and mTOR signaling pathway | Anti-apoptosis      | CDDP  | [66], [67] |
| circITCH   | -           | SCC-6, SCC-9, SCC-25, HN-4 and HN-6 cells; OSCC tumor tissues | miR-421/PDCD4 Axis | Promotes proliferation; Anti-apoptosis | Bortezomib | [68] [69] |

Upregulation, +; Downregulation, -; Cyclin G2, CCNG2; DNA topoisomerase II beta, TOP2B; Doxorubicin, Dox; ETS homologous factor, EHF; Grb-2-related adaptor protein, GRAP; Not mentioned, NM; Phosphatase and tensin homolog, PTEN; Programmed cell death 4, PDCD4; Proto-Oncogene serine/threonine-protein kinase, PIM1; Tissue inhibitor of metalloproteinase-3, TIMP3; Zinc finger and BTB domain-containing 7A protein, ZBTB7A.

evidence suggest that targeted inhibition of several lncRNAs may be a potential therapeutic strategy to improve CDDP resistance in OSCC patients.

3.1.3 | circRNAs and drug resistance of OSCC

It has been suggested that circRNAs are also involved in regulating the progression of drug resistance in cancers. Upregulated circITCH enhanced the toxicity of chemotherapeutic agents against drug-resistant multiple myeloma cells [68]. Similarly, the expression level of circITCH was reduced in OSCC tissues and cell lines compared to adjacent normal tissues and human oral keratinocytes, and circITCH overexpression significantly inhibited OSCC cell proliferation and induced apoptosis [69]. Therefore, it can be speculated that circRNA can also mediate the toxicity of chemotherapeutic drugs on OSCC cells. However, the mechanisms of circRNA-mediated drug resistance of OSCC need to be further explored.

Drug resistance of OSCC cells can be seen to involve a wide range of biological signaling pathways. Fortunately, mitigation of OSCC resistance by modulating ncRNAs is a reliable option. Presumably, signaling pathways identified do not yet address all barriers for drug resistance of OSCC. Therefore, more extensive and comprehensive mechanisms of ncRNAs mediating drug resistance of OSCC are expected to be elucidated.

3.2 | ncRNA-mediated mechanisms of drug resistance of OSCC

Cancers with MDR exhibit several distinctive features, including elevated activity of drug-efflux transporters, high level of apoptotic threshold, enhanced DNA repair, and autophagy-induced drug degradation, which render tumor cells refractory to chemotherapy. Furthermore, the effectiveness of chemotherapeutic agents is constrained by intrinsic or acquired resistance. Differences at the genetic level are also shown in drug resistance of OSCC cells. Five hub genes (NOTCH1, JUN, CTNNB1, CEBPA, and ETS1) were identified by bioinformatic analysis. For example, a high mRNA expression level of NOTCH1 was associated with the EMT phenotype and drug resistance progression. Conversely, the knockdown of NOTCH1 reversed EMT phenotype and drug resistance progression [70]. Additionally, multiple hub genes can be independently regulated by hsa-miR-200c-3p, hsa-miR-200b-3p, hsa-miR-429, and hsa-miR-139-5p in miRNA-mRNA targeting regulatory network [71]. These results suggested that miRNAs can regulate drug resistance-associated hub genes and accelerate or weaken the OSCC drug resistance process. Thus, ncRNAs are destined to be involved in cancer drug-resistant progression. Currently, according to stages of chemotherapeutic drug action, it has been proven that ncRNAs engage in cancer drug resistance across multiple steps, including transmembrane transport proteins (activated drug efflux), epithelial-mesenchymal transition (elevated apoptosis threshold), DNA damage repair (prolonged cell survival), autophagy (enhanced drug degradation) and so on [72]. Notably, ncRNAs are involved in the drug resistance process of OSCC.

3.2.1 | Transmembrane transport proteins

One of the root causes of drug resistance can be attributed to reduced drug concentrations, which are often caused by enhanced expression of drug efflux pump genes. It subsequently generates decreased drug influx, increased efflux, and drug sequestration in intracellular vesicles...
TABLE 2  Expression, mechanisms and clinical implications of ncRNA-mediated drug efflux in multidrug-resistant cancer

| Drug transport protein | Gene       | ncRNA       | Expression | Mechanisms                                                                 | Clinical Responses          | Reference |
|------------------------|------------|-------------|------------|----------------------------------------------------------------------------|-----------------------------|-----------|
| MDR1                   | ABCB1      | circ_0109291| +          | MiR-188-3p, targeting ABCB1, could be sponged by circ_0109291, resulting in enhanced drug resistance. | Promotes proliferation     | [80]      |
|                        |            | MALAT1       | +          | MALAT1 promoted DDP resistance via regulating P-gp, EMT, and the activation of the PI3K/AKT/m-TOR signaling pathway. | Anti-apoptosis              | [81]      |
| MRPI                   | ABCC1      | MALAT1       | +          | MALAT1 decreased DDP sensitivity by upregulating MRPI and MDR1 via STAT3 activation. | Promotes proliferation     | [87]      |
|                        |            | ANRIL        | +          | ANRIL regulated caspase-3/BCL-2 to elevate the MRPI level.                   | Promotes proliferation     | [65]      |
|                        | linc00518  |             | +          | Linc00518 sponged miR-199a and thereby promoted MRPI expression and induced drug resistance. | Anti-apoptosis              | [91]      |
| BCRP                   | ABCG2      | miR-495      | -          | MiR-495 suppressed HOXC6 to inhibit EMT while promoting apoptosis of CSCs in OSCC by inhibiting the TGF-β signaling pathway. | Enhances stemness          | [94]      |
|                        |            | miR-302      | -          | MiR-302 inhibited BCRP expression by targeting the 3’-UTR of BCRP mRNA.      | Anti-apoptosis              | [95]      |
|                        |            | miR-1246     | +          | MiR-1246 enhanced the CSCs via repression of CCNG2.                          | Enhances stemness          | [59]      |

Upregulation, +; Downregulation, -; Cancer stem cells, CSCs; Cyclin G2, CCNG2; Epithelial-mesenchymal transition, EMT.

The expression level of *MDR1* has been found to be upregulated in cells treated with chemotherapeutic agents. Mechanistically, anticancer drugs are capable of triggering epigenetic alterations in the promoter region of the *MDR1* gene, thereby resulting in a high expression level of *P-gp* in tumor cells [76]. Notably, ncRNAs are responsible for the *MDR1*-mediated drug efflux process. For example, miR-491-3p downregulated the expression level of *MDR1* by directly binding to the 3’-UTR of *ABCB1*, thereby enhancing the sensitivity of hepatoma carcinoma cells to drugs, like Dox or vinblastine [77]. In 1997, Jain et al. [78] first estimated immunoreactivity of *P-gp* in oral tissues at different stages of tumorigenesis by flow...

and compartments [73]. Furthermore, ncRNAs are also involved in the drug efflux process (Table 2). These changes mainly involve adenosine triphosphate (ATP) binding cassette (*ABC*) family proteins, which has been divided into seven subfamilies (*ABCA-ABCG*) [74]. It has been well documented that members of the *ABC* transporter protein family were associated with the MDR of OSCC include p-glycoprotein (*P-gp/MDR1/ABCB1*), the MDR-associated protein family (*MRP1/ABCC1*), and breast cancer resistance proteins (*BCRP/ABCG2*) [75]. These *ABC* family proteins have similar transmembrane domains (TMD) that can pump drugs out of cancer cells to reduce the concentration of drugs in cancer cells (Figure 1).
structures and drug efflux mechanisms of three common transmembrane transporter proteins in OSCC. (a). Secondary structure models of drug efflux transporters of the P-gp, BCRP, and MRP1. (b). Drug efflux mechanisms of ABC transporters. ABC transporters exhibit a conformational change upon substrate binding and ATP hydrolysis which drives the transport process of the substrate. (c). MDR caused by drug efflux in OSCC. Abbreviations: ABC: adenosine triphosphate binding cassette; BCRP: breast cancer resistance proteins; MDR: multidrug resistance; MRP1: MDR-associated protein family 1; OSCC: Oral squamous cell carcinoma; P-gp: p-glycoprotein

cytometry and found that its expression was significantly elevated in recurrent OSCC tissues compared to normal tissues. Meanwhile, the expression level of P-gp was higher in T4-stage compared to the T3-stage in recurrent tumors. Another result also confirmed that a high expression level of ABCB1 was correlated with high tumor grades and poor differentiation [79]. It was known that miR-188-3p, an upstream regulator of ABCB1, could be sponged by circ_0109291. Recently, Gao et al. [80] found that circ_0109291 was highly expressed in CDDP-resistant OSCC tissues and cells compared to CDDP-sensitive OSCC tissues and cells. At the same time, miR-188-3p overexpression inhibited CDDP-resistant OSCC tissues and cells compared to CDDP-sensitive OSCC tissues and cells. Furthermore, miR-188-3p inhibitors and ABCB1 overexpression reversed the inhibitory effect of circ_0109291 silencing on CDDP-resistant OSCC cells. Thus, circ_0109291 can increase the expression level of ABCB1 by sponging miR-188-3p and promote CDDP resistance of OSCC cells. In addition, lncRNA metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) that showed high expression level in CDDP-resistant OSCC cells also promoted CDDP resistance by regulating the expression level of P-gp [81].

