Research Article

Active Component of Antrodia cinnamomea Mycelia Targeting Head and Neck Cancer Initiating Cells through Exaggerated Autophagic Cell Death

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Head and neck squamous cell carcinoma (HNSCC) is a highly lethal cancer. Previously, we identify head and neck cancer initiating cells (HN-CICs), which are highly tumorigenic and resistant to conventional therapy. Therefore, development of drug candidates that effectively target HN-CICs would benefit future head and neck cancer therapy. In this study, we first successfully screened for an active component, named YMGKI-1, from natural products of Antrodia cinnamomea Mycelia (ACM), which can target the stemness properties of HNSCC. Treatment of YMGKI-1 significantly downregulated the aldehyde dehydrogenase (ALDH) activity, one of the characteristics of CIC in HNSCC cells. Additionally, the tumorigenic properties of HNSCC cells were attenuated by YMGKI-1 treatment in vivo. Further, the stemness properties of HN-CICs, which are responsible for the malignancy of HNSCC, were also diminished by YMGKI-1 treatment. Strikingly, YMGKI-1 also effectively suppressed the cell viability of HN-CICs but not normal stem cells. Finally, YMGKI-1 induces the cell death of HN-CICs by dysregulating the exaggerated autophagic signaling pathways. Together, our results indicate that YMGKI-1 successfully lessens stemness properties and tumorigenicity of HN-CICs. These findings provide a new drug candidate from purified components of ACM as an alternative therapy for head and neck cancer in the future.

1. Introduction

Head and neck squamous cell carcinoma (HNSCC) represents the sixth most common cancer worldwide and the third most common cancer in developing nations [1]. Despite the recent advancements in the multidisciplinary treatment of HNSCC, prognosis of patients with locally advanced diseases and long-term survival rates remains unsatisfactory [2].
Over the past decade, increasing evidence suggests that the hierarchical model of cancer initiating cells (CICs) or cancer stem cells (CSCs) in that each tumor formation is governed by a rare subpopulation of cells with self-renewal capacity [3]. CICs have been demonstrated to have capacities of promoting tumor growth, tumor regeneration, metastatic progression, and tumor recurrence [4, 5]. It is imperative to uncover new therapeutic drug on targeting cancer stem cells [6].

Previously, we have verified a subpopulation of HNSCC cells (HNSCCs) displaying the characteristics of CICs [5]. In addition, we have identified molecular markers such as CD133 [8] and memGRP78 (membrane anchoring GRP78) [9] for targeting CICs. Others have found that Aldehyde dehydrogenase (ALDH), which has been demonstrated as a breast cancer CIC marker [10], could also be the CIC marker of head and neck cancer and other cancers [11, 12]. More recently, we have successfully verified an epithelial-mesenchymal transition (EMT) related gene expression, S100A4, which is upregulated in HN-CICs. Consequently, we have shown that significant downregulation of ALDH activity comes along with loss of stemness properties in S100A4 knockdown HNSCCs and HN-CICs [13]. Therefore, the cell-based ALDH activity assay is useful as a screening system for drug candidates, which would target the stemness properties of cells. With the assay system, we would be able to screen for active components of natural products from Chinese medicinal herbals on target cellular ALDH activity in vitro.

It has been reported that CICs, bearing highly malignant stem cell traits acquired through processes such as the EMT, would survive during metabolic stress only if they adopt adaptive mechanisms such as autophagy [14]. However, autophagy has a dual role in CICs [15]. Autophagy has been linked to the maintenance of breast cancer stem-like phenotype [16]. In contrast, some studies demonstrate that autophagy plays an essential role in differentiation of glioma-initiating cells [17] and in inhibition of colon cancer stem cell [18]. Thus, the role of autophagy in CICs remains elusive.

*Antrodia cinnamomea*, a rare mushroom of the family Polyporaceae, only grows naturally in Taiwan [19]. For Taiwanese traditional medicinal treatment, *Antrodia cinnamomea* has been applied for diarrhea, intoxication, hypertension, abdominal pain, itchy skin [20], and cancers therapy [21]. The biological or physiological functions of the crude extracts or purified active components of fruiting bodies or submerged cultured mycelia of *Antrodia cinnamomea* have been examined. Empirically, these active components of natural products from *Antrodia cinnamomea* showed antioxidant [20], anti-inflammatory [22], hepatoprotective [23], and antitumor activities. Further, the antitumor activities of *Antrodia cinnamomea* have recently become popular as an alternative therapeutic agent for several types of human cancer [21, 24].

Several pure compounds in the fruiting body of *Antrodia cinnamomea* have been isolated and identified with antitumor activity [25]. Compounds of maleic and succinic acid derivatives from the mycelium of *Antrodia cinnamomea* exhibit significant cytotoxic effects on Lewis lung carcinoma (LLC) tumor cell [26]. However, the inhibitory effect of *Antrodia cinnamomea* on the cancer initiating cells has never been studied. Therefore, we are interested in screening and evaluating the effects of active components from *Antrodia cinnamomea* on targeting CICs and in elucidating the possible biological mechanisms mediating the antitumor activity in vitro and in vivo.

