Cytotoxic Activities of Naturally Occurring Oleanane-, Ursane-, and Lupane-type Triterpenes on HepG2 and AGS Cells

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ABSTRACT
Background: It is well known that the naturally occurring modified triterpenes in plants have a wide diversity of chemical structures and biological functions. The lupane-, oleanane-, and ursane-type triterpenes are the three major members of natural triterpenes with a wide range of biological properties. A systematic approach is necessary to review their structures and biological activities according to the backbones and the different substituents. Objective: Thirty lupane-(L1-7), oleanane-(O1-14), and ursane-type (U1-9) triterpenes with structural diversity were examined to evaluate their cytotoxic activities against two cancer cell lines, human hepatocellular carcinoma (HepG2) and AGS cells. Materials and Methods: They were isolated from Hedera helix, Juglans sinensis, and Pulsatilla koreana using a series of column chromatography methods and were treated to evaluate their cytotoxic activities against HepG2 and AGS human gastric adenocarcinoma cell. Further, two triterpenes showing the most potent activities were subjected to the apoptotic screening assay using flow cytometry. Results: The polar groups, such as an oxo group at C-1, a free hydroxyl at C-2, C-3, or C-23, and a carboxylic moiety at C-28, as well as the type of backbone, explicitly increased the cytotoxic activity on two cancer cells. O5 and U5 showed significantly the potent cytotoxic activity in comparison to other glycosidic triterpenes. In annexin-V/propidium iodide (PI) staining assay, the percentage of late apoptosis (annexin-V/PI+) 12 and 24 h after treatment with O5 and U5 at 25 μM increased from 14.5% to 93.1% and from 46.4% to 49.1%, respectively, in AGS cells. The cytotoxicity induced by O5 showed a significant difference compared to U5 for 12 and 24 h.

Conclusion: In the study, we can suggest the potent moieties which influence their cytotoxic activities against two cancer cells. The polar groups at C-1, C-2, C-3, C-23, and C-28 and the linkage of sugar moieties influenced the different cytotoxic activities.

Key words: Flow cytometry, lupane, oleanane, triterpene, ursane

SUMMARY
• Thirty naturally occurring oleanane-, ursane-, and lupane-type triterpenes were isolated from Hedera helix, Juglans sinensis, and Pulsatilla koreana

INTRODUCTION
Triterpenes are a member of terpenes, which are structurally diverse class of natural compounds derived from the combination of C5 isoprene units, and have been abundantly found in the plant kingdom. They mainly exist in the cyclized form of squalene (30 carbons) which are acyclic hydrocarbons formed by two units of farnesyl diphosphate. Triterpenes are precursors of steroids in both plants and animals which have a wide range of pharmacological properties, such as anti-inflammatory, anti-angiogenesis, anti-oxidative, pro-apoptotic, and re-differentiation effects. Especially, lupane-, oleanane-, and ursane-type triterpenes showed the cytotoxicity against various cancer cell lines and have been considered as promising anti-cancer agents. Their cytotoxic activity on cancer cells is significantly affected by the type of basic backbone and the present number and position of derivatives, such as alkyl, hydroxyl, carboxyl, and amino acids. Among these, triterpene glycosides (i.e., triterpenoidal saponins) occurring by the combination of sugar residues to basic backbone also have significant cytotoxic effects on cancer cells.

To date, it has been mainly demonstrated the structure–activity relationship between synthesized triterpene derivatives on cancer cells. We evaluated the
cytotoxic activities of naturally occurring lupane-, oleanane-, and ursane-type triterpenes isolated from *Juglans sinensis* Dode (Juglandaceae),* Pulsatilla koreana* Nakai (Ranunculaceae),* and Hedera helix* L. (Araliaceae) against human hepatocellular carcinoma (HepG2) and AGS human gastric adenocarcinoma cell. We tried to suggest structurally some key determinants in the structure of triterpene derivatives on the cytotoxicity against cancer cell lines, and also explore their preliminary mechanisms through inhibiting cell cycle and inducing apoptosis.