Since P-gp cannot explain all drug-resistant processes of cancer cells, MRP1, or ABCC1, was firstly identified in small cell lung cancer [82]. Functionally, MDR1 is mainly confined to the extrusion of xenobiotics. MRP1, on the other hand, outputs both endobiotics and xenobiotics, thereby affecting physiological processes beyond drug distribution [83, 84]. The expression level of MRP1 is known to be significantly upregulated in several drug-resistant diseases, including non-small cell lung cancer [85] and epilepsy [86]. In CDDP-resistant A549 cells, expression levels of lncRNA MALAT1 and MRP1 were upregulated compared to A549 cells. It was further found that the upregulated MRP1 was triggered by lncRNA MALAT1, thereby diminishing the sensitivity of cells to CDDP [87]. All of the above evidence points to ncRNA as a potential window in the process of alleviating drug resistance. In OSCC, the expression level of MRP1 was found to be higher in cancerous tissues than in adjacent non-neoplastic tissues. Elevated expression level of MRP1 was detected in CDDP-resistant cells compared with clinical samples, suggesting intrinsic drug resistance in OSCC. Furthermore, a high expression level of MRP1 was significantly associated with OSCC clinical stage, lymph node metastasis, and histological grade [88]. Nakamura et al. [89] also revealed that an elevated expression level of MRP1 was detected in OSCC cell lines treated with CDDP compared to untreated
Epithelial-mesenchymal transition

Epithelial-mesenchymal transition (EMT) is a reversible cellular program characterized by the loss of polarity of epithelial cells and their transformation into mesenchymal cells with the ability to move freely [101]. The EMT process is initiated by EMT-activating transcription factors (EMT-TFs), including three protein families: Snail (Snail/SNAI1 and Slug/SnaI2), basic helix-loop-helix (TWIST1, TWIST2, and TCF3), and zinc-finger E-box-binding homeobox (ZEB1 and ZEB2) [Figure 2] [102, 103]. First, epithelial genes such as E-cadherin, Claudin, cytokeratin, and zona occludens 1 are suppressed whilst mesenchymal phenotype genes including Vimentin, fibronectin, N-cadherin, and matrix metalloproteinases (MMPs) are activated [104]. Then, epithelial cells lose their typical polygonal, pebbled appearance and acquire a spindle-shaped mesenchymal morphology. Subsequently, the epithelial actin architecture restructures and cells acquire motility and invasiveness by forming lamellipodia, filopodia, and invadopodia, as well as by producing MMPs to degrade extracellular matrix (ECM) proteins [105]. Further, the mesenchymal cells produced are also able to reversibly return to back to their epithelial state in a process known as mesenchymal-epithelial transformation (MET) [106]. It has been established that the EMT process plays influential roles in specific processes such as embryonic development, tissue formation, and wound healing [107]. Furthermore, activation of inappropriate EMT process contributes to malignant progression of several cancers, such as cervical cancer, OSCC, and more [108, 109].

EMT is known to encompass multiple molecules and signaling pathways, and more recently ncRNAs have been shown to serve as crucial regulators of expressions and functions of EMT-TFs in OSCC pathologic processes (Table 3). For example, the expression level of circIGHG was found to be significantly upregulated in OSCC tissues compared to adjacent non-tumor tissues and was positively associated with EMT phenotype. It was further found that circIGHG induced EMT program via targeting miR-142-5p and thus, promoted OSCC progression [111]. Emerging evidence also suggested that EMT, particularly EMT-TFs, could affect the development of tumor drug resistance. EMT-TFs can bind to promoters of certain ABC transporter genes, thereby, activating the EMT program and enhancing drug resistance [112]. It can be seen that there were broad and profound mechanisms by which ncRNAs can mediate the EMT process and promote tumor drug resistance. LncRNA TINCR, a spliced lncRNA, is essential for normal epidermal differentiation. Compared with sensitive cells, the expression level of LncRNA TINCR was significantly increased in drug-resistant cells. Mechanistically, LncRNA TINCR, sponging miR-125b, stimulated human epidermal growth factor receptor 2 (HER-2) release and induced drug resistance. Furthermore, Snail-1 was a target gene of miR-125b. LncRNA TINCR silencing reversed drug resistance and EMT process by the regulation of miR-125b targeting HER-2 and Snail-1, respectively [113]. Therefore, the EMT program and tumor drug-resistant processes regulated by ncRNAs involve a variety of complex molecules and signaling pathways, with ncRNAs often acting as upstream regulators. It can be postulated that the expression levels of ncRNAs can be determinants of the EMT program and cell drug-resistance progression, and ncRNAs can serve as a potential target regulatory window in the inhibition of EMT program and tumor drug resistance.
Currently, accumulating evidence suggests complex associations between the EMT program and drug resistance of OSCC. It has been found that MDR tumor cells induced by various chemotherapeutic agents such as CDDP and epidermal growth factor receptor (EGFR) inhibitors frequently possess EMT phenotype. For example, EMT phenotype was observed in cetuximab-resistant OSCC cells with a concomitant loss of EGFR expression [116]. CDDP-resistant OSCC cell lines manifested an EMT phenotype, with decreased expression levels of E-cadherin and increased expression levels of TWIST and N-cadherin [117]. Sun et al. [118] constructed stabilized CDDP-resistant TSCC cell models and corroborated that CDDP-resistant cells exhibited mesenchymal phenotype compared to parental cells. Further results revealed that decreased expression levels of miR-200b and miR-15b contributed to chemotherapy-induced TSCC EMT phenotype. These results indicated that drug-resistant OSCC cells induced by chemotherapeutic agents tended to exhibit EMT phenotype, which predisposed residual cancer cells for being more aggressive. Notably, the EMT program also increased MDR emergence in OSCC. Snail overexpression inhibited expression levels of E-cadherin and β-catenin in OSCC cells and promoted the EMT process, as well as enhanced cellular resistance to erlotinib [119]. Lin et al. [120] found that OSCC cells with EMT phenotype induced by PAK1 (p21 (RAC1) activated kinase 1) exhibited a CDDP resistance. PAK1 took part in the invasion, migration, and cytoskeletal remodeling of OSCC cells [121]. In addition, miR-485-5p overexpression suppressed PAK1, to further reversing the EMT process. Thus, the EMT process and chemoresistance in OSCC can often coexist and mutually reinforce each other, which may involve common molecules and signaling pathways in both. Chemotherapy can induce the EMT process to promote tumor invasion and metastasis, which in turn can lead to MDR. These feedback loops collectively contribute to the malignant development of OSCC. Moreover, the paramount regulatory roles of ncRNAs in the EMT program and drug resistance of OSCC should also be considered. A better understanding of how ncRNAs regulate the EMT process and MDR can shed light on more effective ways to improve OSCC prognosis.
### Table 3  The ncRNAs mediating the EMT process and associated mechanisms

| ncRNA         | Location | Expression | Molecular mechanisms of action in EMT                                                                 | Clinical Responses                  | Reference |
|---------------|----------|------------|-----------------------------------------------------------------------------------------------------|-------------------------------------|-----------|
| circPTK2      | 8q24.3   | -          | Via sponging miR-429/miR-200b-3p, circPTK2 promoted TGF-β-induced EMT and cancer cell invasion by targeting TIF1γ. | Promotes metastasis                 | [110]     |
| circIGHG      | 6q32     | +          | CircIGHG directly bound miR-142-5p and consequently elevated IGF2BP3 activity.                        | Enhances invasion                   | [111]     |
| lncRNA TINCR  | 19p13.3  | +          | LncRNA TINCR sponged miR-125b and released HER-2, and miR-125b promoted Snail-1 transcription.       | Promotes metastasis                 | [113]     |
| lncRNA SATB2-AS1 | 2q33.1 | -          | After downregulation of SATB2 expression, HDAC1 failed to recruit in Snail promoter and Snail transcription was promoted. | Enhances invasion                   | [114]     |
| miR-483-3p    | 11p15.5  | -          | MiR-483-3p directly targeted integrin β3, and thus repressed downstream FAK/Erk signaling pathway.   | Promotes metastasis                 | [115]     |
| miR-200b      | 1p36.33  | -          | MiR-200b promoted EMT through upregulation of BMI1 and enhanced tumor metastasis in chemotherapy-resistant TSCC. | Promotes metastasis                 | [118]     |
| miR-15b       | 3q25.33  | -          | MiR-15b promoted EMT through upregulation of BMI1 and enhanced tumor metastasis in chemotherapy-resistant TSCC. | Promotes metastasis                 | [118]     |
| miR-485-5p    | 14q32.31 | -          | Downregulation of miR-485-5p promoted PAK1 protein expression, which in turn stimulated EMT.          | Enhances invasion                   | [120]     |

Upregulation, +; Downregulation, -; B lymphoma Mo-MLV insertion region 1 homolog, BMI1; Epithelial-mesenchymal transition, EMT; Not mentioned, NM; p21 (RAC1) activated kinase 1, PAK1.

### 3.2.3 DNA damage repair

Small molecules binding to specific DNA sites are considered as potential chemotherapeutic agents. Currently, the mechanisms of many chemotherapeutic agents are to block tumor cell division by damaging DNA, such as CDDP, carboplatin, and 5-FU [122]. However, cancer cells have developed enhanced DNA repair capacity to diminish the sensitivity of cancer cells to therapy [123]. Under normal physiological conditions, multiple DNA damage responses exist for repairing damaged DNA to maintain genomic stability in cells, including base excision repair (BER), nucleotide excision repair (NER), mismatch repair (MMR), homologous recombination (HR), and non-homologous end joining (NHEJ) [124, 125]. In addition, cells can mobilize low-fidelity trans-lesion synthesis (TLS) DNA polymerase to bypass damaged DNA lesions and prevent cell death through the TLS system [126]. These repair mechanisms are of great value in maintaining normal cell survivals. Nevertheless, in cancer cells, these mechanisms...
DNA damage repair mediated by ncRNAs. Three common DNA damage repair mechanisms (NER, HR, and NHEJ) involved in ncRNA are shown here. A variety of different ncRNAs regulate OSCC drug resistance progression by inhibiting or promoting DNA damage repair mechanisms in response to multiple key molecules. Abbreviations: HR: homologous recombination; ncRNAs: noncoding RNAs; NER: nucleotide excision repair; NHEJ: non-homologous end-joining

apparently are hindrances to the efficacy of chemotherapeutic agents. Growing studies suggest that following the use of chemotherapeutic agents, enhanced DNA repair mechanisms are identified as a primary mechanism of drug resistance [127–129].

Intriguingly, alteration of molecules in signaling pathways that regulate DNA repair capacity contributes to the sensitivity and resistance of chemotherapeutic agents (Figure 3). *RAD51*, an important HR repair protein, can repair DNA damage caused by chemotherapeutic agents, thereby reducing the efficacy of drugs and thus, trigger MDR [130, 131]. Enhanced DNA repair capacity was observed in drug-resistant gastric cancer cells [132]. Further results showed that interferon regulatory factor-1, a tumor suppressor, could directly suppress the expression level of *RAD51* via binding the *RAD51* promoter, thereby impairing DNA damage repair and ultimately reversing chemoresistance [132]. The tumor suppressor complex breast cancer susceptibility gene 1; BRCA1-associated RING domain protein 1 (*BRCA1-BARD1*) repairs DSBs by HR signaling pathway [133]. Zhao et al. [134] indicated that the *BRCA1-BARD1* complex promoted the assembly of synaptic complexes, which are key intermediates in *RAD51*-mediated DNA repair. It is therefore clear that *BRCA1* and *BARD1* are key regulators of the functions of *RAD51*. These results suggest that targeting molecules that regulate DNA repair signaling pathways could be an effective approach to reversing MDR in OSCC.