Herein, we exploited the perivous in vitro cell-based ALDH activity assay system to screen for the active components from *Antrodia cinnamomea* Mycelia extracts (ACMEs) with effective cytotoxicity toward to CICs. We used both in vitro assays and in vivo xenograft mouse models to examine the cytotoxic effects of a purified component of ACME, YMGKI-1 (a derivative of maleic and succinic acid), on the HN-CICs. Strikingly, YMGKI-1 treatment diminished stemness properties and tumorigenicity of HN-CICs both in vitro and in vivo. Finally, we demonstrated that the induction of cell death in YMGKI-1-treated HN-CICs is through dysregulatory autophagic signaling, which is regulated mainly by mTOR molecular pathway. Thus, our study indicates that we have successfully isolated active compounds from natural products of *Antrodia cinnamomea* Mycelia (ACM) with selective and effective cytotoxicity on HN-CICs. Further, our findings provide a new drug candidate as an alternative therapy for head and neck cancer in the future.

2. Materials and Methods

2.1. Extraction, Isolation, Purification, and Structure Determination of Single Compounds from the ACM. YMGKI-1 (3-[4-(3-Methylbut-2-enyloxy)phenyl]-4-isobutyl-N-hydroxy- pyrrole-2,5-dione; Mw: 329 Da) from ACMs and the chemical structure of the purified chemicals were performed and determined by Dr. Chien-Chih Chen (Hungkuang University, Taichung, Taiwan). The purification procedure of YMGKI-1 from ACMs was described in Supplementary Materials and Methods available online at http://dx.doi.org/10.1155/2013/946451. Dimethyl sulfoxide (DMSO) (Sigma-Aldrich D2650, Saint Louis, MO) is used as a drug solvent.

2.2. Cell Lines. Human tongue carcinoma cells (SAS) were obtained from the Japanese Collection of Research Bioresources (Tokyo, Japan) [27]. Human gingival squamous carcinoma cells (OECM-1) were provided from Dr. C. L. Meng (National Defense Medical College, Taipei, Taiwan). Primary culture of normal human oral keratinocytes (NHKOS) was as described [5]. Additionally, the amniotic fluid stem cells-2 (AFSC-2) and hematopoietic stem cell (HSC) were isolated by Dr. Shiu-Huey Chou at Fu-Jen University. The isolation protocols of AFSC-2 and HSC were described in Supplementary Materials and Methods.

2.3. Antibodies and Reagents. Anti-Oct-4 (MAB430), anti-Nanog (AB9220), anti-CD133 (MAB4310), and anti-GAPDH (MAB574) were obtained from Millipore Corporation (Billerica, MA). Anti-GRP78 (610978) and anti-E-cadherin (610182) were obtained from BD Bioscience (San Jose, CA). Anti-Notch2 (sc-5545) and anti-Pi3K (sc-1637) were obtained from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA). Anti-LC3
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2.4. Cell Lines Cultivation and Enrichment of HN-CICs. In brief, SAS and OECM1 were grown in DMEM or in RPMI supplemented with 10% FBS (Grand Island, NY), respectively. The SAS cells were then cultured in tumor sphere medium consisting of serum-free DMEM/F12 medium (GIBCO), N2 supplement (GIBCO), 10 ng/mL human recombinant basic fibroblast growth factor-basic (FGF), and 10 ng/mL Epidermal Growth Factor (EGF) (R&D Systems, Minneapolis, MN). Cells were plated at a density of $7.5 \times 10^4$ live cells/10 mm dish, and the medium was changed every other day until the tumor sphere formation was achieved to enrich the SAS-HNCICs or OECM1-HN-CICs in about 4 weeks [5].

2.5. ALDH Activity Assay. The ALDEFLUOR kit (Stem Cell Technologies, Durham, NC, USA) was used to examine the ALDH enzymatic activity [13]. Single cell suspension obtained from cancer cells with or without treatment was suspended in ALDEFLUOR assay buffer containing ALDH substrate (BAAA, 1 $\mu$mol/L per 1 $\times 10^6$ cells) and incubated during 40 minutes at 37°C. As negative control, for each sample of cells an aliquot was treated with 50 mmol/L diethylamino benzaldehyde (DEAB), a specific ALDH inhibitor. This resulted in a significant decrease in the fluorescence intensity of ALDH-positive cells and was used to identify the ALDH-positive cells. The amount of intracellular fluorescence was measured by FACS Calibur apparatus (Becton Dickinson, San Diego, CA).