**MATERIALS AND METHODS**

**General**

Compounds L1-7, O1-14, and U1-9 were isolated from *J. sinensis* Dode (Juglandaceae) (L1, O1-4, O10-14, and U1-9),* P. koreana* Nakai (Ranunculaceae) (L2-7, O6, and O9),* and *H. helix* L. (O5, O7, and O8) using repeated column chromatography and high-performance liquid chromatography. The structures of these compounds were unequivocally determined by diverse spectroscopic analyses, such as 1D, 2D nuclear magnetic resonance (NMR) experiments, mass spectrometry analyses, as well as by comparison with the literature for the known compounds. The 1H and 13C NMR, 1H-1H correlation spectroscopy, Heteronuclear single quantum coherence spectroscopy (HSQC), and heteronuclear multiple bond correlation spectra were recorded on a Bruker AMX 500 (Bruker BioSpin GmbH, Karlsruhe, Germany) or 500 (Bruker BioSpin GmbH, Karlsruhe, Germany) spectrometer in pyridine-d$_5$. High- and low-resolution FABMS (Fast Atom Bombardment Mass Spectrometry) were obtained on a JEOL JMS-AX505WA. Detection of DNA cycle and apoptosis on hepatic stellate cells-T6 cells was conducted by flow cytometry (BD Biosciences, FACSCalibur, Franklin Lakes, NJ, USA).

**Cell lines**

HepG2 and AGS (human gastric adenoma carcinoma cells) cell lines were purchased from ATCC (Manassas, VA, USA). HepG2 and AGS cells were maintained in Dulbecco's modiﬁed eagle medium and Roswell Park Memorial Institute 1640 medium, respectively, supplemented with 10% (v/v) fetal bovine serum, penicillin (100 U/mL), and streptomycin (100 µg/mL). They were incubated in a humidified atmosphere of 5% CO$_2$ gas at 37°C.

**Cytotoxicity assay**

The cytotoxicity of L1-7, O1-14, and U1-9 against two cancer cell lines was evaluated by modified 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. All samples were dissolved in dimethyl sulfoxide (final concentration, 0.1%) and diluted in distilled water. Cells were seeded in 96-well plates at a density of 3 x 10$^4$ cells/well for the cytotoxicity assay. After 24 h incubation, serum-free medium was changed and two cell lines were treated with vehicle or L1-7, O1-14, and U1-9 at the concentration of 100 µM for 24 or 48 h and incubated with 2 mg/mL of MTT for 2 h. Reduction of MTT to formazan was assessed in an ELISA plate reader at 540 nm. Doxorubicin (Sigma-Aldrich, St. Louis, MO, USA) was used as a positive control. Cell viability rate was calculated as the percentage of MTT absorption as follows: % survival = (mean experimental absorbance/mean control absorbance) x 100. Data were expressed as the mean of three independent experiments.

**Flow cytometry for analyzing apoptosis and caspase-3 activity**

AGS cells were seeded in 6-well plates at a density of 6 x 10$^4$ cells/well for measuring the DNA cycle, apoptosis, and caspase-3 using flow cytometry. After 24 h, cells were treated with O5 and U5 at 10 and 25 µM, respectively. After 12 or 24 h, cells were trypsinized, washed twice with PBS buffer, and centrifuged at room temperature. For evaluating the DNA cycle, cell pellets were suspended in ice-cold 70% ethanol for fixation of the stage of DNA cycle at 4°C. After overnight, cells were centrifuged and resuspended in 500 µL of PI/RNASE staining buffer (BD Pharmingen, Franklin Lakes, NJ, USA), incubated at room temperature for 30 min, and analyzed. For measuring apoptosis using a fluorescein isothiocyanate (FITC)-Annexin V apoptosis detection kit (BD Pharmingen, Franklin Lakes, NJ, USA), cells were stained with annexin V and PI, subsequently, according to the manufacturer's manual, and measured by flow cytometry. The alteration of caspase-3 activity by O5 and U5 was carried out according to the manufacturer's manual (FITC Active Caspase-3 kit, BD Pharmingen, Franklin Lakes, NJ, USA). All experiments were performed by flow cytometry (BD Biosciences, FACSCalibur, Franklin Lakes, NJ, USA).

**Statistical analysis**

The evaluation of statistical significance was determined by one-way ANOVA test, with P < 0.001 and P < 0.01 considered to be statistically significant.