It is becoming clear that ncRNAs affect protein stability and induce drug resistance in tumor cells by regulating DNA damage-responsive genes (Table 4). As a central role in NHEJ, X-Ray Cross Complementing 4 (*XRCC4*) was abundant on DNA ligase 4 and bound with *XRCC4*-like factor to form complexes that formed alternate helical filament DNA to help cells survive [135]. Reduced expression level of miR-151a was found in temozolomide (TMZ)-resistant cells. Mechanistically, miR-151a overexpression can sensitize TMZ-resistant cells by repressing *XRCC4*-mediated DNA repair [137]. More recently, mounting evidence revealed that altered expression levels of lncRNAs were present in tumor cells and that lncRNA expression profiling could correlate with the evolution of tumor drug resistance [138, 139]. By using a microarray screening, Sharma et al. [140] identified a new lncRNA caused by DNA damage, termed DNA damage-sensitive RNA1 (DDSR1). LncRNA DDSR1 was further found to be...
### TABLE 4 ncRNAs involved in the network of DNA damage repair

| ncRNA       | Expression | ncRNA targets          | Mechanisms of DNA damage repair                                                                 | Reference |
|-------------|------------|------------------------|-------------------------------------------------------------------------------------------------|-----------|
| miR-140     | -          | FEN1                   | Enhancing FEN1-mediated DNA repair                                                               | [136]     |
| miR-151a    | -          | XRCC4                  | Activating XRCC4-mediated NHEJ signaling pathway                                                | [137]     |
| lncRNA DCSR1| +          | BRCA1                  | Sequestering BRCA1-RAP80 complex via direct interactions with BRCA1; Activating HR signaling pathway | [140]     |
| lncRNA Inc-R1| +         | RAD51; miR-193a-3p     | Activating HR signaling pathway via sponging miR-193a-3p                                         | [141]     |
| lncRNA MEG3 | +          | p53                    | Triggering HR signaling pathway via elevating p53 levels                                         | [142]     |
| circ_0001946| -          | miR-7-5p, miR-671-5p, miR-1270 and miR-3156-5p | Activating NER signaling pathway                                                               | [145]     |
| circ_0062020| +          | TRIP13; miR-615-5p     | Activating NHEJ signaling pathway via sponging miR-615-5p                                       | [146]     |

Upregulation, +; Downregulation, -; Flap endonuclease 1, FEN1; Temozolomide, TMZ; Thyroid hormone receptor interactor 13, TRIP13.

involved in HR signaling pathway to enhance DNA repair. Lnc-R1 silencing interfered with HR signaling pathway, increased DSBs level, and decreased the expression level of RAD51. It was further found that miR-193a-3p could bind to Inc-R1 and RAD51 mRNA and block their expression. Thus, Inc-R1 competitively recognized miR-193a-3p to increase the expression level of RAD51 and stabilized HR signaling pathway [141]. The expression levels of circRNAs, a type of young ncRNAs, were detected as disordered in a variety of cancers and were considered as key regulators of cancer development, invasion, and metastasis [143, 144]. In recent years, circRNAs have also been found to participate in the progression of tumor drug resistance. In prostate cancer, circ_0062020 overexpression promoted the expression level of thyroid hormone receptor-interacting protein 13 (TRIP13) via sponging miR-615-5p [146]. NHEJ signaling pathway was thus actuated, leading to enhanced DNA repair and decreased drug sensitivity of tumor cells [147]. Although the biological function of circRNAs is not fully understood, explorations of a few circRNAs have provided navigation to mechanisms of tumor drug resistance. Research on multimolecular interactions (i.e. miRNAs-lncRNAs-circRNAs) could help to fully elucidate the drug resistance mechanisms of OSCC.

#### 3.2.4 Autophagy

Autophagy is a highly conserved cellular process of intracellular lysosomal degradation and organelle recycling controlled by more than 40 autophagy-related genes [148, 149]. Autophagy can be categorized as at least three distinct forms: macroautophagy [150], microautophagy (wrapping and degrading cytoplasmic components by bending and folding inwards of lysosomal membranes) [151], and chaperone-mediated autophagy (direct transport of misfolded proteins recognized by translocons into lysosomes) [152]. The most typical type of autophagy is macroautophagy, so the term “autophagy” is often used to refer to macroautophagy. Autophagy plays roles in health and disease states, such as embryonic development [153] and neurodegenerative diseases [154]. Notably, autophagy is involved in cancer progression. Intriguingly, autophagy can exert both pro- and anticancer effects. Taking cancer promotion as an example, when cells are starved, autophagy is enhanced to meet the energy and material needs of cancer cells. The neighbor of BRCA1 gene 1 (NBR1), an autophagy-related receptor, could steer MHC I from the surface of cancer cells into the cytoplasm. As a result, autophagy can reduce the expression level of MHC I on the surface of cancer cells, thereby impeding antigen presentation [155]. In mouse models, blocking the autophagy process restored the expression level of MHC I on the surface of cancer cells, which in turn enhanced antigen presentation and thus, weaken the immune escape of cancer cells [156].

As a “junk cleaner”, the autophagy process is also detected in OSCC. Elevated expression level of LC3II (an autophagy-related gene) was observed in OSCC, whereas migration and invasion of OSCC cells were silenced by
autophagy gene blockers [157]. It is well known that ncRNAs have been shown to function as cancer suppressors or oncogenes in OSCC progression. Then, whether autophagy and ncRNAs have synergistic or antagonistic effects in OSCC has attracted the attention of scholars. Gao et al. [158] found that hypoxia enhanced both the expression level of circCDR1as and the autophagy process in OSCC cells. Mechanistically, circCDR1as facilitated the autophagy process via targeting multiple key regulators. Meanwhile, circCDR1as enhanced the autophagy process in OSCC cells by sponging miR-671-5p, inhibiting mTOR, and upregulating AKT and ERK signaling pathways. Thus, ncRNAs serve as upstream messengers along signaling pathways responsible for the regulation of OSCC by autophagy, indicating that targeting ncRNAs could alleviate the autophagy-related OSCC process (Figure 4). However, it remains largely unknown as to how autophagy is regulated by other ncRNAs in OSCC.

Autophagy and ncRNAs are not only involved in cancer progression, but in recent years the two have been found to jointly regulate the development of cancer drug resistance (Table 5). For example, miR-519a enhanced TMZ-induced autophagy and apoptosis processes, while inhibition of miR-519a decreased cellular autophagy and promoted TMZ resistance. In vivo, miR-519a sensitized cells to TMZ and enhanced apoptosis by boosting the cellular autophagy process [159]. In OSCC, autophagic flux was higher in drug-resistant cells compared to parental cells, and inhibition of autophagy led to decreased stemness, suggesting that autophagy enhanced CDDP-induced stemness and chemoresistance in OSCC cells [79]. Interestingly, ncRNAs are also implicated in the autophagy-mediated drug-resistant process in OSCC cells. The expression levels of ATG3 and ATG7 were reduced after IncRNA HOTAIR blocking, which inhibited autophagy. In parallel, the rate of apoptosis increased and the sensitivity of OSCC cells to CDDP was enhanced [67]. Ectopic expression of IncRNA highly upregulated in liver cancer (HULC) induced autophagy process in cancer cells, while IncRNA HULC silencing sensitized cancer cells to anti-tumor drugs by inhibiting the autophagy process. Importantly, IncRNA HULC was aberrantly upregulated in OSCC cell lines compared to normal cells and inhibition of IncRNA HULC expression suppressed cancer cell proliferation and drug resistance [160, 161]. Thus, inhibition of IncRNA HULC to alleviate OSCC resistance may be partially mediated by the autophagic signaling pathways. Collectively, inhibition of the autophagy process modulated by expression levels of ncRNAs can impair resistance of OSCC cells to chemotherapeutic drugs, thereby, promoting apoptosis and alleviating OSCC progression. Autophagy inhibitors may be promising adjuvant approaches in the fight against...
TABLE 5  Studies on the role of ncRNA-mediated autophagy in multidrug-resistant cancer

| ncRNA       | Expression | Sample(s)                                      | Mechanism in autophagy                                                                                     | Reference |
|-------------|------------|------------------------------------------------|-------------------------------------------------------------------------------------------------------------|-----------|
| circCDR1as  | +          | OSCC tissues; Tca-8113, SCC-15, and HOK cells | Sponging miR-671-5p, inhibiting mTOR and upregulating AKT and ERK½ signaling pathways                    | [158]     |
| miR-519a    | -          | U87-MG cells (glioblastoma)                     | Enhancing chemosensitivity and promoting autophagy by targeting STAT3/Bcl2 signaling pathway                | [159]     |
| IncRNA HOTAIR | +            | CAL-27 cells                                   | Upregulating expression of MAP1LC3B, beclin1, ATG3 and ATG7                                              | [67]      |
| IncRNA HULC | +          | OSCC tissues; SCC-9, SCC-15, SCC-25 and CAL-27 cells | Enhancing EMT process                                                                                     | [161]     |
| IncRNA CASC9 | +            | OSCC tissues; SCC-15 and CAL-27 cells            | Enhancing AKT/mTOR signaling pathway                                                                    | [162]     |
| linc00160   | +          | HCC tissues; MHCC-97, HCCLM-3, Hep-3B, and Huh-7 cells | Promoting autophagy and drug resistance by regulating miR-132-targeted PIK3R3                          | [163]     |

Upregulation, +; Downregulation, -; Hepatocellular carcinoma, HCC; Microtubule-associated protein 1 light chain 3B, MAP1LC3B; Phosphoinositide-3-kinase regulatory subunit 3, PIK3R3.

the drug resistance of OSCC. Therefore, there is an urgent need to develop new autophagy modulators with higher efficacy and lower toxicity for the treatment of drug resistance of OSCC.