2.6. Side Population Analysis. Cells were resuspended at 1 $\times 10^6$/ml in prewarmed DMEM with 2% FCS. Hoechst 33342 dye was added at a final concentration of 5 $\mu$g/mL in the presence or absence of fumitremorgin C (FTC) (10 $\mu$M; Sigma, St Louis, MO, USA) and was incubated at 37°C for 90 min with intermittent shaking. At the end of the incubation, the cells were washed with ice-cold HBSS with 2% FCS and centrifuged down at 4°C and resuspended in ice-cold HBSS containing 2% FCS. Propidium iodide at a final concentration of 2 $\mu$g/mL was added to the cells to gate viable cells. The cells were filtered through a 40 $\mu$m cell strainer to obtain single cell suspension before analysis. The Hoechst 33342 dye was excited at 357 nm and its fluorescence was dual-wavelength analyzed (blue, 402–446 nm; red, 650–670 nm). Analyses were done on FACS Aria (Becton Dickinson, San Diego, CA).

2.7. Cell Viability Assay. Cell viability was measured by MTT assay based on the ability of live cells to convert tetrazolium salt into purple formazan. Cells were seeded onto 24-well dishes with medium containing YMGKI-1 at concentrations ranging from 0 to 35 $\mu$g/mL for 24 hr. After treatment for the indicated times, 5 $\mu$L of MTT solution (4 mg MTT/ml PBS) was added to each well and the plate was incubated at 37°C for 3 h. After the removal of medium, 100 $\mu$L of DMSO was added to each well and the plate was gently shaken for 30 min. The absorbance was determined at 560 nm. The cell viability was calculated as OD of experimental groups/OD of control groups x 100%.

2.8. Apoptotic Assay. Apoptotic cells were detected with an Annexin V-APC kit (Calbiochem, Darmstadt, Germany) according to manufacturer’s guidelines. After staining, the cells were incubated with 20 $\mu$g/mL propidium iodide (PI) to gate the dead cells. Consequently, the fluorescent signals from Annexin V or PI were analyzed by FACS Canto apparatus (Becton Dickinson, San Diego, CA, USA).

2.9. Matrigel Invasion Assay. For transwell migration assays, 2 $\times 10^5$ cells were plated into the top chamber of a transwell (Corning, Acton, MA) with a porous transparent polyethylene terephthalate membrane (8.0 $\mu$m pore size). Cells were plated in medium with lower serum (0.5% FBS), and medium supplemented with higher serum (10% FBS) was used as a chemoattractant in the lower chamber. The cells were incubated for 24 h and cells that did not migrate through the pores were removed by a cotton swab. Cells on the lower surface of the membrane were stained with Hoechst 33258 (Sigma-Aldrich) to show the nuclei; fluorescence was detected at a magnification of 100X using a fluorescence microscope (Carl Zeiss, Oberkochen, Germany). The number of fluorescent cells in a total of five randomly selected fields was counted.

2.10. Anchorage Independent Growth Assay. Each well (35 mm) of a six-well culture dish was coated with 2 mL bottom agar (Sigma-Aldrich) mixture (DMEM, 10% (v/v) FBS, 0.6% (w/v) agar). After the bottom layer was solidified, 2 mL top agar-medium mixture (DMEM, 10% (v/v) FBS, 0.3% (w/v) agar) containing 2 $\times 10^4$ cells was added, and the dishes were incubated at 37°C for 2 weeks. Plates were stained with 0.005% Crystal Violet and then the colonies were counted. The number of total colonies was counted over five fields per well for a total of 15 fields in triplicate experiments.

2.11. Acridine Orange (AO) Staining Assay. Cells were washed with Hank’s buffered salt solution (HBSS) twice, followed by staining with 1 $\mu$g/mL acridine orange (Sigma, A 6014), and diluted in HBSS containing 5% FBS for 15 min at 37°C. After staining, cells were washed ice-cold HBSS with 5% FBS and centrifuged down at 4°C and resuspended in ice-cold HBSS containing 5% FBS. The stained cells were observed under a red filter fluorescence microscope. To quantify the AVO formation, the acridine orange stained cells were resuspended in HBSS containing 5% FBS and analyzed by FACS Canto apparatus (Becton Dickinson, San Diego, CA, USA) [28].

2.12. In Vivo Tumorigenic Assay. All the animal practices in this study were approved and in accordance with the Institutional Animal Care and Use Committee (IACUC) of National
Yang-Ming University, Taipei, Taiwan (IACUC approval nos. 1001223 and 991235). The effect of YMGKI-1 on anti-tumorigenic activity was examined in 8-week-old nude BALB/c nu/nu mice (n = 3 per group). SAS cells exposed to YMGKI-1 (1 × 10^6) were subcutaneously injected into the subcutaneous of nude mice. Tumors were generally palpable at 5 to 7 days after inoculation and tumor volumes were measured (using a caliper and calculated as length × width² × 0.5) twice weekly until 28 days after injection. To determine the therapeutic effect of YMGKI-1 on regression, SAS cells (5 × 10⁵ cells in 0.1 mL DMEM) were inoculated subcutaneously into the back of nude BALB/c nu/nu mice (n = 3 per group). Tumors were allowed to develop to be palpable. Then, the YMGKI-1 was administered intraperitoneally at days 11, 13, and 17 after SAS cell inoculation, and tumor volumes were determined twice weekly. The tumor volume was calculated according to the formula: (Length × Width²)/2 [5].

