Induction of apoptosis by *Armillaria mellea* constituent armillarikin in human hepatocellular carcinoma

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Abstract: *Armillaria mellea* is a honey mushroom often used in the traditional Chinese medicine “Tianma”. Currently, this medicinal mushroom is also used as a dietary supplement in numerous Western and Eastern countries. Armillarikin was isolated from *A. mellea*, and we previously discovered that it induced cytotoxicity in human leukemia cells. In this study, we further investigated the cytotoxicity of armillarikin against liver and intrahepatic bile duct cancer cells. Armillarikin was cytotoxic against human hepatocellular carcinoma Huh7, HA22T, and HepG2 cells based on the 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulphophenyl)-2H-tetrazolium and alamarBlue® assays. Armillarikin treatment also induced the collapse of the mitochondrial transmembrane potential of these cells. Furthermore, armillarikin-induced apoptotic cell death was demonstrated by sub-G1 chromosomal DNA formation by using flow cytometry. In addition, the apoptosis was inhibited by the pan-caspase inhibitor, Z-VAD-fmk. Immunoblotting also revealed the armillarikin-induced activation of procaspase-3, -8, and -9 and upregulation of the apoptosis- and cell cycle arrest-related phospho-histones 2 and 3, respectively. Moreover, reactive oxygen species scavengers also inhibited the armillarikin-induced apoptosis in human hepatocellular carcinoma, suggesting that reactive oxygen species formation played an important role in the armillarikin-induced apoptosis of human hepatocellular carcinoma. In conclusion, our study indicates the potential of armillarikin as an effective agent for hepatoma or leukemia therapies.

Keywords: armillarikin, *Armillaria mellea* (*A. mellea*), hepatocellular carcinoma (HCC), apoptosis, reactive oxygen species (ROS)

Introduction

Armillaria mellea is a golden colored honey mushroom that belongs to the Tricholomataceae family of fungi. It is grown in numerous areas of the world, including the US. It is an edible mushroom that is used for its medicinal and health-promoting properties worldwide. For example, the fermented mycelia extracts are used as dietary supplement products. In Chinese herbal medicine, *A. mellea* is an important component of “Tianma” (“Tian-ma” or “Tien-ma”, literally translated as “heavenly hemp”) because it grows symbiotically with *Gastrodia elata* blume. The orchid *G. elata* relies on *A. mellea* to provide its nutrition and both have become common components of Tianma. In Chinese herbal medicine, *A. mellea* is indicated for the treatments of palsy, dizziness, hypertension, headache, insomnia, vertigo, and neurasthenia.

Over the past few decades, hepatocellular carcinoma (HCC, also called malignant hepatoma) has been one of the five most deadly cancers worldwide⁴ and is especially prevalent in less developed countries. For example, liver cancer was the second most

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deadly cancer in males in 2012,⁴ and the first or second most deadly cancer in a decade in Taiwan. HCC is more prevalent in Asian countries, but its incidence has also increased in Western countries due to chronic hepatitis or cirrhosis associated with hepatitis B virus, hepatitis C virus, alcoholic fatty liver disease, and other risk factors, such as aflatoxin, obesity, type II diabetes, and smoking.^{4,5} Current aggressive therapies include surgical treatment, chemotherapy, transhepatic arterial chemoembolization, radiation therapy, radiofrequency ablation, and targeted therapy with multiple drugs. However, effective cures for HCC are urgently needed because of the associated poor prognosis with a relatively low 1-year survival rate and less than 10% 5-year survival rate.^{6–8}

Several compounds found in *A. mellea* have potential antitumor or cytotoxicity effects, including armillarikin,^{2,9,10} armillaridin,^{3,11–13} other melleolides,^{9,12,13} and armamial.¹⁴ Other biological activities of *A. mellea* extracts or components include induction of maturation in human dendritic cells,¹⁵ upregulation of concanavalin A- or lipopolysaccharide-stimulated lymphocyte proliferation,¹⁶ antifungal activity, and antioxidant properties.¹⁷

Programmed cell death, such as apoptosis, autophagy, and necroptosis, is a common form of anticancer drug-induced cytotoxicity. In our previous study, we demonstrated reactive oxygen species (ROS)-mediated apoptosis of human leukemia cells induced by armillarikin.² Therefore, we speculated whether armillarikin would also be useful for hepatoma treatment and, here, demonstrated its cytotoxicity and apoptotic effects against HCC cells.