In general, with the progression of chemotherapeutic drug action on cancer cells and conditions of cancer cells, drug resistance mechanisms can be mainly attributed to drug efflux mediated by transmembrane transporter proteins, elevated apoptosis threshold induced by EMT, enhanced DNA repair capacity, and drug degradation due to autophagy process. Certainly, ncRNAs can also regulate the OSCC resistance process in other ways besides the aforementioned drug-resistant mechanisms. The glycolytic pathways can also be mediated by ncRNAs to trigger drug resistance of OSCC. Wang et al. [164] found that Inc-p23154 promoted Glut1 expression, triggered glycolytic dysregulation, and induced drug resistance of OSCC by directly targeting the 3’UTR of miR-378a-3p. Importantly, the mechanisms of different drug resistance of OSCC appear to be independent, but in fact the mechanisms interact with each other. For example, EMT-TFs could recognize ABC transporter genes to regulate drug efflux, and the BRCA1 gene is involved in both DNA damage repair and the autophagy process. At the same time, ncRNAs play connecting roles like bridges. Consequently, a better understanding of the mechanisms of ncRNA-mediated drug resistance of OSCC can improve the basis for developing approaches to target ncRNAs to alleviate OSCC resistance.

4 | NCRNA-CENTERED APPROACHES TO MITIGATE DRUG-RESISTANCE OF OSCC

Exploring the mechanisms of ncRNA-mediated drug resistance of OSCC could allow us to discover novel ways of attenuating or blocking drug-resistant processes. Many approaches have shown success in rescuing chemotherapeutic agents from acquired or intrinsic drug resistance. Currently, major approaches for targeting ncRNAs to alleviate drug resistance of OSCC include targeting oral cancer stem cells, the use of adjuvant drugs, and interfering with signaling pathways.

4.1 | Targeting oral cancer stem cells

Oral cancer stem cells (OCSCs) are capable of self-renewal within a long period of time and reproduce the different cell lineages found in the primary cancers [165, 166]. One of the characteristics of OCSCs is the development of drug resistance in cancer cells, including conventional chemotherapies and immunotherapies. Therefore, OCSC-targeted therapies are promising treatment approaches to overcome the drug resistance of OSCC. Among miRNA families, let-7c is widely viewed as a tumor suppressor. The expression level of let-7c was found downregulated in OCSCs, while let-7c overexpression weakened stemness hallmarks and reversed chemoresistance [167]. Moreover,
let-7c overexpression inhibited IL-8 secretion, suggesting that the upregulation of let-7c could weaken the stemness of OCSCs, thereby, enhancing the cytotoxicity of CDDP [168]. The expression level of CD133 was found elevated in OCSCs and drug resistance was enhanced. Notably, combination therapies with targeting CD133 and administering CDDP inhibited OCSC-mediated OSCC initiation [169]. Dysregulated expression level of the transcription factor SOX2 regulated drug resistance of cancer cells to existing cancer therapies [170]. The expression level of SOX2 was upregulated in OSCC, and OCSC property was strengthened. SOX2 silencing suppressed expression levels of drug-resistant genes in OSCCs. Meanwhile, the knockdown of SOX2 combining with CDDP treatment attenuated drug resistance and improved the survival of OSCC mice [171]. Natural compound honokiol diminished self-renewal of OCSCs. At the same time, honokiol potentiated the effect of CDDP and inhibited cancer stemness of OCSCs, suggesting that honokiol may be an adjunct to the treatment of OSCC [172]. Taken together, to alleviate drug resistance of OSCC, it is essential to improve knowledge on OCSCs, with a particular focus on molecular features.

4.2 Applying adjuvant drugs

Combinations of multiple drugs have also been used to induce apoptosis in drug-resistant cells via enhancing cytotoxicity. Aldo-keto reductase (AKR) 1C family has been found to be associated with drug resistance [173]. AKR1Cs (including AKR1C1, AKR1C2, AKR1C3, and AKR1C4) were shown to be upregulated in CDDP-resistant OSCC cells. Mefenamic acid, an inhibitor of AKR1Cs, restored the sensitivity of drug-resistant cells to CDDP and 5-FU [174]. Isomahanine was able to induce endoplasmic reticulum stress in drug-resistant OSCC cells, ultimately inducing apoptosis [175]. Moreover, ursolic acid (UA) inhibited the phosphorylation of the AKT/BAD signaling pathway in drug-resistant OSCC cells, which in turn activated intrinsic apoptotic mechanisms [176]. Additionally, UA attenuated cancer cell stemness and thus reversed chemoresistance by interfering with miR-149-5p [177]. As such, drugs targeting ncRNAs hold potential in alleviating the drug resistance of OSCC. However, it is unknown whether the effect of these drugs in combination with first-line chemotherapeutic agents will amplify side effects in patients. Hence, this provides a direction for future exploration of multiple combination therapies.

4.3 Interfering with drug resistance-associated signaling pathways

Targeted inhibition of EGFR and related signaling pathways has been used as a treatment option for cancers. EGFR and its downstream signaling pathways were confirmed to be associated with CDDP sensitivity. EGFR inhibitors sensitized OSCC cells to 5-FU and CDDP. It has been proposed that inhibition of the EGFR signaling pathway may serve as a reasonable strategy for the treatment of drug-resistant OSCC patients [178, 179]. Furthermore, Li et al. [180] detected a higher expression level of β-catenin in CDDP-treated OSCC cells compared to controls. The sensitivity of OSCC cells to CDDP was enhanced by β-catenin silencing. In addition, the Wnt signaling pathway was also inhibited by β-catenin silencing. This indicates that blocking the Wnt/β-catenin signaling pathway could reverse drug resistance of OSCC cells. A variety of ncRNAs are known to be involved in the Wnt/β-catenin signaling pathway, including miR-106a and IncRNA placenta-specific protein 2 (PLAC2) [181, 182]. On the other hand, it has been suggested that vitamin D can reduce the risk of many cancers. Also, in OSCC, vitamin D sensitized cancer cells to CDDP. Mechanistically, vitamin D inhibited the activation of the NF-κB signaling pathway, thereby enhancing CDDP toxicity in OSCC cells [183]. It is well established that ncRNAs and NF-κB signaling pathway are involved in OSCC process while it is unclear whether there are ncRNAs which regulate vitamin D-mediated inhibition of NF-κB signaling pathway. Therefore, interfering with drug-resistant signaling pathways mediated by ncRNAs can be an intervention to reverse the drug resistance of OSCC and enhance chemotherapeutic drug toxicity.

At the current stage, the development of ncRNA-based therapeutic approaches for drug resistance of OSCC has been correspondingly successful. However, how to modify ncRNAs to enhance the toxicity of chemotherapeutic drugs still needs to be further explored, especially in combination with emerging technologies.

5 FUTURE PERSPECTIVES

The systemic toxicity and low bioavailability of chemotherapeutic agents are current challenges in the treatment of OSCC. Hence, the development of advanced drug delivery strategies is urgently needed (Figure 5). It has been established that the toxic effects of CDDP on cancer cells can be scavenged by GSH catalyzed by GST (GSH-S-transferase). Therefore, it can be hypothesized that down-regulated GST could restore OSCC cell apoptosis induced by CDDP. Based on the above view, Han et al. [184] constructed a GST inhibitor (ethacrynic acid (ECA))-loadable nanomaterial termed MPEG-PLA-SS-ECA, and modified into nanoparticles carrying pingyangmycin and carboplatin. ECA, pingyangmycin, and carboplatin could all be released uniformly. In their results, MPEG-PLA-SS-ECA nanoparticles were shown to restore the chemosensitivity of drug-resistant OSCC cell lines. Meanwhile, Yang
et al. [185] also developed Pt(IV)-NPs assembled from biotin-labeled Pt(IV) prodrug derivative and cyclodextrin-functionalized IR780. Since IR780 acted as a targeting ligand and for mitochondria, Pt(IV)-NPs could localize in mitochondria and release CDDP, inducing mitochondrial DNA damage, thereby, downregulating GSH level and inhibiting DNA repair mechanisms. Furthermore, a new drug delivery system (FA-PEG-S-S-PCL@PTX, FA-NPs) was able to deliver PTX, inhibit GSH level, effectively mitigating OSCC progression [186]. In addition to interfering with GSH level, nano-delivery platforms that blocked the tumor cell cycle have been developed. Synthesized polyethylene glycol-graphene quantum dots-Pt (Gpt) sensitized OSCC cells to chemotherapeutic agents, ultimately blocking the S-phase cell cycle and promoting apoptosis of OSCC cells [187]. The coloaded high-density lipoprotein-mimicking nanoparticles (HMNs) comprising NLS-Dox/anti-miR21 restored drug sensitivity in cancer cells with greater cytotoxicity [188]. Considering that miR-21 was also involved in the mechanisms of OSCC resistance, it could be speculated that HMNs could reverse the drug resistance of OSCC.

Direct modification of ncRNAs, instead of loading ncRNAs, has also been proposed for targeting cancer genes. For example, siRNAs that chemically bind to carriers form carrier-siRNA conjugates, and lipid and PEG molecules modify siRNA to form self-assembled lipid nanoparticles [72]. Given the complex intracellular microenvironment, it is recommended that targeting ncRNAs with two or more different vectors improve drug delivery. Altogether, nanoparticles have shown superiority against tumor cell chemoresistance, and therefore, they can serve as a potential strategy against OSCC-resistant patients.

In recent years, CRISPR systems have also been expanded for application in tumor drug resistance therapy. On the one hand, key drivers of drug-resistant tumors can be screened by CRISPR systems. By using genomewide CRISPR/Cas9 library screening, phosphoglycerate dehydrogenase (PHGDH) was identified as a key driver of sorafenib resistance in hepatocellular carcinoma, suggesting that PHGDH could serve as a target for alleviating sorafenib resistance [189]. This application could support precision medicine for tumor drug resistance. On the
other hand, Rosenblum et al. [190] used lipid nanoparticles (LNP) as a delivery vehicle to encapsulate Cas9 mRNA and sgRNAs. As a result, this delivery system was validated to significantly inhibit tumor growth and increase survival rate by 80% in mouse models of glioblastoma and metastatic ovarian cancer. These strategies provided a clue to treating OSCC resistance by using CRISPR systems to screen for key drivers of OSCC resistance, followed by combining CRISPR systems with multiple technologies (like nanotechnology) to construct targeted drug delivery platforms. This concept will be the direction of future research.