**RESULTS AND DISCUSSION**

**Cytotoxic activities of 30 triterpenes against HepG2 and AGS cell lines**

We evaluated the cytotoxicity of 30 triterpenes derivatives including lupane-(L1-7), oleanane-(O1-14), and ursane-type (U1-9) against HepG2 and AGS cell lines by MTT cell viability assay in 24 and 48 h. All triterpenes

| No. | Lupane-type | Oleanane-type | Ursane-type |
|-----|-------------|---------------|-------------|
| L1  | Lupeol      | Oleanolic acid| U1          |
| L2  | Anemoside A | Arjunolic acid| U2          |
| L3  | 23-hydroxy-3β-[(O-α-L-rhamnopyranosyl-(1→2)-α-L-arabinopyranosyl)oxy]lup-20(29)-en-28-oic acid 28-Oβ-D-glucopyranosyl ester | 1-oxy-3β, 23-dihydroxy-olean-12-en-28-oic acid | U3          |
| L4  | Anemoside B | 2α, 3β, 23-trihydroxy-olean-12-en-28-oic acid 28-Oβ-D-glucopyranoside | 2α, 3α, 23-trihydroxy-urs-12-en-28-oic acid |
| L5  | Pulsatilloside E | 1-oxy-3β-hydroxy-olean-12-en-28-oic acid 28-Oβ-D-glucopyranoside | 3β, 23-dihydroxy-urs-12-en-28-oic acid |
| L6  | Cusssasopan C | Virgatic acid | 3β, 22a-dihydroxy-urs-12-en-28-oic acid |
| L7  | 3β-[(O-α-L-rhamnopyranosyl-(1→2)-O-β-D-glucopyranosyl-(1→4)]-α-L-arabinopyranosyl)oxy]lup-20(29)-en-28-oic acid 28-O-α-L-rhamnopyranosyl- (1→4)-O-β-D-glucopyranosyl-(1→6)-β-D-glucopyranosyl ester | Hederagenin 28-O-β-D-glucopyranoside | 1-oxy-3β-hydroxy-olean-18-ene |
| L8  | Hederagenin 28-O-β-D-glucopyranoside | Loniceria japonica saponin 9 | Ursolic acid |
| L9  | Pulsatilloside C | Pulsatilla saponin H | 23-hydroxyursolic acid |
| L10 | Pulsatilla saponin | Hederacide C | Corosolic acid |
| L11 | α-hederin | 2α, 3α, 23-trihydroxy-urs-12-en-28-oic acid | 3-oxy-23-hydroxy-urs-12-en-28-oic acid |
| L12 | Hederacide C | 3β, 23-dihydroxy-urs-12-en-28-oic acid 28-Oβ-D-glucopyranoside | 3β, 22a-dihydroxy-urs-12-en-28-oic acid |
| L13 | 3β-acetoxy-olean-18-en-28-oic acid | 2α, 3α, 23a-trihydroxy-urs-12-en-28-oic acid | Asiatic acid |
| L14 | 1-oxy-3β-hydroxy-olean-18-ene | 2α, 3α, 23a-trihydroxy-urs-12-en-28-oic acid 28-Oβ-D-glucopyranoside | 28-Oβ-D-glucopyranoside |
were isolated from *J. sinensis* Dode (Juglandaceae) (L1, O1-4, O10-14, and U1-9), *P. koreana* Nakai (Ranunculaceae) (L2-7, O6, and O9), and *H. indica* L. (O5, O7, and O8). Their names and structures are listed and provided in Table 1 and Figure 1, respectively.

### Relationship of structures and activities of 30 triterpenes

The cytotoxic activity of L1-7, O1-14, and U1-U9 at the concentration of 100 μM against HepG2 and AGS cells for 24 and 48 h was plotted in Figure 2. In lupane-type triterpenes (L1-7), L2 including an O-linkage of two glycosides at C-3, a carboxylic group at C-17, and a hydroxyl group at C-23 showed the most significant cytotoxic activity than other lupane-type triterpenes against two cancer cell lines. The substituents of sugar moieties at C-28 unambiguously decreased the cytotoxicity against cancer cells (L2 vs. L3-7). In addition, L1 with a methyl group instead of a carboxylic group at C-17 was essential for cytotoxic activity.