2.13. Statistics. An unpaired t-test was used for the statistical analysis. The results were considered to be statistically different when the P value was < 0.05.

3. Results

3.1. Effective Targeting of HN-CICs by YMGKI-1 Treatment. Cancer-initiating cells (CICs) are a rare subpopulation of cancer cells, which are responsible for tumor growth and cancer recurrence during conventional chemotherapy or radiotherapy [3, 4]. High ALDH activity has been used as a selection marker to isolated breast cancer CICs or head and neck CICs (HN-CICs) [29]. Therefore, we first utilized the in vitro cell-based aldehyde dehydrogenase (ALDH) activity assay system to screen for the active components from Antrodia cinnamomea mycelia extract (ACME) with effective reduction on ALDH activity of cells. Of the tested compounds purified from ACME, YMGKI-1 (a malic and succinic acid derivative (Figure 1(a))) significantly reduced the ALDH enzymatic activity of YMGKI-1-treated SAS cells in a dose-dependent manner (Figure 1(b)). We also found that YMGKI-1 diminished the ALDH enzyme activity in other human cancer cell lines (Supplementary Table 1). Next, treatment with YMGKI-1 also reduced the percentage of memGRP78+ cells in HNSCC cells, where memGRP78 has been used as a cell surface marker for isolation of HN-CICs (Figure 1(c)) [9]. In Figure 1(d), we showed that treatment of YMGKI-1 in OECMI cells caused the decreasing of the side population (SP) cells, which is also one of the characteristics of CICs [30]. To further evaluate the inhibitory effect of YMGKI-1 treatment on different cell lines including normal human oral keratinocytes (NHOKs), amniotic fluid stem cells-2 (AFSC-2), hematopoietic stem cell (HSC), parental SAS, and SAS-HN-CICs, the MTT cell proliferation assay was applied. Interestingly, we observed a massive decrease in viability of the SAS-HN-CICs but no significant inhibition on the other cells (Figure 1(e); **P < 0.01; ***P < 0.001). Hence, the previous findings suggest that YMGKI-1 might effectively and selectively target HN-CICs.

3.2. Reduction of Xenograft Tumor Growth on YMGKI-1-Treated HNSSC Cells. Existence of HN-CICs is critical for the tumor growth of HNSCC in vivo [5, 8, 9, 13]. In Figure 1, we showed that YMGKI-1 treatment effectively diminished the stemness properties of HN-CICs in HNSCCs. Therefore, we wanted to determine if YMGKI-1 treatment could attenuate the tumor initiating activity of HNSCC in vivo. As shown in Figure 2(a), nude mice injected with YMGKI-1 pre-treated SAS cells, a tumorigenic cell line, displayed significant reduction of the tumor growth as compared to those mice injected with untreated SAS cells (*P < 0.05; **P < 0.01). Next, whether YMGKI-1 could function as a therapeutic reagent on attenuating the tumor growth was examined as follow. We first subcutaneously implanted parental SAS cells into nude mice and allowed the tumors to be established. Mice bearing palpable tumor were then intraperitoneally injected with YMGKI-1 or DMSO (as control). Effectively, tumor-bearing mice receiving YMGKI-1 treatment afterward displayed reduced tumor size and the tumor growth curves in comparison with the control mice (Figure 2(b)) (*P < 0.05; **P < 0.01). In the meantime, we did not observe severe side effect but with little body weight loss from mice with YMGKI-1 treatment (data not shown).

3.3. Diminished Stemness Properties and Enhanced Differentiation Ability in YMGKI-1-Treated HN-CICs. To further evaluate whether treatment of YMGKI-1 would hinder the stemness properties of HN-CICs, we examined the following stemness properties of YMGKI-1-treated HN-CICs. We first observed that YMGKI-1 treatment significantly decreased the percentage of memGRP78+ or CD133+ cells in enriched SAS-HN-CICs (Figures 3(a) and 3(b)), where memGRP78+ or CD133+ cells have been demonstrated as CICs with higher stemness properties [5, 9]. Inversely, SAS-HN-CICs with the YMGKI-1 treatment displayed enhanced expression of epithelial differentiation marker, cytokeratin 18 (CK-18) (Figure 3(c)). Immunoblot analyses also showed that the expression of “cancer stemness” proteins (Oct-4, Nanog and Notch2) and GRP78 was diminished in YMGKI-1-treated SAS-HN-CICs (Figure 3(d)). In contrast, the YMGKI-1-treated SAS-HN-CICs displayed increased expression of epithelial-like protein (E-Cadherin) (Figure 3(d)). Together, YMGKI-1 treatment diminished the stemness properties but enhanced the differentiation ability of HN-CICs.