Besides the two strategies mentioned above, the “self-cellular drug delivery” platform has recently acquired considerable attention. Both glucosamine and glucose were known to be recognized by glucose transporters on the surface of red blood cells. By conjugating insulin to glucosamine, the red blood cells thus served as the insulin-carrying van. Under high blood glucose levels, glucose competitively bound transporter proteins, resulting in insulin freeing and thus lowering blood glucose levels. Notably, red blood cells can be replaced by nanoparticles modified with glucose transporters, which will provide solutions for the construction of other bionic cells [191]. This system is a biocompatible “intelligent drug delivery system” that can autonomously regulate drug levels according to different conditions. Furthermore, rumenic acid (RA), an anticancer fatty acid, and Dox prodrug were enveloped in adipocytes. Modified adipocytes acted as a Trojan horse for anticancer drug delivery through lipid metabolism of tumor cells, thereby, enhancing drug transport efficiency [192]. Moreover, Ci et al. [193] used liquid nitrogen to prepare dead acute myeloid leukemia (AML) cells and constructed drug delivery vehicles that encapsulated Dox. Due to the fact that dead cells have similar protein expression as the source cells, dead cells kept their bone marrow homing capability of live AML cells. The results of in vivo experiments revealed that cryo-shocked cancer cells prolonged the blood half-life of drugs and improved enrichment of chemotherapeutic drugs in the bone marrow. In addition, the therapy significantly prolonged the survival of AML mice when combined with immune adjuvants.

Compared to drug delivery platforms constructed with synthetic nanomaterials, drug carriers based on self-cells maintained cellular targeting as well as good biocompatibility. However, the self-cellular drug delivery platform remains in initial stages, and it will be possible to modify dead cells carrying multiple anticancer drugs or to loading different drugs in different locations of the dead cells at later stages. Collectively, several platforms based on nanodelivery and self-cellular drug delivery hold promise for the treatment of many drug-resistant cancers, especially OSCC.

6 | CONCLUSIONS
Growing evidence suggests the involvement of ncRNAs in drug resistance of OSCC. The important roles of ncRNAs in drug resistance make them potential targets for cancer therapies. This review highlighted the functions and mechanisms of ncRNAs such as miRNAs, lncRNAs, and circRNAs in the drug-resistant process of OSCC, and ultimately elucidated that treatment targeting these aberrantly expressed ncRNAs would be a promising approach to reverse drug resistance. Therapeutic interventions based on ncRNAs in combination with conventional chemotherapies may be an ideal option to address drug resistance in OSCC patients. However, it remains a challenge to screen out key ncRNAs from the large number of ncRNAs. Although ncRNAs have been extensively studied in OSCC, their roles as therapeutic targets for drug resistance of OSCC remains to be explored in depth. Moreover, studies have mainly focused on miRNAs and lncRNAs, and exploration of circRNA-mediated drug resistance of OSCC is still relatively rare. A new space for cancer drug resistance treatment has opened by the combination of multiple technologies for the development of chemotherapy drug delivery platforms, such as nanomaterials, genome editing, and modifying self-cells. Altogether, strategies are proposed to accelerate the pace from the lab to the bedside, and ncRNAs are expected to become novel targets for OSCC drug-resistant therapies.

DECLARATIONS
AUTHORS’ CONTRIBUTIONS
Manuscript writing (original draft): MX, LQY, YWY
Conceptualization/funding acquisition: XT, ZL
Manuscript writing, review, and editing: MX, WYR, CR, WL
Manuscript writing, vetting, and final approval: MX, LQY, YWY, WYR, CR, WL, XT, ZL

ACKNOWLEDGEMENTS
We thank XT and ZL for their scientific advice and critical reading of the manuscript.

ETHICS APPROVAL AND CONSENT TO PARTICIPATE
Not applicable.

CONSENT FOR PUBLICATION
Not applicable.

CONFLICT OF INTEREST STATEMENT
The authors declare that they have no conflict of interest.
FUNDING
The present work was supported by the National Natural Science Foundation of China [Nos. 81700522]; the Natural Science Foundation of Anhui Province [1808085MH235, 1908085QH328]; the Grants for Scientific Research of BSKY from Anhui Medical University [XJ201706].

DATA AVAILABILITY STATEMENT
Not applicable.

ORCID
Xiang Meng https://orcid.org/0000-0003-1137-8482
Lei Zhang https://orcid.org/0000-0003-2424-7019

REFERENCES
1. Loganathan SK, Schleicher K, Malik A, Quevedo R, Langille E, Teng K, et al. Rare driver mutations in head and neck squamous cell carcinomas converge on NOTCH signaling. Science. 2020;367(6483):1264-9.

2. Li Q, Dong H, Yang G, Song Y, Mou Y, Ni Y. Mouse Tumor-Bearing Models as Preclinical Study Platforms for Oral Squamous Cell Carcinoma. Front Oncol. 2020;10:212.

3. Bloebaum M, Poort L, Böckmann R, Kessler P. Survival after curative surgical treatment for primary oral squamous cell carcinoma. J Cranio-maxillofac Surg. 2014;42(8):1572-6.

4. Garcia-Mayea Y, Mir C, Masson F, Paciucci R, ME LL. Insights into new mechanisms and models of cancer stem cell multidrug resistance. Semin Cancer Biol. 2020;60:166-80.

5. Adams BD, Parsons C, Walker L, Zhang WC, Slack FJ. Targeting noncoding RNAs in disease. J Clin Invest. 2017;127(3):761-7.

6. Sun Q, Hao Q, Prasanth KV. Nuclear Long Noncoding RNAs: Key Regulators of Gene Expression. Trends Genet. 2018;34(2):142-57.

7. Piccoli MT, Gupta SK, Viereck J, Foinquinos A, Samolovac S, Kramer FL, et al. Inhibition of the Cardiac Fibroblast-Enriched IncRNA Meg3 Prevents Cardiac Fibrosis and Diastolic Dysfunction. Circ Res. 2017;121(5):575-83.

8. Uroda T, Anastasakou E, Rossi A, Teulon JM, Pellequer JL, Kramer FL, et al. Inhibition of the Cardiac Fibroblast-Enriched IncRNA Meg3 Prevents Cardiac Fibrosis and Diastolic Dysfunction. Circ Res. 2017;121(5):575-83.

9. Kaikkonen MU, Adelman K. Emerging Roles of Non-Coding RNA Transcription. Trends Biochem Sci. 2018;43(9):654-67.

10. Anastasiadou E, Jacob LS, Slack FJ. Non-coding RNA networks in cancer. Nat Rev Cancer. 2018;18(5):5-18.

11. Pan L, Yang H, Xu C, Chen S, Meng Z, Li K, et al. ZNF750 inhibited the malignant progression of oral squamous cell carcinoma by regulating tumor vascular microenvironment. Biomed Pharmacother. 2018;105:566-72.

12. Hazawa M, Lin DC, Handral H, Xu L, Chen Y, Jiang YY, et al. ZNF750 is a lineage-specific tumour suppressor in squamous cell carcinoma. Oncogene. 2017;36(16):2243-54.

13. Jiang Y, Jiang YY, Xie JJ, Mayakonda A, Hazawa M, Chen L, et al. Co-activation of super-enhancer-driven CCAT1 by TP63 and SOX2 promotes squamous cancer progression. Nat Commun. 2018;9(1):3619.
29. Guha N, Warnakulasuriya S, Vlaanderen J, Straif K. Betel quid chewing and the risk of oral and oropharyngeal cancers: a meta-analysis with implications for cancer control. Int J Cancer. 2014;135(6):1433-43.

30. Arora M, Shrivastava S, Mishra VK, Mathur MR. Use of Betel Quid in India from 2009 to 2017: An Epidemiological Analysis of the Global Adult Tobacco Survey (GATS). Subst Use Misuse. 2020;55(9):1465-71.

31. Ndiaye C, Mena M, Alemany L, Arbyn M, Castellsagué X, Laporte L, et al. HPV DNA, E6/E7 mRNA, and p16INK4a detection in head and neck cancers: a systematic review and meta-analysis. Lancet Oncol. 2014;15(12):1319-31.

32. Mello FW, Miguel AFP, Dutra KL, Porporatti AL, Warnakulasuriya S, Guerra ENS, et al. Prevalence of oral potentially malignant disorders: A systematic review and meta-analysis. J Oral Pathol Med. 2018;47(7):633-40.

33. Brouns E, Baart J, Karagözoglu K, Aartman I, Bloemena E, van der Waal I. Malignant transformation of oral leukoplakia in a well-defined cohort of 144 patients. Oral Dis. 2014;20(3):e19-24.

34. Warnakulasuriya S. White, red, and mixed lesions of oral mucosa: A clinicopathologic approach to diagnosis. Periodontol 2000. 2019;80(1):89-104.

35. Chang YA, Weng SL, Yang SF, Chou CH, Huang WC, Tu SJ, et al. A Three-MicroRNA Signature as a Potential Biomarker for the Early Detection of Oral Cancer. Int J Mol Sci. 2018;19(3):758.

36. Hazawa M, Lin DC, Kobayashi A, Jiang YY, Xu L, Dewi FRP, et al. ROCK-dependent phosphorylation of NUP62 regulates p63 nuclear transport and squamous cell carcinoma proliferation. EMBO Rep. 2018;19(1):73-88.

37. Hazawa M, Yoshino H, Nakagawa Y, Shimizu M, Nitta K, Sato Y, et al. Karyopherin-β1 Regulates Radiosensitivity and Radiation-Increased Programmed Death-Ligand 1 Expression in Human Head and Neck Squamous Cell Carcinoma Cell Lines. Cancers (Basel). 2020;12(4):908.

38. Takamizawa S, Ishiki H, Shimoi T, Shimizu M, Satomi E. Neoadjuvant Cisplatin in BRCA Carriers With HER2-Negative Breast Cancer. J Clin Oncol. 2020;38(23):2699-2700.

39. Patil V, Noronha V, Dhumal SB, Joshi A, Menon N, Bhatcharjee A, et al. Low-cost oral metronomic chemotherapy versus intravenous cisplatin in patients with recurrent, metastatic, inoperable head and neck carcinoma: an open-label, parallel-group, non-inferiority, randomised, phase 3 trial. Lancet Glob Health. 2020;8(9):e1213-22.

40. Bjerring AW, Fossa SD, Haugnes HS, Nome R, Stokke TM, Haugaa KH, et al. The cardiac impact of cisplatin-based chemotherapy in survivors of testicular cancer: a 30-year follow-up. Eur Heart J Cardiovasc Imaging. 2020;jeaa289. https://doi.org/10.1093/ehjci/jeaa289.