We investigated the changes in progress of apoptosis induced by O5 (IC_{50} 7.6 μM in AGS cells and 15.2 μM in HepG2 cells) and U5 (IC_{50} 11.3 μM in AGS cells and 24.5 μM in HepG2 cells), which showed the most significant inhibitory activity against two cancer cells using the flow cytometry [Figure 3]. While the structure of U5 includes all the factors, such as ursane-type aglycone, a free hydroxyl group at C-2, C-3, and C-23, and a free carboxylic group at C-28 (O3, O4, O7-9, and O12) showed no potent cytotoxic activities compared to those with a free carboxylic acid (O1, O2, O5, O10, O11, and O13). Interestingly, introduction of sugar moiety at C-3 did not influence the loss of cytotoxic activity (O5). The presence of a hydroxyl moiety at C-23 could significantly increase cell growth inhibition (O10 vs. O11). In terms of ursane-type triterpenes, the cytotoxic results of HepG2 and AGS cell lines treated with U1-9 generally showed more potent inhibition than lupane- and oleanane-type triterpenes on cell proliferation. In addition, some modifications, such as hydroxylation at C-2 (U3 vs. U5 and U4 vs. U6) or C-22 (U7 vs. U8), oxidation from hydroxyl to oxo at C-3 (U4 vs. U9), and glycosylation at C-28 (U4 vs. U7), did not significantly change the cell viability. However, the configuration of hydroxyl group at C-3 had an important influence on the anti-tumor activity. U1 and U2 which have α-hydroxyl group at C-3 showed no cytotoxic activity compared to others which have β-hydroxyl group (U3-9). On the basis of the above results, it was suggested that a free hydroxyl group at C-2, C-3, and C-23, and a free carboxylic group at C-28 were essential for the cell growth inhibition, and the addition of sugar moiety at C-3 or C-28 reduced the cytotoxic effect as the length of glycosides was increased. In comparison between oleanane- and ursane-type triterpenes having same modifications, ursane-type triterpenes had slightly more potent cytotoxicity than oleanane-type triterpenes (U3 vs. O1, U6 vs. O2, and U7 vs. O3) as reported in the literature.\(^{[17-20]}\)

### Apoptotic activities of O5 and U5 against HepG2 and AGS cell lines

We investigated the changes in progress of apoptosis induced by O5 (IC_{50} 7.6 μM in AGS cells and 15.2 μM in HepG2 cells) and U5 (IC_{50} 11.3 μM in AGS cells and 24.5 μM in HepG2 cells), which showed the most significant inhibitory activity against two cancer cells using the flow cytometry [Figure 3]. While the structure of U5 includes all the factors, such as ursane-type aglycone, a free hydroxyl group at C-2, C-3, and C-23, and a free carboxylic group at

### Table 1: Structures of L1-L7, O1-O14, and U1-U9

| No. | R_1 | R_2 | R_3 | R_4 |
|-----|-----|-----|-----|-----|
| L1  | H   | CH_3 |     |     |
| L2  | α‑Ara_2‑α‑Rha | OH | COOH |     |
| L3  | α‑Ara_2‑α‑Rha | OH | COO‑β‑Glc |     |
| L4  | α‑Ara_2‑α‑Rha | OH | COO‑β‑Glc_2‑β‑Glc_4‑α‑Rha |     |
| L5  | α‑Ara_2‑α‑Rha | H  | COO‑β‑Glc_2‑β‑Glc_4‑α‑Rha | α‑Rha |
| L6  | α‑Ara_4‑α‑Rha | H  | COO‑β‑Glc_2‑β‑Glc_4‑β‑Glc_6‑α‑Rha | α‑Rha |
| L7  | α‑Ara_4‑α‑Rha | OH | COO‑β‑Glc_2‑β‑Glc_4‑β‑Glc_6‑α‑Rha | α‑Rha |