3.4. Reduction of Cell Survival and Malignancy of HN-CICs by YMGKI-1 Treatment In Vitro. Herein, we wanted to determine whether YMGKI-1 treatment affects cell survival and malignancy of HN-CICs or HNSCCs in vitro. Cells were first treated with YMGKI-1 and analyzed by flow cytometry, and the cells stained positively with propidium iodide (PI) were accounted as “dead” cells. Interestingly, YMGKI-1 treatment significantly induced cell death in SAS-HN-CICs whereas no significant cytotoxicity or morphological change was observed in YMGKI-1-treated parental SAS cells (Supplementary Figures SIA and SIB). Collectively, these results suggest that YMGKI-1 effectively suppresses the growth of SAS-HN-CICs but not parental SAS cells. Further, the in vitro
Figure 1: Continued.
malignancy of YMGKI-1-treated SAS-HN-CICs including matrigel invasion ability and anchorage independent growth was evaluated. As shown in Figures 3(b) and 3(c), the invasiveness and soft agar colony formation abilities of YMGKI-1 treated SAS-HN-CICs were significantly reduced.

3.5. Induction of Autophagic Cell Death in YMGKI-1-Treated HN-CICs. Due to the observation of induced cell death in YMGKI-1-treated HN-CICs (Supplementary Figures S1A and S1B), next, we wanted to examine whether the dying cells of YMGKI-1-treated HN-CICs were undergoing apoptosis. Apoptosis is designated as type I programmed cell death, and the apoptotic cells are determined by costaining with Annexin-V (AV) and propidium iodide (PI) [31]. When SAS-HN-CICs were cultivated with YMGKI-1 of 35 μg/mL, there was a slight decrease in early apoptotic cells (AV⁺/PI⁻) (Figure 5(a); untreated: 2.1% versus treated: 1.8%) but a significant increase in later apoptotic cell death (AV⁺/PI⁺) (Figure 5(a); untreated: 3.7% versus treated: 23.6%). Nevertheless, many YMGKI-1-treated cells could die from the other programmed cell death such as autophagy (untreated: (AV⁻/PI⁺: 6.7%) versus treated: (AV⁻/PI⁺: 40.8%)) (Figure 5(a)).

To further understand the molecular mechanism by which to cause the YMGKI-1 induced cell death in HN-CICs, we wanted to determine whether YMGKI-1 treatment induced autophagy in HN-CICs. Acidification of autolysosomes caused by lysosomes fusion with autophagosomes is regarded as an indication of cell autophagy, and autophagy is characterized by AVO formation, which can be detected and measured by acridine orange staining [32]. Using acridine orange staining, we showed progressive increasing in formation of AVO in a dose-dependent manner of YMGKI-1 treated HN-CICs (Figure 5(b)). Further, under the treatment with 10, 25, or 35 μg/mL concentrations of YMGKI-1, we detected that around 12%, 20%, or 43% of HN-CICs underwent autophagy by flow cytometry analyses, respectively (Figure 5(c)).

Autophagy is defined as type II programmed cell death and could be also determined by the induction of LC3-II [33]. Thus, we wanted to utilize the amount of LC3-II or the LC3-II/LC3-I ratio, which is positively correlated with the number of autophagosomes, to identify whether the YMGKI-1-induced dying cells were undergoing autophagy. As shown in Figure 5(d), an increase of LC3-II/LC3-I ratio in YMGKI-1 treated cells was observed in a dose-dependent manner. To further confirm that autophagic cell death was responsible for the cell death of YMGKI-1-treated HN-CICs, we evaluated the protective effect of 3-methyladenine (3-MA; a prototypic autophagy inhibitor [33]) in HN-CICs with combined treatment with YMGKI-1 and 3-MA. As expected, 3-MA cotreatment attenuated the cell death and the LC3-II accumulation induced by YMGKI-1 treatment in HN-CICs (Figures 5(e) and 5(f)).

We also examined the effects of YMGKI-1 treatment on the expression of apoptosis-related proteins in HN-CICs (Supplementary Figure S2A). HN-CICs treated with YMGKI-1 resulted in cleavage of caspase-3 and poly (ADP-ribose) polymerase (PARP) at 35 μg/mL but not at 10 or 25 μg/mL. Recent data support that the execution of apoptosis could be dependent on the occurrence of autophagy [33, 34]. In summary, our results suggest that YMGKI-1-treated HN-CICs were undergoing the activation of autophagic cell death by which to cause apoptosis.

