Synthesis and Pro-Apoptotic Activity of Novel Glycyrrhetinic Acid Derivatives

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Introduction

Organic molecules synthesized by plants constitute a rich reservoir of biologically active compounds. For centuries extracts from various plants have been extensively used in traditional medicines for the treatment of a wide variety of human ailments; even today, many cultures still employ them directly for medicinal purposes.[1–4] Among the classes of recognized therapeutically useful products, pentacyclic triterpenoids have been studied intensively for their diverse biological, pharmacological, and medicinal activities, which are similar to those of retinoids and steroids.[5, 6] However, these triterpenoids exhibit only weak effects on the biological activity of their molecular targets; therefore these compounds have been used as building blocks for the synthesis of more active analogues.[6]

Oleanolic acid, an abundantly occurring triterpene, has been converted into 2-cyano-3,12-dioxoolean-9(11),1(2)-dien-30-oic acid (CDDO) and other structurally related analogues (CDDO-Me, CDDO-Im, CDDO-CN; Scheme 1 A). All of these synthetic derivatives were reported to display various bioactivities: cytoprotection, cancer cell growth inhibition, apoptosis induction, and inhibition of the production of NO induced by INF-γ in mouse macrophages.[7–12] CDDO and CDDO-Me are currently in clinical trials for cancer treatment, and have been shown to effectively suppress the growth of a broad spectrum of solid and hematologic cancer cell types, both in vitro and in mouse models bearing xenografted human tumors.[8–12] During the development of CDDO, it was found that the 2-cyano-1-en-3-one in ring A, and the 9(11)-en-12-one in ring C are essential for the biological activity of CDDO and its analogues.[13–15]

18β-H-Glycyrrhetinic acid (Scheme 1 B), the aglycon of glycyrrhizin, is abundant in licorice root (Glycyrrhiza glabra and Glycyrrhiza uralensis Fischer). The glycyrrhizin content in triterpene extracts from licorice root amounts to 90 %. Recent reviews have described the wide spectrum of glycyrrhetinic acid bioactivity, such as anti-inflammatory, antiviral, hepatoprotective, antitumor, and immunomodulatory activities.[16, 17] Several studies have reported that glycyrrhizin and glycyrrhetinic acid have moderate cytotoxic and apoptotic effects on cancer cells, although most reported only moderate or low potency.

Scheme 1. A) Structure of CDDO and its structurally related analogues. B) Structure of glycyrrhetinic acid.

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In attempts to prepare more-potent analogues of glycyrrhetinic acid, we synthesized compounds 2–12, similar to CDDO-Me (1), by introducing modification at both rings A and C (Scheme 1A). We investigated the effects of the novel derivatives on the growth of human cancer cells, and we identified methyl 2-cyano-3,12-dioxo-18βH-olean-9(11),1(2)-dien-30-oate 12 as a compound displaying significant antiproliferative activity toward cancer cells; the other glycyrrhetinic-acid derivatives did not display this activity. We compared 12 and CDDO-Me on several cell lines under the same conditions, and we showed that IC₅₀ was lower for 12 than CDDO-Me for all cell lines. Compound 12 induced cell-cycle arrest, the translocation of phosphatidylserine to the cell surface, and fragmentation of the nucleus. It also caused a dramatic dissipation of the mitochondrial potential, and induced activation of the caspase cascade; these effects were more pronounced for 12 than for CDDO-Me. The data indicate that 12 induces the death of cancer cells by the intrinsic caspase-dependent apoptosis pathway.

Results and Discussion

Chemical synthesis

The reaction sequence to introduce the 2-cyano-1-en-3-one and 9(11)-en-12-one in A and C rings of glycyrrhetinic acid is shown in Scheme 2.