Materials and methods

Cell culture and treatment

The human HCC cell lines Huh7, HepG2, and HA22T were cultured in Dulbecco's Modified Eagle's Medium (Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% fetal calf serum (Hyclone, Logan, UT), nonessential amino acids (Gibco), and penicillin/streptomycin (Gibco) at 37°C in an atmosphere of 5% CO₂. The armillarikin was purified as previously described.^{2,9,18} In brief, the ethanol (EtOH) extract of *A. mellea* was separated by using silica gel column chromatography, which was eluted with *n*-hexane:ethyl acetate (EtOAc, 1:1). The final extract obtained was dissolved in dimethyl sulfoxide (DMSO, Sigma-Aldrich Co., St Louis, MO, USA) as a 1,000-fold stock solution. The armillarikin powder was stable at 4°C for at least 2 years as shown by the similar growth inhibition we observed after this period in the 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2*H*-tetrazolium, inner salt (MTS) viability

assays of the same cell line. For drug treatment, the HCC cell lines were treated with the vehicle DMSO or various concentrations of armillarikin for 1–3 days as indicated. For some experiments, the armillarikin-treated cells were also incubated in the presence or absence of 20 µM of the broad-spectrum caspase inhibitor, Z-VAD-fmk (R&D Systems, Inc., Minneapolis, MN, USA) or the antioxidant reagent, *N*-acetyl cysteine (NAC, 2.5 mM, Sigma-Aldrich Co., St Louis, MO, USA). The Z-VAD-fmk and NAC were dissolved in DMSO or water as 1,000- and 20-fold stock solutions, respectively.

Cell viability assay

The HCC cells were treated with 6.25–50 µM armillarikin for 24–72 hours and then the viability was measured by using the MTS assay (Promega Corporation, Fitchburg, WI, USA) following the manufacturer's instructions. The absorbance optical density values of the DMSO vehicle-treated control cells were considered as 100% activity.

Cell morphology and micronuclei observation

After incubation with armillarikin, the morphology of the HCC cells was photographed directly or after fixation with methanol and staining with Giemsa (Gibco). For the micronuclei observation, the cells were fixed with methanol and stained with Hoechst 33342 2.5 µg/mL (Thermo Fisher Scientific) as described in the manual of the fluorescence microscopy software (AxioVision, Zeiss, Oberkochen, Germany).

Flow cytometry

The HCC cells were treated with DMSO or armillarikin, collected, and then resuspended in hypotonic buffer (0.1% each of sodium citrate and Triton X-100) containing 5 µg/mL propidium iodide (Sigma) as described by Nicoletti et al¹⁹ to detect the apoptotic cells with hypodiploid chromosomal DNA (sub-G1 or DNA laddering) by using flow cytometry (fluorescence-activated cell sorting, FACSCalibur).²⁰ The percentage of the sub-G1 cells was calculated. To detect the collapsed mitochondrial transmembrane potential of the cells by using flow cytometry, they were mixed with 0.02 µM 3,3'-dihexyloxacarbocyanine iodide (Molecular Probes, Eugene, OR, USA) and analyzed according to the manufacturer's instructions. All the flow cytometric data were analyzed by using the FlowJo software (Tree Star, Inc., Ashland, OR, USA).

Immunoblotting

After treatment with armillarikin, the HCC cells were collected and dissolved in lysis buffer (Sigma-Aldrich) containing protease inhibitors (Roche Biochemicals, Indianapolis, IN, USA). After the debris had been removed and quantified, 50 µg of the cell lysates was separated by using 12.5% sodium dodecyl sulfate-polyacrylamide gel electrophoresis. Then, the separated proteins were immunoblotted with antibodies against caspase-8 (BD Biosciences, San Jose, CA, USA), caspase-3 (Active Motif), cleaved nuclear poly adenosine diphosphate-ribose polymerase (Cell Signaling, Beverly, MA, USA), caspase-9, apoptosis-related phospho-histone H2A.X (phospho-histone 2, Upstate), and phospho-histone 3 (Santa Cruz Biotechnology Inc., Dallas, TX, USA) while the loading control was actin (Sigma-Aldrich).