3.6. Affected Signaling Pathways in YMGKI-1-Treated HN-CICs. The activation of mTOR-mediated signaling pathway
is known to inhibit autophagy. In addition, the mTOR pathway is activated for self-renewal, cell survival, and malignancy of CICs [35]. In addition, HER2 overexpression is reported to be able to increase the stem/progenitor cell population in malignant mammary cells. Increased levels of HER2 can activate the PI3K/Akt pathway which is related to self-renewal ability in stem cells [36]. Further, activation of phosphor-AMPK can inactivate the mTOR complex-1, inversely, to cause the activation of autophagy [37].

Herein, we examined the effect of YMGKI-1 on these signaling molecules by immunoblot analyses. As shown in Figures 6(a) and 6(b), YMGKI-1 treatment in HNCICs effectively decreased the expression level of phosphor-mTOR, HER2, phosphor-EGFR, Phosphatidylinositol 3-kinases (PI3K), phosphor-p44/42 MAPK (Thr202/Tyr204), and phosphor-AMPK but not phosphor-p38 MAPK in a dose-dependent manner. Additionally, cotreatment of 3-MA attenuated the expression of phospho-AMPK, which is
Figure 3: YMGKI-1 treatment diminishes stemness properties and enhances differentiation capability in HN-CICs. SAS-HN-CICs were treated with YMGKI-1 at different concentration for 24 hr, then stained with anti-GRP78 (a), anti-CD133 (b), or anti-CK18 (c) antibodies, respectively, and quantitated by flow cytometry. Data are means ± SD of triplicate samples from three experiments (***P < 0.005). Dead cells were excluded by gating on the propidium-iodide- (PI-) positive cell fraction. (d) Crude cell extract proteins of YMGKI-1-treated SAS-HN-CICs were collected, electrophores, and analyzed by immunoblotting against anti-Oct-4, anti-Nanog, anti-Notch2, anti-GRP78, anti-E-cadherin, or anti-GAPDH antibodies as indicated. The immunoactive signal of GAPDH protein of different crude cell extracts was referred as loading control. The same concentration (0.03%) of vehicle (DMSO) was added to all control groups.
Figure 4: YMGKI-1 treatment reduces cell malignancy of HN-CICs in vitro. (a) SAS-HN-CICs were treated with 0, 10, 25, and 35 µg/mL of YMGKI-1 for 24 hr, afterward, plated onto transwell, and analyzed as Materials and Methods. Dead cells were excluded by trypan blue dye. (b) SAS-HN-CICs were treated with 0, 10, 25, and 35 µg/mL of YMGKI-1 for 24 hr, afterward, plated onto soft agar for 12 day. The colony formation ability of previous cells was examined (see Section 2). Data are means ± SD of triplicate samples from three experiments (**P < 0.005). The same concentration (0.03%) of vehicle (DMSO) was added to all control groups.
Figure 5: YMGKI-1 treatment induces autophagic cell death in HN-CICs. (a) SAS-HN-CICs treated with 35 μg/mL of YMGKI-1 were costained with Annexin V and Propidium iodide (PI) and examined by flow cytometry. (b) SAS-HN-CICs were treated with YMGKI-1 at different concentrations for 24 hr, afterward, stained with acridine orange, and observed under a red filter fluorescence microscope. (c) Acridine orange (AO) positively stained cells of SAS-HN-CICs in (b) were quantitated by flow cytometric analysis (upper panel). The mean fluorescence intensity of AO stained cells was calculated (lower panel). Data are means ± SD of triplicate samples from three experiments (**P < 0.005). (d) Crude cell extract proteins of YMGKI-1-treated SAS-HN-CICs were collected, electrophorized, and further analyzed by immunoblotting against anti-LC3 or anti-GAPDH serum as indicated. (e) SAS-HN-CICs cells were either singly treated with 0, 25, or 35 μg/mL of YMGKI-1, or cotreated with 3-Methylamphetamine (3-MA), an autophagy inhibitor, for 24 hr, afterward, stained with propidium iodide (PI), and then examined by flow cytometry. Data are means ± SD of triplicate samples from three experiments (**P < 0.005). (f) Crude cell extract proteins of SAS-HN-CICs singly treated with YMGKI-1 or cotreated with 3-MA were isolated and analyzed by immunoblotting against anti-LC3 or anti-GAPDH serum as indicated. The immunoreactive signal of GAPDH protein of different crude cell extracts was referred as loading control. The same concentration (0.03%) of vehicle (DMSO) was added to all control groups.
Figure 6: YMGKI-1 treatment dysregulates the autophagy and diminishes the sphere formation ability in HN-CICs. (a) Crude cell extract proteins of YMGKI-1-treated SAS-HN-CICs were collected, electrophorized, and analyzed by immunoblotting against anti-LC3, anti-phosphor-mTOR, anti-HER2, PI3K, anti-phosphor-p44/p42 MAPK, anti-phosphor-p38, or anti-GAPDH serum as indicated. (b) Crude cell extract proteins of SAS-HN-CICs singly treated with YMGKI-1 or cotreated with 3-MA were isolated and analyzed by immunoblotting against anti-phosphor-AMPK or anti-GAPDH serum as indicated. (c) SAS-HN-CICs were treated with metformin (10 mM) or rapamycin (100 nM) for 96 hr. Consequently, immunoblotting analysis was performed by against anti-phosphor-AMPK, anti-phosphor-mTOR, and anti-LC3 or anti-GAPDH serum as indicated. The immunoactive signal of GAPDH protein of different crude cell extracts was referred as loading control. (d) Single cell suspension of SAS-HN-CICs was treated with YMGKI-1 or cotreated with 3-MA for 24 hr, and the sphere formation ability of YMGKI-1 or cotreated with 3-MA-treated HNCICs cells was examined. Arrows indicated the sphere cells. Data are means ± SD of triplicate samples from three experiments (** P < 0.005). The same concentration (0.03%) of vehicle (DMSO) was added to all control groups.
induced by YMGKI-1 treatment in HN-CICs (Figure 6(b)). It has been reported that AMPK activator like metformin or EGCG selectively kills cancer stem cells [38, 39]. To address the role of AMPK and mTOR on mediating HN-CICs through autophagy, we treated HN-CICs with AMPK activators metformin (Mf) or mTOR inhibitors rapamycin (Rapa), respectively. As shown in Figure 6(c), HN-CICs treated with metformin or rapamycin displayed slight induction of LC3-II along with upregulation of phosphor-AMPK or downregulation of phosphor-mTOR, respectively (Figure 6(c)). In the meantime, we did not observe significant cell death from HN-CICs under metformin or rapamycin treatment (Supplementary Figure S3B). These data suggest that YMGKI-1 treatment in HN-CICs simultaneously induced dysregulatory autophagic cell death through multiple mechanisms.