41. Goodsell DS. The molecular perspective: Cisplatin. Stem Cells. 2006;24(3):514-5.

42. Li X, Guo S, Xiong XK, Peng BY, Huang JM, Chen MF, et al. Combination of quercetin and cisplatin enhances apoptosis in OSCC cells by downregulating XIAP through the NF-κB pathway. J Cancer. 2019;10(19):4509-21.

43. Horowitz J, Charagheff E. Massive incorporation of 5-fluorouracil into a bacterial ribonucleic acid. Nature. 1959;184:1213-5.

44. Ji N, Jiang L, Deng P, Xu H, Chen F, Liu J, et al. Synergistic effect of honokiol and 5-fluorouracil on apoptosis of oral squamous cell carcinoma cells. J Oral Pathol Med. 2017;46(3):201-7.

45. Abu Samaan TM, Samec M, Liskova A, Kubatka P, Büsselberg D. Paclitaxel’s Mechanistic and Clinical Effects on Breast Cancer. Biomolecules. 2019;9(12):789.

46. Myoung H, Hong SD, Kim YY, Hong SP, Kim MJ. Evaluation of the anti-tumor and anti-angiogenic effect of paclitaxel and thalidomide on the xenotransplanted oral squamous cell carcinoma. Cancer Lett. 2001;163(2):191-200.

47. Sawatani Y, Komiyama Y, Nakashiro KI, Uchida D, Fukushima C, Shimura M, et al. Paclitaxel Potentiates the Anticancer Effect of Cetuximab by Enhancing Antibody-Dependent Cellular Cytotoxicity on Oral Squamous Cell Carcinoma Cells In Vitro. Int J Mol Sci. 2020;21(17):6292.

48. Zhou J, Kang Y, Chen L, Wang H, Liu J, Zeng S, et al. The Drug-Resistance Mechanisms of Five Platinum-Based Antitumor Agents. Front Pharmacol. 2020;11:343.

49. Goodall GJ, Wickramasinghe VO. RNA in cancer. Nat Rev Cancer. 2021;21(2):22-36.

50. Yan ZY, Luo ZQ, Zhang LJ, Li J, Liu JQ. Integrated Analysis and MicroRNA Expression Profiling Identified Seven miRNAs Associated With Progression of Oral Squamous Cell Carcinoma. J Cell Physiol. 2017;232(8):2178-85.

51. Troiano G, Mastrangelo F, Caponio VCA, Laino L, Cirillo N, Lo Muzio L. Predictive Prognostic Value of Tissue-Based MicroRNA Expression in Oral Squamous Cell Carcinoma: A Systematic Review and Meta-analysis. J Dent Res. 2018;97(7):759-66.

52. Yu ZW, Zhong LP, Ji T, Zhang P, Chen WT, Zhang CP. MicroRNAs contribute to the chemoresistance of cisplatin in tongue squamous cell carcinoma lines. Oral Oncol. 2010;46(4):317-22.

53. Yeh LY, Yang CC, Wu HL, Kao SY, Liu CJ, Chen YF, et al. The miR-372-ZBTB7A Oncogenic Axis Suppresses TRAIL-R2 Associated Drug Sensitivity in Oral Carcinoma. Front Oncol. 2020;10:47.

54. Liu T, Chen G, Sun D, Lei M, Li Y, Zhou C, et al. Exosomes containing miR-21 transfer the characteristic of cisplatin resistance by targeting PTEN and PDCC4 in oral squamous cell carcinoma. Acta Biochim Biophys Sin. 2017;49(9):808-16.

55. Huang WC, Jang TH, Tung SL, Yen TC, Chan SH, Wang LH. A novel miR-365-3p/EHF/keratin 16 axis promotes oral squamous cell carcinoma metastasis, cancer stemness and drug resistance via enhancing β3-integrin/c-met signaling pathway. J Exp Clin Cancer Res. 2019;38(1):89.

56. Du L, Ma S, Wen X, Chai J, Zhou D. Oral squamous cell carcinoma cells are resistant to doxorubicin through upregulation of miR-221. Mol Med Rep. 2017;16(3):2659-67.

57. Lin SC, Wu HL, Yeh LY, Yang CC, Kao SY, Chang KW. Activation of the miR-371/372/373 miRNA Cluster Enhances Oncogenicity and Drug Resistance in Oral Carcinoma Cells. Int J Mol Sci. 2020;21(24):9442.

58. Lu M, Wang C, Chen W, Mao C, Wang J. miR-654-5p Targets GRAP to Promote Proliferation, Metastasis, and Chemoresistance of Oral Squamous Cell Carcinoma Through Ras/MAPK Signaling. DNA Cell Biol. 2018;37(4):381-8.

59. Lin SS, Peng CY, Liao YW, Chou MY, Hsieh PL, Yu CC. miR-1246 Targets CCNG2 to Enhance Cancer Stemness and Chemoresistance in Oral Carcinomas. Cancers (Basel). 2018;10(8):272.

60. Thomson DW, Dinger ME. Endogenous microRNA sponges: evidence and controversy. Nat Rev Genet. 2016;17(5):272-83.
61. Niu X, Yang B, Liu F, Fang Q. LncRNA HOXA11-AS promotes OSCC progression by sponging miR-98-5p to upregulate YBX2 expression. Biomed Pharmacother. 2020;121:109623.

62. Wang X, Li H, Shi J. LncRNA HOXA11-AS Promotes Proliferation and Cisplatin Resistance of Oral Squamous Cell Carcinoma by Suppression of miR-214-3p Expression. Biomed Res Int. 2019;2019:8645153.

63. Xing F, Liu Y, Wu SY, Wu K, Sharma S, Mo YY, et al. Loss of XIST in Breast Cancer Activates MSN-c-Met and Reprograms Microglia via Exosomal miRNA to Promote Brain Metastasis. Cancer Res. 2018;78(15):4316-30.

64. Fang Z, Zhao J, Xie W, Sun Q, Wang H, Qiao B. LncRNA UCA1 promotes proliferation and cisplatin resistance of oral squamous cell carcinoma by suppressing miR-184 expression. Cancer Med. 2017;6(12):2897-2908.

65. Zhang D, Ding L, Li Y, Ren J, Shi G, Wang Y, et al. Midkine derived from cancer-associated fibroblasts promotes cisplatin-resistance via up-regulation of the expression of lncRNA ANRIL in tumour cells. Sci Rep. 2017;7(1):16231.

66. Tao D, Zhang Z, Liu X, Zhang Z, Fu Y, Zhang P, et al. LncRNA HOTAIR promotes the invasion and metastasis of oral squamous cell carcinoma through metastasis-associated gene 2. Mol Carcinog. 2020;59(4):353-64.

67. Wang X, Liu W, Wang P, Li S. RNA interference of long noncoding RNA HOTAIR suppresses autophagy and promotes apoptosis and sensitivity to cisplatin in oral squamous cell carcinoma. J Oral Pathol Med. 2018;47(10):930-7.

68. Liu J, Du F, Chen C, Li D, Chen Y, Xiao X, et al. Circular RNA ITCH regulates the miR-615-3p/PRKCD axis in multiple myeloma. Life Sci. 2020;262:118506.

69. Hao C, Wangzhou K, Liang Z, Liu C, Wang L, Gong L, et al. Circular RNA ITCH Suppresses Cell Proliferation but Induces Apoptosis in Oral Squamous Cell Carcinoma by Regulating miR-421/PDCD4 Axis. Cancer Manag Res. 2020;12:5651-8.

70. Zeng D, Liang YK, Xiao YS, Wei XL, Lin HY, Wu Y, et al. Inhibition of Notch1 reverses EMT and chemoresistance to cisplatin via direct downregulation of MCAM in triple-negative breast cancer cells. Int J Cancer. 2020;147(2):490-504.

71. Wu HT, Chen WT, Li GW, Shen JX, Ye QQ, Zhang ML, et al. Analysis of the Differentially Expressed Genes Induced by Cisplatin Resistance in Oral Squamous Cell Carcinomas and Their Interaction. Front Genet. 2019;10:1328.

72. Wang WT, Han C, Sun YM, Chen TQ, Chen YQ. Noncoding RNAs in cancer therapy resistance and targeted drug development. J Hematol Oncol. 2019;12(1):55.

73. Peetla C, Vijayaraghavalu S, Labhasetwar V. Biophysics of cell membrane lipids in cancer drug resistance: Implications for drug transport and drug delivery with nanoparticles. Adv Drug Deliv Rev. 2013;65(13-14):1686-98.

74. Fletcher JI, Haber M, Henderson MJ, Norris MD. ABC transporters in cancer: more than just drug efflux pumps. Nat Rev Cancer. 2010;10(2):147-56.

75. Chen Z, Shi T, Zhang L, Zhu P, Deng M, Huang C, et al. Mammalian drug efflux transporters of the ATP binding cassette (ABC) family in multidrug resistance: A review of the past decade. Cancer Lett. 2016;370(1):153-64.

76. Baker EK, Johnstone RW, Zalcberg JR, El-Osta A. Epigenetic changes to the MDR1 locus in response to chemotherapeutic drugs. Oncogene. 2005;24(54):8061-75.

77. Zhao Y, Qi X, Chen J, Wei W, Yu C, Yan H, et al. The miR-491-3p/Spi3/ABCB1 axis attenuates multidrug resistance of hepatocellular carcinoma. Cancer Lett. 2017;408:102-11.

78. Jain V, Das SN, Luthra K, Shukla NK, Balhara R. Differential expression of multidrug resistance gene product, P-glycoprotein, in normal, dysplastic and malignant oral mucosa in India. Int J Cancer. 1997;74(1):128-33.

79. Naik PP, Mukhopadhyay S, Panda PK, Sinha N, Das CK, Mishra R, et al. Autophagy regulates cisplatin-induced stemness and chemoresistance via the upregulation of CD44, ABCB1 and ADAM17 in oral squamous cell carcinoma. Cell Prolif. 2018;51(1):e12411.

80. Gao F, Han J, Wang Y, Jia L, Luo W, Zeng Y. Circ_0109291 Promotes the Cisplatin Resistance of Oral Squamous Cell Carcinoma by Sponging miR-188-3p to Increase ABCB1 Expression. Cancer Biother Radiopharm. 2020;https://doi.org/10.1089/cbr.2020.3928.
91. Chang L, Hu Z, Zhou Z, Zhang H. Linc00518 Contributes to Multidrug Resistance Through Regulating the MiR-199a/MRP1 Axis in Breast Cancer. Cell Physiol Biochem. 2018;48(1):16-28.