Ara: L‑arabinopyranose; Rha: L‑rhamnopyranoside; Glc: D‑glucopyranose

| No. | R_1 | R_2 | R_3 | R_4 |
|-----|-----|-----|-----|-----|
| O1  | OH  | H   | H   | H   |
| O2  | OH  | OH  | H   | OH  |
| O3  | OH  | OH  | β‑Glc | H  |
| O4  | OH  | OH  | β‑Glc | OH  |
| O5  | α‑Ara_2‑α‑Rha | OH | H   | H   |
| O6  | α‑Ara_2‑α‑Rha | OH | β‑Glc | H   |
| O7  | α‑Ara_2‑α‑Rha | OH | β‑Glc_2‑β‑Glc_4‑α‑Rha | H  |
| O8  | α‑Ara_2‑α‑Rha | H  | β‑Glc_2‑β‑Glc_4‑α‑Rha | H  |
| O9  | α‑Ara_2‑α‑Rha | OH | β‑Glc_2‑β‑Glc_4‑β‑Glc_6‑α‑Rha | H  |

Ara: L‑arabinopyranose; Rha: L‑rhamnopyranoside; Glc: D‑glucopyranose

| No. | R_1 | R_2 |
|-----|-----|-----|
| O10 | H   | H   |
| O11 | OH  | H   |
| O12 | OH  | β‑Glc |

Glc: D‑glucopyranose

Figure 1: (a-f) Structures of L1-L7, O1-O14, and U1-U9.
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C-28, which essentially influence the cell viability against cancer cells in the cytotoxic assay, O5 as oleanane-type has not only a hydroxyl moiety at C-23 and a free carboxylic acid at C-28, but also two sugars at C-3. O5 and U5 at concentrations of 10 and 25 μM were treated in AGS cells for 12 and 24 h. The annexin-V/PI double staining results described that two compounds, O5 and U5, showed different aspects on the stage of apoptosis in AGS cells. When treated with 10 and 25 μM for 12 h incubation, U5 had much more potent cytotoxicity than O5 (Figure 3; b1 vs. d1 and c1 vs. e1). While O5 at 10 μM for 24 h did not induce the significant apoptosis compared to control group, the population of the apoptotic cells treated with O5 at 25 μM for 24 h in late apoptosis stage (annexin-V+/PI+) was dramatically increased to about 93.1% compared to U5 at the same concentration (49.1%). Our finding suggested that the delayed cytotoxicity of O5 might arise from the alteration of cell permeability by two glycosides at C-3 in structure. In the case of U5, the percentage of late apoptotic cells (annexin-V+/PI+) treated with 10 μM has doubled from 20.9% in 12 h to 43.6% in 24 h, while the number of late apoptotic cells induced by 25 μM showed no significant change as time passed (46.4% in 12 h and 49.1% in 24 h).

Changes of caspase-3 activity by O5 and U5 in AGS cell line

To further understand the mechanism of O5 and U5-induced apoptosis, we evaluated the caspase-3 activity that plays a pivotal role in the process of apoptosis in AGS cells. O5 and U5 gradually increased the percentage of the activated caspase-3 activity in a concentration-dependent manner, when the cells were incubated with samples for 24 h [Figure 4]. The percentages of the activated caspase-3 treated with O5 and U5 compared to those of the control cells increased gradually with increasing the concentrations from 10 (O5; 41.1% and U5; 45.0%) to 25 μM (O5; 56.6% and U5; 55.2%). These results suggested that active caspase-3 played an important role in executing apoptosis by O5 and U5 in AGS cells.

CONCLUSION

In the present study, we examined the cytotoxic activities of 30 lupane-, oleanane-, and ursane-type triterpenes against HepG2 and AGS cell lines. The types of backbones and simple structural modifications, such as hydroxylation and glycosylation, significantly influenced their biological activities. O5, an oleanolic acid derivative with hydroxyl group at C-23 and two sugar groups at C-3 in structure, was the most potent apoptotic activities on the two cancer cell lines via caspase-3 regulation. Although further studies are needed to clarify the mechanism related with apoptosis by O5 and U5 on the inhibition of cancer cells proliferation, the structure–activity relationship of 30 triterpenes including O5 and U5 might give an idea for developing the therapeutic agents consisting of a triterpenes-concentrated preparation.

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Conflicts of interest
There are no conflicts of interest.

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Figure 4: The flow cytometry analysis of O5 and U5-induced caspase-3 activity in AGS cells. The cells were incubated for 24 h with vehicle (control, white color), O5 or U5 at the indicated concentration (red color). Values are represented as percentage. Data are representative of three independent experiments.