Results and discussion

Growth inhibition of HCC cells by armillarikin

We demonstrated the armillarikin-induced cytotoxicity and apoptosis in human leukemia cells in our previous study. To evaluate the cytotoxicity of armillarikin against HCC cells, we treated the human HCC cell lines, Huh7, HepG2, and HA22T cells with DMSO (vehicle control) or 6.25–50 µM armillarikin

for 1–3 days. Ethical approval was not required by the IRB of Chang Jung Christian University due to observations of cytotoxic activity of armillarikin in HCC cell lines in our study. The 0.1% DMSO showed no obvious effect on cell viability and morphology compared to the untreated control (data not shown). The cell viability was measured by using the MTS (Figure 1) and alamarBlue® (Invitrogen, Carlsbad, CA, USA) (Figure S1) assays. Armillarikin inhibited the cell growth of the three HCC cell lines in a time- and dose-dependent manner. The highest antiproliferative interval was 48–72 hours for most of the three HCC cells. To determine whether the decreased cell viability was induced by the cell death, we performed the following experiments. We observed the morphology of the HA22T and Huh7 HCC cells treated with armillarikin to confirm its cytotoxicity in these cells (Figure 2). In addition, armillarikin induced the collapse of the mitochondrial transmembrane potential in armillarikin-treated HepG2 and HA22T HCC cells (Figure 3), indicating induction of HCC cell death. These data suggest that armillarikin was cytotoxic against the HCC cells.

Induction of apoptosis by armillarikin in human hepatoma cells

Armillarikin caused HCC cell death and, therefore, we subsequently determined whether it also induced apoptosis.

Figure 1 Armillarikin inhibits hepatoma cell proliferation.

Notes: (A) Huh7, (B) HA22T, and (C) HepG2 HCC cells were treated with DMSO or armillarikin for 24–72 h. The relative number of viable cells was detected by MTS assay. OD values of DMSO-treated control cells were designated as 100% activity as a standard. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$ (Student's *t*-test) compared with DMSO control. Data are mean $\pm$ SD.

Abbreviations: DMSO, dimethyl sulfoxide; HCC, hepatocellular carcinoma; MTS, 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium; OD, optical density.

Figure 2 Morphology of armillarikin-treated cells.

Notes: (A) HA22T and (B) Huh7 cells were treated with DMSO or 50 μ M armillarikin for 3 days, and cell morphology was observed directly (A, upper panel) or after staining (A, lower panel and B). A: 100 $\times$, B: 400 $\times$.

Abbreviations: DMSO, dimethyl sulfoxide; MTP, mitochondrial transmembrane potential.

As shown in Figure 4, armillarikin induced a high percentage of hypoploid (or sub-G1) cells in the Huh7 and HepG2 HCC cell lines according to cell cycle analysis using flow cytometry, indicating that armillarikin was apoptotic against the majority of the cells. To further confirm this result, we examined the micronuclei morphology. As shown in Figure 4C, fragmented nuclei were observed in the immunofluorescence nuclei staining images after armillarikin treatment of Huh7 cells.

Caspases play key roles in the apoptosis process and, therefore, the armillarikin-induced apoptosis was further investigated using the broad-spectrum caspase inhibitor, Z-VAD-fmk. As shown in Figure 5, Z-VAD-fmk attenuated the growth inhibition induced by armillarikin in both HA22T and Huh7 HCC cells, indicating that caspase-associated death pathways mediated the armillarikin-induced apoptosis. To further confirm this result, we detected the cleavage of caspases and apoptosis-related protein expression by using Western blotting.

Figure 3 Armillarikin-induced loss of mitochondrial integrity in human hepatocellular carcinoma cells.

Notes: After DMSO, or 12.5 or 25 μM armillarikin treatment for 3 days, (A) HepG2 and (B) HA22T cells were stained with DiOC6(3). MTP of cells was analyzed using flow cytometry (right panel). Percentage of cells with low MTP is expressed as mean ± SD. *** $P < 0.001$ (Student's *t*-test) compared with DMSO control.