18β-Glycyrrhetinic acid acetate 2, obtained from a licorice extract, was used as the starting material. Compound 2 was esterified at 0–8°C with ethereal diazomethane to give methyl glycyrrhetinate acetate 3, which was reduced by Zn/HCl in dioxane at 5–10°C. The resulting methyl ester of 11-deoxoglycyrrhetinic acid acetate 4 was converted into 12-oxo derivative 5 by treating with hydrogen peroxide in acetic acid at 80°C. The formation of the 9,11-double bond was achieved by bromination–dehydrobromination of ketone 5 with bromine in acetic acid at 80°C. Finally the 9(11)-en-12-one moiety in the C ring was obtained. Deprotection of acetate group by KOH in methanol (reflux) freed the 3-hydroxy group, then Jones oxidation gave the ketone 8. Subsequent formylation at C₂ was performed by condensation with HCO₂Et/NaOMe in benzene, and the resulting hydroxymethylene derivative 9 was cyclized into isoxazole 10 by reacting with hydroxylamine hydrochloride in aqueous ethanol (reflux). Opening of the isoxazole ring at the N–O bond was promoted by NaOMe routinely to deliver the 2-cyano group in 11. The new 1,2 double bond was formed by dehydrogenation with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) in benzene (reflux) to complete the synthesis of the 2-cyano-1-en-3-one moiety in A ring in 12.

It should be noted that the synthesis scheme for our end-product 12 has been described by Chadalapaka et al.[18] Our investigations were conducted independently and in parallel. In addition to the synthesis scheme, a detailed description of the synthesis and physicochemical properties of the end product (and of the intermediates) is presented in this work (see the Experimental Section).

Biological studies

Cell-viability inhibition: Inhibitors of cell growth are potentially useful as chemopreventive and chemotherapeutic agents. The
in vitro cytotoxicity of the novel derivatives of glycyrrhetinic acid against human epidermoid cancer cell-line KB-3-1 was determined by using the MTT assay, a colorimetric technique for the determination of cell viability which was developed for the initial stages of drug screening. The assay quantifies the reduction of the yellow tetrazole 3-(4,5-Dimethylthiazol-2-yl)-2,5-di-phenyltetrazolium bromide (MTT) to its purple formazan derivative by mitochondrial dehydrogenase. It assumes cell viability to be proportional to the production of formazan, and thus low IC$_{50}$ values imply high cytotoxicity or antiproliferation activity.

Glycyrrhetinic acid, 7, 8, 9, 10 and 12 were tested; the other derivatives were found to be insoluble in dimethylsulfoxide (DMSO). Cells were exposed to the compounds for 24 h and then assayed for growth by the MTT method. The cells were also incubated in the presence of CDDO-Me 1, whose ability to inhibit cancer-cell growth was established earlier (reviewed by Liby et al.).\cite{19} Figure 1 shows the dose–response curves for 1 and 12 with KB-3-1 cells, and the IC$_{50}$ values for all the tested compounds (for the inhibition of KB-3-1 cell growth) are presented in Table 1. Compound 12 displayed the highest activity. The in vitro IC$_{50}$ values (the concentrations required for 50% growth inhibition) were 0.3 and 1.2 $\mu$m for 12 and 1, respectively (Table 1). IC$_{50}$ values for the other glycyrrhetinic acid derivatives were greater than 10 $\mu$m.

We compared the effects of 1 and 12 on the growth of different human cancer cell-lines: KB-3-1 epidermoid carcinoma cells, KB-8-5 multidrug-resistant cancer cells (a derivative of KB-3-1), HeLa cervical epithelioid carcinoma cells, MCF-7 breast adenocarcinoma cells, and SKNMC neuroblastoma cells. The dose–response curves for 12 with the different cell lines are displayed in Figure 2. Compound 12 induces concentration-dependent cell death in all cell lines tested. IC$_{50}$ values for 12 and 1 are displayed in Table 2. The IC$_{50}$ values for compounds were similar for all cell lines, with the exception of MCF-7, for which IC$_{50}$ was more than ten times higher than for KB-3-1 (5 $\mu$m vs 0.3 $\mu$m for 12). IC$_{50}$ values for 12 were lower than for 1 for all tested cell lines.

One of the reasons for the failure of chemotherapy-based treatment is multidrug resistance (MDR). We tested the ability of 12 to suppress the growth of multidrug-resistant KB-8-5 cells. This cell line is characterized by overexpression of the MDR1 gene, which encodes P-glycoprotein, an ATP-dependent membrane pump that efficiently decreases the intracellular concentrations of various compounds. Treatment of KB-8-5 cells with 12 significantly decreased the number of living cells (Figure 2); the IC$_{50}$ value for this cell line was only four times higher than the IC$_{50}$ for the drug-sensitive KB-3-1 (0.3 $\mu$m for KB-3-1 vs 1.2 $\mu$m for KB-8-5). Thus, this glycyrrhetinic acid derivative is not targeted at P-glycoprotein, and might be efficient against tumors exhibiting the P-glycoprotein-dependent MDR phenotype.