92. Allikmets R, Schriml LM, Hutchinson A, Romano-Spica V, Dean M. A human placenta-specific ATP-binding cassette gene (ABCP) on chromosome 4q22 that is involved in multidrug resistance. Cancer Res. 1998;58(23):5337-9.

93. Doyle LA, Yang W, Abruzzo LV, Krogmann T, Gao Y, Rishi AK, et al. A multidrug resistance transporter from human MCF-7 breast cancer cells. Proc Natl Acad Sci USA. 1998;95(26):15665-70.

94. You X, Zhou Z, Chen W, Wei X, Zhou H, Luo W. MiRNA-495 inhibits confers inhibitory effects on cancer stem cells in oral squamous cell carcinoma through the HOXC6-mediated TGF-β signaling pathway. Stem Cell Res Ther. 2020;11(1):117.

95. Wang Y, Zhao L, Xiao Q, Jiang L, He M, Bai X, et al. miR-302a/b/c/d cooperatively inhibit BCRP expression to increase drug sensitivity in breast cancer cells. Gynecol Oncol. 2016;141(3):592-601.

96. Naito H, Wakabayashi T, Kidoya H, Muramatsu F, Takara K, Eino D, et al. Endothelial Side Population Cells Contribute to Tumor Angiogenesis and Antiangiogenic Drug Resistance. Cancer Res. 2016;76(11):3200-10.

97. Zhang P, Zhang Y, Mao L, Zhang Z, Chen W. Side population in oral squamous cell carcinoma possesses tumor stem cell phenotypes. Cancer Lett. 2009;277(2):227-34.

98. Liu Y, Cui P, Chen J, Li W. Isolation and phenotypic characterization of side population cells in oral squamous cell carcinoma. Mol Med Rep. 2015;11(5):3642-6.

99. Olteanu GE, Mihai IM, Bojin F, Gavriliuc O, Paunescu V. The natural adaptive evolution of cancer: The metastatic ability of cancer cells. Bosn J Basic Med Sci. 2020;20(3):303-9.

100. Pan G, Liu Y, Zhang L, Zhou F, Yang S. EMT-associated microRNAs and their roles in cancer stemness and drug resistance. Cancer Commun (Lond). 2021;41(3):199-217.

101. Lu W, Kang Y. Epithelial-Mesenchymal Plasticity in Cancer Progression and Metastasis. Dev Cell. 2019;49(3):361-74.

102. Lamouille S, Xu J, Derynck R. Molecular mechanisms of epithelial-mesenchymal transition. Nat Rev Mol Cell Biol. 2014;15(3):178-96.

103. Dongre A, Weinberg RA. New insights into the mechanisms of epithelial-mesenchymal transition and implications for cancer. Nat Rev Mol Cell Biol. 2019;20(2):69-84.

104. Thiery JP, Acloque H, Huang RY, Nieto MA. Epithelial-mesenchymal transitions in development and disease. Cell. 2009;139(5):871-90.
123. Stefanaki CD, Keffler K, McClintock S, Milac L, Prosperi JR. APC loss affects DNA damage repair causing doxorubicin resistance in breast cancer cells. Neoplasia. 2019;21(12):1143-50.

124. Curtin NJ. DNA repair dysregulation from cancer driver to therapeutic target. Nat Rev Cancer. 2012;12(12):801-17.

125. Tian H, Gao Z, Li H, Zhang B, Wang G, Zhang Q, et al. DNA damage response—a double-edged sword in cancer prevention and cancer therapy. Cancer Lett. 2015;358(1):8-16.

126. Patel SM, Dash RC, Hadden MK. Translesion synthesis inhibitors as a new class of cancer chemotherapeutics. Expert Opin Investig Drugs. 2021;30(1):13-24.

127. Sarkaria JN, Kitange GJ, James CD, Plummer R, Calvert H, Weller M, et al. Mechanisms of chemoresistance to alkylating agents in malignant glioma. Clin Cancer Res. 2008;14(10):2900-8.

128. O’Grady S, Finn SP, Cuffe S, Richard DJ, O’Byrne KJ, Barr MP. The role of DNA repair pathways in cisplatin resistant lung cancer. Cancer Treat Rev. 2014;40(10):1161-70.

129. Masuda H, Ozols RF, Lai GM, Fojo A, Rothenberg M, Hamilton TC. Increased DNA repair as a mechanism of acquired resistance to cis-diaminedichloroplatinum(II) in human ovarian cancer cell lines. Cancer Res. 1998;48(20):5713-6.

130. Bhat KP, Cortez D. RPA and RAD51: fork reversal, fork protection, and genome stability. Nat Struct Mol Biol. 2018;25(6):446-53.

131. Laurini E, Marson D, Fergmilia A, Aulici S, Fergmilia M, Pricl S. Role of Rad51 and DNA repair in cancer: A molecular perspective. Pharmacol Ther. 2020;208:107492.

132. Tan L, Yuan J, Zhu W, Tao K, Wang G, Gao J. Interferon regulatory factor-1 suppresses DNA damage response and reverses chemotherapy resistance by downregulating the expression of RAD51 in gastric cancer. Am J Cancer Res. 2020;10(4):1255-70.

133. Li Q, Saito TT, Martinez-Garcia M, Deshong AJ, Nadarajan S, Lawrence KS, et al. The tumor suppressor BRCA1-BARD1 complex localizes to the synaptonemal complex and regulates recombination under meiotic dysfunction in Caenorhabditis elegans. PLoS Genet. 2018;14(1):e1007701.

134. Zhao W, Steinfeld JB, Liang F, Chen X, Maranon DG, Jian Ma, et al. Hsa_circ_0001946 inhibits Lung Cancer Progression and Mediates Cisplatin Sensitivity in Non-small Cell Lung Cancer via the Nucleotide Excision Repair Signaling Pathway. Front Oncol. 2019;9:508.

135. Brouwer I, Sitters G, Candelli A, Heerema SJ, Heller I, de Melo AJ, et al. ESCRT machinery mediates selective microautophagy of endoplasmic reticulum in yeast. Embo J. 2019;39(2):e102586.

136. Kaushik S, Cuervo AM. The coming of age of chaperone-mediated autophagy. Nat Rev Mol Cell Biol. 2020;21:1-10.

137. Luo R, Su LY, Li G, Yang J, Liu Q, Yang LX, et al. Dissecting dynamic expression of autophagy-related genes during human fetal digestive tract development via single-cell RNA sequencing. Autophagy. 2019;15(7):1296-1308.

138. Schäfer JA, Schuessner JP, Bircham PW, Tsuji T, Funaya C, Pajonk O, et al. ESCRT machinery mediates selective macroautophagy of endoplasmic reticulum in yeast. Embo J. 2019;39(2):e102586.

139. Shihabudeen Haider Ali MS, Cheng X, Moran M, Haemming S, Naldrett MJ, Alvarez S, et al. LncRNA Meg3 protects endothelial function by regulating the DNA damage response. Nucleic Acids Res. 2019;47(3):1505-22.

140. Sharma V, Khurana S, Abdelmohsen K, Oberdoerffer P, Gorospe M, et al. A BRCA1-interacting lncRNA regulates homologous recombination. EMBO Rep. 2015;16(11):1520-34.

141. Shen L, Wang Q, Liu R, Chen Z, Zhang X, Zhou P, et al. LncRNA Inc-RI regulates homologous recombination repair of DNA double-strand breaks by stabilizing RAD51 mRNA as a competitive endogenous RNA. Nucleic Acids Res. 2018;46(2):717-29.

142. Zhang L, Meng X, Zhu XW, Yang DC, Chen R, Jiang Y, et al. MicroRNA-140 impedes DNA repair by targeting FEN1 and enhances chemotherapeutic response in breast cancer. Oncotarget. 2020;11(1):87-96.

143. Zhong Y, Du Y, Yang X, Mo Y, Fan C, Xiong F, et al. Circular RNAs function as ceRNAs to regulate and control human cancer progression. Mol Cancer. 2018;17(1):79.

144. Kristensen LS, Andersen MS, Stagsted LV, Ebbsen KK, Hansen TB, Kjems J. The biogenesis, biology and characterization of circular RNAs. Nat Rev Genet. 2019;20(11):675-91.

145. Huang MS, Liu JY, Xia XB, Liu YZ, Li X, Yin JY, et al. Hsa_circ_0001946 Inhibits Lung Cancer Progression and Mediates Cisplatin Sensitivity in Non-small Cell Lung Cancer via the Nucleotide Excision Repair Signaling Pathway. Front Oncol. 2019;9:508.

146. Levine B, Kroemer G. Biological Functions of Autophagy Genes: A Disease Perspective. Cell. 2019;176(1-2):11-42.

147. Mizushima N. The ATG conjugation systems in autophagy. J Cell Biol. 2016;212(1):25-37.

148. Levine B, Kroemer G. Biological Functions of Autophagy. Curr Opin Cell Biol. 2019;63:1-10.

149. McWilliams TG, Prescott AR, Villarejo-Zori B, Ball G, Boya P, Ganley IG. A comparative map of macroautophagy and mitophagy in the vertebrate eye. Autophagy. 2019;15(7):1296-1302.

150. Sharma V, Khurana S, Abdelmohsen K, Oberdoerffer P, Gorospe M, et al. A BRCA1-interacting lncRNA regulates homologous recombination. EMBO Rep. 2015;16(11):1520-34.

151. Schäfer JA, Schuessner JP, Bircham PW, Tsuji T, Funaya C, Pajonk O, et al. ESCRT machinery mediates selective macroautophagy of endoplasmic reticulum in yeast. Embo J. 2019;39(2):e102586.

152. Kaushik S, Cuervo AM. The coming of age of chaperone-mediated autophagy. Nat Rev Mol Cell Biol. 2018;19(6):365-81.

153. McWilliams TG, Prescott AR, Villarejo-Zori B, Ball G, Boya P, Ganley IG. A comparative map of macroautophagy and mitophagy in the vertebrate eye. Autophagy. 2019;15(7):1296-1308.

154. Yamamoto K, Venida A, Yano J, Biancur DE, Kakiuchi M, Hansen TK, et al. The tumor suppressor BRCA1-BARD1 complex localizes to the synaptonemal complex and regulates recombination under mitotic dysfunction in Caenorhabditis elegans. PLoS Genet. 2018;14(1):e1007701.