Abbreviations: DMSO, dimethyl sulfoxide; DiOC6(3), dihexyloxacarbocyanine iodide; HCC, hepatocellular carcinoma; MTP, Mitochondrial transmembrane potential; SD, standard deviation.

As shown in Figure 6, the full-length procaspase-9, -3, and -8 expression levels were decreased while the cleaved form of poly adenosine diphosphate-ribose polymerase (a caspase-3 substrate) and the cell cycle arrest-related phospho-histone 3 and DNA fragmentation-related phospho-H2A.X were increased following armillarikin treatment of human HCC cells.²¹ Furthermore, the vehicle control at 24, 48, and 72 hours confirmed there were no obvious differences in the proteins we detected (data not shown). These data confirmed the armillarikin-induced apoptosis of the HCC cells.

ROS-mediated apoptosis by armillarikin in human HCC cells

To further investigate the other signaling molecules likely involved in the armillarikin-induced apoptosis, we used the ROS scavenger, NAC. Cotreatment with NAC attenuated both armillarikin-induced apoptosis and growth inhibition in HA22T cells (Figure 7A–C). These data suggest that ROS

production occurs upstream of armillarikin-induced apoptosis in human HCC cells.

Armilarikin is known to have cytotoxic effects against HCC, leukemia, lung, and colon cancers,^{2,9,10} but our current findings clearly elucidated that the mechanism underlying the ROS-mediated apoptosis induced by armillarikin in HCC is similar to that operating in human leukemia cells.² Furthermore, this study demonstrates an example of the regulation of ROS-mediated apoptosis induced by DNA-damaging reagents,^{22,23} similar to that shown by natural products, such as marine algae and Baizhu extracts, and caffeic acid phenethyl ester from propolis.^{24–27} Similar to armillarikin, these compounds act as pro-oxidant agents for ROS production, which correlated with apoptosis induction. The detailed mechanism of the potential pro-oxidant effects of armillarikin will be elucidated in future studies. Because the production of ROS is deleterious to healthy cells, the induction of oxidative stress that leads to apoptosis is favorable in

Figure 4 (Continued)

Figure 4 Armilarikin-induced apoptotic cell death of human HCC cells.

Notes: (A and B) Huh7 and (B, right panel) HepG2 cells were treated with DMSO or armilarikin for 3 days, and the percentage of apoptotic cells (hypodiploid cells) with DNA ladders was detected using flow cytometry. $**P < 0.01$ and $***P < 0.001$ (Student's *t*-test) compared with DMSO control. (C) Armilarikin-induced micronuclei morphology. After Huh7 cells had been treated with armilarikin for 3 days, they were fixed and stained with Hoechst. Fluorescent and cell images were captured; arrow heads indicate micronuclei 400 $\times$.

Abbreviations: DMSO, dimethyl sulfoxide; HCC, hepatocellular carcinoma.

cancer tumor treatment.²⁸ Therefore, antioxidant/pro-oxidant compounds might have contradictory effects on normal and tumor cells.

Recently, we also published a report that armillaridin is an *A. mellea* constituent with both antitumor effects on human esophageal carcinoma in vivo³ and inhibitory effects on the differentiation of human activated macrophages.¹¹ Therefore,

the effects of both armillarikin and armillaridin support the potential usefulness of *A. mellea* as an antitumor medicinal mushroom worth future development.

Conclusion

In this study, we discovered that armillarikin inhibited the growth of human HCC Huh7, HepG2, and HA22T cells.

Figure 5 Z-VAD-fmk inhibited cell death in armilarikin-treated human HCC cells.

Notes: (A) HA22T and (B) Huh7 HCC cells were treated with DMSO or 6.25–50 μ M armilarikin with or without of 20 μ M Z-VAD-fmk (pan-caspase inhibitor) for 48 or 72 hours. The relative number of viable cells was detected using MTS assay. $*P < 0.05$, $**P < 0.01$, and $***P < 0.001$ (Student's *t*-test) compared with Z-VAD-fmk treatment.

Abbreviations: DMSO, dimethyl sulfoxide; HCC, hepatocellular carcinoma; MTS, 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium.