3.7 Attenuation of Inhibitory Effect of YMGKI-1 by Autophagy Repressor on Sphere Formation Ability of HN-CICs. To further investigate whether the “antistemness” properties of YMGKI-1 are autophagy dependent, the sphere formation ability of YMGKI-1-treated HN-CICs was analyzed. As shown in Figure 6(d), YMGKI-1 treatment significantly inhibited the sphere formation ability of HN-CICs. Further, cotreatment of 3-MA lessened the inhibitory effect of YMGKI-1 on sphere formation ability. The results indicated that YMGKI-1 treatment inhibited the stemness properties through autophagy activation in HN-CICs.

3.8. Autophagic Pathway and Putative Molecules Regulated by YMGKI-1. It has been known that cells undergoing epithelial-mesenchymal transition (EMT) can gain stem cell properties [40]. We also observed that YMGKI-1 treatment not only increased the expression of CK-18, an epithelial differentiation marker, but also reduced the expression of “cancer stemness” genes and markers in HN-CICs (Figures 1(b)–1(d), Figures 3(a)–3(d)). Intriguingly, Tseng et al. predict that Himanimide-C, a YMGKI-1 analogue, might be a potential inhibitor of cyclooxygenase-2 (COX-2) and 5-lipoxygenase (5-LOX) [7], which play important roles in EMT [40, 41]. Collectively, we hypothesize that YMGKI-1 might suppress EMT and stemness properties of HN-CICs by inhibiting 5-LOX or COX-2. Therefore, we proposed a molecular signaling pathway in HN-CICs responding to YMGKI-1 treatment (Figure 7). Overall, YMGKI-1 would directly or indirectly regulate the autophagic pathway mediated by mTOR and AMPK, in addition to the aforementioned putative COX-2 or 5-LOX molecules, by which YMGKI-1 regulates the stemness, tumorigenicity, differentiation ability, and cell death of HN-CICs. The future research to delineate the function of YMGKI-1 or YMGKI-1 derivatives on HN-CICs will benefit future cancer therapeutics.

4. Discussion

In this study, we utilized the in vitro cell-based ALDH activity assay system to screen drugs on targeting cancer initiating cells. YMGKI-1, a purified component of natural product, Antrodia cinnamomea mycelia extract, decreased ALDH activity in oral cancer and other cancer cell lines (Figure 1 and Supplementary Table 1). We also showed that YMGKI-1 treatment reduced the stemness properties of parental SAS cells (Figures 1(c) and 1(d)). In addition, we demonstrated that YMGKI-1 suppressed tumor growth in preventive and therapeutic model in vivo (Figure 2). Moreover, YMGKI-1 treatment in SAS-HN-CICs abrogated the in vitro malignancy and downregulated the stemness properties (Figures 3 and 4). Finally, YMGKI-1 was able to inhibit the growth of HN-CICs by dysregulatory autophagic cell death (Figure 5). Of note, coaddition of YMGKI-1 and autophagy inhibitor (3-MA) to HN-CICs attenuated the cell death, LC3-II accumulation, AMPK activation, and the morphological change of spheroid cells induced by YMGKI-1 treatment (Figure 6). All of these suggest that YMGKI-1 can function as a prospective drug for targeting CICs, and the future research to delineate the function of other YMGKI-1 analogues on HN-CICs will be of benefit for future cancer therapeutics.