155. de Souza ASC, Gonçalves LB, Lepique AP, de Araujo-Souza PS. The Role of Autophagy in Tumor Immunology—Complex Mechanisms That May Be Explored Therapeutically. Front Oncol. 2020;10:603661.

156. Yamamoto K, Venida A, Yano J, Biancur DE, Kakiuchi M, Hansen TK, et al. The tumor suppressor BRCA1-BARD1 complex localizes to the synaptonemal complex and regulates recombination under meiotic dysfunction in Caenorhabditis elegans. PLoS Genet. 2018;14(1):e1007701.
MENG ET AL.

157. Fan T, Chen Y, He Z, Wang Q, Yang X, Ren Z, et al. Inhibition of ROS/NUPR1-dependent autophagy antagonises repeated cadmium exposure-induced oral squamous cell carcinoma cell migration and invasion. Toxicol Lett. 2019;314:142-52.

158. Gao L, Dou ZC, Ren WH, Li SM, Liang X, Zhi KQ. CircCDR1as upregulates autophagy under hypoxia to promote tumor cell survival via AKT/ERK(½)/mTOR signaling pathways in oral squamous cell carcinomas. Cell Death Dis. 2019;10(10):745.

159. Li H, Chen L, Li J, Jiang S, Li X, He J, et al. LncRNA-164. Wang Y, Zhang X, Wang Z, Hu Q, Wu J, Li Y, et al. LncRNA-166. Baniebrahimi G, Mir F, Khanmohammadi R. Cancer stem cells.

160. Xiong H, Ni Z, He J, Jiang S, Li X, He J, et al. miR-519a enhances chemosensitivity and promotes autophagy in glioblastoma by targeting STAT3/Bcl2 signaling pathway. J Hematol Oncol. 2018;11(1):70.

161. Su W, Tang J, Wang Y, Sun S, Shen Y, Yang H. Long non-coding RNA highly up-regulated in liver cancer promotes epithelial-to-mesenchymal transition process in oral squamous cell carcinoma. J Cell Mol Med. 2019;23(4):2645-55.

162. Yang Y, Chen D, Liu H, Yang K. Increased expression of lncRNA CASC9 promotes tumor progression by suppressing autophagy-mediated cell apoptosis via the AKT/mTOR pathway in oral squamous cell carcinoma. Cell Death Dis. 2019;10(2):41.

163. Zhang W, Liu Y, Fu Y, Han W, Xu H, Wen L, et al. Long non-coding RNA LINC00160 functions as a decoy of microRNA-132 to mediate autophagy and drug resistance in hepatocellular carcinoma via inhibition of PIK3R3. Cancer Lett. 2020;478:22-33.

164. Peng CY, Wang TY, Lee SS, Hsieh PL, Liao YW, Tsai LL, et al. Targeting CD133 in oral squamous cell carcinoma. Biomed Res Int. 2016;2016:5378567.

165. Chien CS, Wang ML, Chu PY, Chang YL, Liu WH, Yu CC, et al. Lin28B/Let-7 regulates expression of Oct4 and Sox2 and reprograms oral squamous cell carcinoma cells to a stemlike state. Cancer Res. 2015;75(12):2553-65.

166. Peng CY, Wang TY, Lee SS, Hsieh PL, Liao YW, Tsai LL, et al. Let-7c restores radiosensitivity and chemosensitivity and impairs stemness in oral cancer cells through inhibiting interleukin-8. J Oral Pathol Med. 2018;47(6):590-7.

167. Yu CC, Hu FW, Yu CH, Chou MY. Targeting CD133 in the enhancement of chemosensitivity in oral squamous cell carcinoma-derived side population cancer stem cells. Head Neck. 2016;38(1):E231-8.

168. Novak D, Hüser L, Elton JJ, Urmansky V, Altevogt P, Utikal J. SOX2 in development and cancer biology. Semin Cancer Biol. 2020;67(Pt 1):74-82.

169. Chou MY, Hu FW, Yu CH, Yu CC. Sox2 expression involvement in the oncogenicity and radiochemoresistance of oral cancer stem cells. Oral Oncol. 2015;51(1):31-9.

170. Chang MT, Lee SP, Fang CY, Hsieh PL, Liao YW, Lu MY, et al. Chemosensitizing effect of honokiol in oral carcinoma stem cells via regulation of IL-6/Stat3 signaling. Environ Toxicol. 2018;33(11):1105-12.

171. Bortolozzi R, Bresolin S, Rampazzo E, Paganin M, Maule F, Mariotto E, et al. AKR1C enzymes sustain therapy resistance in paediatric T-ALL. Br J Cancer. 2018;118(7):985-94.

172. Shiiba M, Yamagami H, Yamamoto A, Minakawa Y, Okamoto A, Kasamatsu A, et al. Mefenamic acid enhances anticancer drug sensitivity via inhibition of aldo-keto reductase IC enzyme activity. Oncol Rep. 2017;37(4):2025-32.

173. Utaipan T, Athipornchai A, Suksamrarn A, Chunsriviroth S, Chunglok W. Isomahanine induces endoplasmic reticulum stress and simultaneously triggers p38 MAPK-mediated apoptosis and autophagy in multidrug-resistant human oral squamous cell carcinoma cells. Oncol Rep. 2017;37(2):1243-52.

174. Chen CF, Yang JS, Chen WK, Lu CC, Chiang JH, Chiu HY, et al. Ursolic acid elicits intrinsic apoptotic machinery by downregulating the phosphorylation of AKT/BAD signaling in human cisplatin-resistant oral cancer CAR cells. Oncol Rep. 2018;40(3):1752-60.

175. Chen Q, Luo J, Wu C, Lu H, Cai S, Bao C, et al. The miRNA-149-5p/MyD88 axis is responsible for ursolic acid-mediated attenuation of the stemness and chemoresistance of non-small cell lung cancer cells. Environ Toxicol. 2020;35(5):561-9.

176. Ohnishi Y, Yasui H, Kakudo K, Nozaki N, Mietzsch M. Cetuximab-resistant oral squamous cell carcinoma cells become sensitive in anchorage-independent culture conditions through the activation of the EGFR/AKT pathway. Int J Oncol. 2015;47(6):2165-72.

177. Chen YJ, Chen SY, Lovel R, Ku YC, Lai YH, Hung CL, et al. Enhancing chemosensitivity in oral squamous cell carcinoma by lentivirus-vector-mediated RNA interference targeting EGFR and MRP2. Oncol Lett. 2016;12(3):2107-14.

178. Li L, Liu HC, Wang C, Liu X, Hu FC, Xie N, et al. Overexpression of β-Catenin Induces Cisplatin Resistance in Oral Squamous Cell Carcinoma. Biomed Res Int. 2016;2016:5378567.

179. Zhang N, Wei ZL, Yin J, Zhang L, Wang J, Jin ZL. MiR-106a* inhibits oral squamous cell carcinoma progression by directly targeting MeCP2 and suppressing the Wnt/β-catenin pathway. Am J Transl Res. 2018;10(11):3542-54.

180. Chen F, Qi S, Zhang X, Wu J, Yang X, Wang R. lncRNA PLAC2 activated by H3K27 acetylation promotes cell proliferation and invasion via the activation of Wnt/β-catenin pathway in oral squamous cell carcinoma. Int J Oncol. 2019;54(4):1183-94.

181. Huang Z, Zhang Y, Li H, Zhou Y, Zhang Q, Chen R, et al. Vitamin D promotes the cisplatin sensitivity of oral squamous cell carcinoma by inhibiting LCN2-modulated NF-κB pathway activation through RPS3. Cell Death Dis. 2019;10(12):936.

182. Han B, Wang Y, Wang L, Shang Z, Wang S, Pei J. Preparation of GST Inhibitor Nanoparticle Drug Delivery System and Its Reversal Effect on the Multidrug Resistance in Oral Carcinoma. Nanomaterials (Basel). 2015;5(4):1571-87.

183. Yang GG, Pan ZY, Zhang DY, Cao Q, Ji LN, Mao ZW. Precisely Assembled Nanoparticles against Cisplatin Resistance via Cancer-Specific Targeting of Mitochondria and Imaging-Guided Chemo-Photothermal Therapy. ACS Appl Mater Interfaces. 2020;12(39):43444-55.
taxel to enhance oral squamous cell carcinoma therapy. J Mater Chem B. 2020;8(15):3113-22.

187. Wei Z, Yin X, Cai Y, Xu W, Song C, Wang Y, et al. Antitumor effect of a Pt-loaded nanocomposite based on graphene quantum dots combats hypoxia-induced chemoresistance of oral squamous cell carcinoma. Int J Nanomedicine. 2018;13:1505-24.

188. Rui M, Qu Y, Gao T, Ge Y, Feng C, Xu X. Simultaneous delivery of anti-miR21 with doxorubicin prodrug by mimetic lipoprotein nanoparticles for synergistic effect against drug resistance in cancer cells. Int J Nanomedicine. 2017;12:217-37.

189. Wei L, Lee D, Law CT, Zhang MS, Shen J, Chin DW, et al. Genome-wide CRISPR/Cas9 library screening identified PHGDH as a critical driver for Sorafenib resistance in HCC. Nat Commun. 2019;10(1):4681.

190. Rosenblum D, Gutkin A, Kedmi R, Ramishetti S, Veiga N, Jacobi AM, et al. CRISPR-Cas9 genome editing using targeted lipid nanoparticles for cancer therapy. Sci Adv. 2020;6(47):eabc9450.

191. Wang C, Ye Y, Sun W, Yu J, Wang J, Lawrence DS, et al. Red Blood Cells for Glucose-Responsive Insulin Delivery. Adv Mater. 2017;29(18). https://doi.org/10.1002/adma.201606617.

192. Wen D, Wang J, Van Den Driessche G, Chen Q, Zhang Y, Chen G, et al. Adipocytes as Anticancer Drug Delivery Depot. Matter. 2019;1(5):1203-14.

193. Ci T, Li H, Chen G, Wang Z, Wang J, Abdou P, et al. Cryo-shocked cancer cells for targeted drug delivery and vaccination. Sci Adv. 2020;6(50):eabc3013.

How to cite this article: Meng X, Lou Q-y, Yang W-y, Wang Y-r, Chen R, Wang L, et al. The role of non-coding RNAs in drug resistance of oral squamous cell carcinoma and therapeutic potential. Cancer Commun. 2021;41:981–1006. https://doi.org/10.1002/cac2.12194