Figure 6 Armillarikin activated apoptosis-related caspases and other proteins in HCC cells.

Notes: (A) Huh7, (B) HepG2, and (C) HA22T cells were treated with DMSO or 12.5 or 25 μ M armillarikin for 24–72 hours as indicated. Cell lysates were collected, subjected to 12% SDS-PAGE, and analyzed using immunoblotting with antibodies against cleaved caspase-3 substrate, PARP (cleaved PARP), procaspase-3 (Cas 3), procaspase-8 (Cas 8), procaspase-9 (Cas 9), phospho-histone 3 (p-His 3), apoptosis-related phospho-histone 2 (p-His 2), and β -actin (loading control). 200 $\times$ magnification.

Abbreviations: ADP, poly adenosine diphosphate; Cas 3, procaspase-3; Cas 8, procaspase-8; DMSO, dimethyl sulfoxide; HCC, hepatocellular carcinoma; p-His 2, phospho-histone 2; p-His 3, phospho-histone 3; SDS-PAGE, sodium dodecyl sulfate-polyacrylamide gel electrophoresis; PARP, poly ADP ribose polymerase.

This phenomenon was accompanied by cell death and the collapse of the mitochondria due to loss of mitochondrial transmembrane potential after armillarikin treatment. In addition, we found a high percentage of hypodiploid cells (with sub-G1 DNA), detected by the DNA content determination. We further demonstrated the induction of apoptosis by examining the micronuclei, and this was further confirmed by the interference in the apoptotic effect induced by the broad caspase inhibitor, Z-VAD-fmk. Moreover, armillarikin-induced apoptosis was reconfirmed by

the immunoblotting of apoptosis-related proteins; cleavage of procaspase-9, -3, and -8; increased cleavage of the caspases 3 substrate, poly adenosine diphosphate-ribose polymerase; and an increase in the cell cycle arrest-related phospho-histone 3 and phosphorylation of the apoptotic marker, H2A.X phosphorylation.²¹ Finally, apoptosis induction by armillarikin was mediated by ROS, as the ROS scavenger, NAC, hindered both the high percentage of hypodiploid cells and growth inhibitory effect of armillarikin.

Figure 7 Human HCC cells were protected against armillarikin-induced cytotoxicity by ROS scavenger NAC.

Notes: (A and B) HA22T, (C) Huh7, and HepG2 cells were treated with (A and B) dimethyl sulfoxide (DMSO) or 25 μM or (C) 12.5–50 μM armillarikin with or without 2.5 mM NAC for 48 hours. Then, (A and B) percentage of cells with hypodiploid DNA content and (C) relative number of viable cells were detected and calculated using (A and B) flow cytometry and (C) MTS assay, respectively. OD values of DMSO-treated control cells were designated as 100% activity as a standard. *, $P < 0.05$ and **, $P < 0.01$ (Student's t-test) compared with Armillarikin plus NAC.

Abbreviations: DMSO, dimethyl sulfoxide; HCC, hepatocellular carcinoma; ROS, reactive oxygen species; MTS, 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium; NAC, N-acetyl cysteine.

In conclusion, the armillarikin-induced apoptosis was mediated by ROS and accompanied by mitochondrial collapse and activation of caspase-8 and -3 in HCC cells. Further experiments need to be conducted to determine the potential antitumor effects of armillarikin. Our study demonstrated that armillarikin may be considered as a potential candidate for further development as a therapy or adjuvant treatment for HCC.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary material

Figure S1 Armillarikin inhibits hepatoma cell proliferation.

Notes: (A) Huh7, (B) HA22T, and (C) HepG2 HCC cells were treated with 0.1% DMSO (vehicle), or 6.25–50 μM armillarikin dissolved in DMSO for 24–72 hours. The relative number of viable cells was detected using alamarBlue® assay. Fluorescence intensity values of DMSO-treated control cells were designated 100% activity as a standard. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$ (Student's *t*-test) compared with DMSO control. Fluorescence intensity values of DMSO-treated control cells were designated 100% activity as a standard. Data are mean $\pm$ SD.

Abbreviations: DMSO, dimethyl sulfoxide; HCC, hepatocellular carcinoma.

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