Recently, the role of autophagy in the biology of stem cell and cancer initiating cell has been controversial. Cufí and colleagues report that autophagy is positively linked to the maintenance of breast cancer stem-like phenotype [42]. Conversely, others demonstrate that autophagy promotes the differentiation of glioma-initiating cells [17]. Herein, our results indicated that YMGKI-1 can not only induce autophagy but also downregulate the stemness properties and malignancy.
of HN-CICs (Figures 3 and 5). Although basal autophagic activity in some cells contributes to cytoprotective process against environmental stress [15], our findings suggest that the dysregulatory autophagy induced by YMGKI-1 treatment may effectively contribute to the enhanced cell death in HN-CICs but not in majority HNSCC cells (Figures 3(a) and 5 and Supplementary Figures S1 and S3). Further, we showed that YMGKI-1 induced autophagic cell death in HN-CICs through multiple molecular mechanisms including activation of AMPK and downregulation of PI3K-mTOR pathway (Figure 6). Of note, the aforementioned two signaling pathways center the regulation of autophagy. In addition, there are various signaling cascades that regulate mTOR including the HER2/PI3K (class I) pathway; inhibition of HER2/PI3K (class I) pathway has been shown to induce autophagy in cancer [43].

Under sufficient supply of growth factors and nutrients, the active mTORC1 stimulates growth-related processes such as protein translation, for example, by phosphorylation of S6K1 and 4E-BP, while simultaneously inhibiting self-consuming processes such as autophagy [44]. In opposite, the AMPK activity is enhanced when the intracellular ATP/AMP ratio is decreased. Further, AMPK activation induces autophagy through inhibition of mTORC1 [37]. Therefore, both AMPK and mTOR pathways play the crucial roles simultaneously in YMGKI-1-mediated autophagic HNSCC cell death. To further address the role of AMPK and mTOR on mediating HN-CICs through autophagy, we treated HN-CICs with AMPK activators metformin (Mf) or mTOR inhibitors rapamycin, respectively. As shown in Figure 6(c), HN-CICs treated with metformin or rapamycin displayed induction of autophagy. Nevertheless, we did not observe cell death from HN-CICs under single treatment with either metformin or rapamycin (Supplementary Figure S3B). Because of lack of induced cell death in HN-CICs by single treatment with metformin or rapamycin, it is worth note that YMGKI-1 induces dysregulatory autophagic cell death through multiple mechanisms. It will be of interest to determine how multiple mechanisms regulate autophagic cell death in HN-CICs.

Charafe-Jauffret et al. and Todaro et al. demonstrate that cancer stem cells highly express tumorigenic and metastatic activity [45, 46]. Although cancer stem cells may vary among different types of cancer, YMGKI-1 might have common effect on them. Other than HNSCC, here, we had tested the cytotoxic effect of YMGKI-1 treatment on colon, breast, and lung cancer cell lines. As shown in Supplementary Table 1, the more tumorigenic or metastatic cancer cell lines displayed higher reduction of ALDH1 cells under the YMGKI-1 treatment. Together, our data evidences that YMGKI-1 can significantly affect CICs population without discrepancy on different types of tumor.

Conventional anticancer treatments can often transiently shrink tumor volume instead of targeting or killing CICs, which lead to treatment failure, tumor recurrence, and patient death. Cancer initiating cells model suggests that tumors consist of CICs and differentiated cancer cells [3]. In the future, the main treatment strategy will be based on depletion of both the CICs pool and the residual majority cancer cells with combinatorial treatment including CIC-targeted anticancer drugs and conventional cancer therapeutic drugs [47]. Furthermore, in Figure 5(b), we observed that YMGKI-1 effectively inhibited the activation of HER2 but inversely enhanced the expression of phosphor-Src activity (data not shown). The Src kinase activity is closely related to cancerous malignancy [48]. Additionally, Zhang et al. discover that increased Src activation confers considerable Herceptin resistance in breast cancer cells and correlated with Herceptin resistance in breast cancer patients [49]. It is plausible that a combination with YMGKI-1 and Src kinase inhibitor will be a highly effective therapeutic protocol for head and neck cancer treatment. Further research effort with regard to the aforementioned phenomenon is needed in this area.

Together, our present research showed that YMGKI-1, a natural compound from ACME, can specifically and effectively target HN-CICs and diminish the CICs properties and activate dysregulatory autophagic cell death. Therefore, YMGKI-1 treatment might be a potential therapeutic target for HNSCC by eliminating CICs.

**Conflict of Interests**

No potential conflict of interests was disclosed.

**Authors’ Contribution**

Ching-Wen Chang, Chien-Chih Chen, and Meng-Ju Wu have equal contribution for the first authorship.

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