![Figure 1. Effects of 1 and 12 on the viability of KB-3-1 cells. Cells were incubated for 24 h in the presence of 1 and 12 (0.1, 5 and 10 $\mu$m). Cell viability was measured by MTT assay as described in the Experimental Section. The results are expressed as percentages of viable cells observed after treatment, relative to control cells (100%) incubated in the presence of DMSO (0.1 % (v/v)). The data were obtained from three separate experiments in triplicate.](image1)

**Table 1.** IC$_{50}$ values of glycyrrhetinic acid, its derivatives 1, 7, 8, 9, 10, and 12$^{[a]}$

| Compounds       | IC$_{50}$ $\mu$m | Compounds | IC$_{50}$ $\mu$m |
|-----------------|------------------|-----------|------------------|
| glycyrrhetinic   | > 10             | 8         | > 10             |
| acid            | 1.2 ± 0.16       | 10        | > 10             |
|                 | 7 > 10           | 12        | 0.3 ± 0.08       |

$^{[a]}$ IC$_{50}$ was defined as the compound concentration that resulted in 50% cell survival as measured by the MTT assay (see Experimental Section). Incubation time: 24 h.

![Figure 2. Dose–response curves for 12 with different human cancer cell lines. Cells were incubated for 24 h with increasing concentrations of 12 (0.1 to 100 $\mu$m). Cell viability was measured by the MTT assay as described in the Experimental Section. The results are expressed as percentages of viable cells observed after treatment, relative to control cells (100%) incubated in the presence of DMSO (0.1 % (v/v)). Data were obtained from three separate experiments in triplicate.](image2)

**Table 2.** IC$_{50}$ values for 12 and oleanolic acid derivative 1 with human cancer cell lines$^{[b]}$

| Cell line | IC$_{50}$ [m] | IC$_{50}$ [m] |
|-----------|---------------|---------------|
| KB-3-1    | 0.3 ± 0.08    | 1.2 ± 0.16    |
| KB-8-5    | 1.2 ± 0.12    | 3.1 ± 0.29    |
| HeLa      | 1.3 ± 0.28    | 2.8 ± 0.37    |
| MCF-7     | 5 ± 0.34      | > 10          |
| SKNMC     | 0.8 ± 0.1     | 4.9 ± 0.6     |

$^{[b]}$ IC$_{50}$ was defined as the compound concentration that resulted in 50% cell survival as measured by the MTT assay (see Experimental Section). Incubation time: 24 h.
The effect of antioxidants on the cytotoxicity of 12: Similarly to CDDO and many other synthetic triterpenoids, 12 has potential electrophilic Michael acceptor sites at positions 1 and 9 of the triterpenoid nucleus (Figure 3A). It is known that the presence of Michael acceptor groups at specific positions is essential for inhibition of proliferation, promotion of differentiation, and induction of apoptosis in various cell lines. This arises from the ability of Michael electrophiles to target specific nucleophiles, and to affect selective biological functions. The involvement of the Michael electrophiles in a particular biological process can be proved by inhibition of their activity with antioxidants, for example glutathione (GSH).

We investigated whether reducing the nucleophilic agents would abrogate the cytotoxicity of 12. Cells were incubated in the presence of GSH (1, 5, 15 or 45 μM), either alone or in combination with 12 (1 μM; Figure 3B). Incubation of cells in the presence of 12 with GSH (5 or 15 μM) decreased the cytotoxicity of 12. The IC_{50} value in the presence of GSH was 3 μM, but only 0.3 μM for 12 alone. Higher concentrations of GSH were toxic: incubation in the presence of 45 μM glutathione led to 95% cell death (data not shown). It should be noted that incubation with ascorbic acid did not decrease the cytotoxicity of compound 12 (not shown). Thus, we demonstrated that 12 displays biologically active Michael acceptors.

Cell-cycle arrest: Flow cytometry was employed to determine whether 12 caused stage-specific inhibition of the cell cycle (Table 3). After 18 h incubation in the absence of 12, the number of cells with sub-G1 (apoptotic) peak was insignificant. An increase in the concentration of 12 (0.3 to 1 μM) yielded a corresponding increase in the population of cells in sub-G1 (19.2 to 51.8%; values relative to the control) in a concentration-dependent manner (n = 3; p < 0.05). The increase in the population of cells in the sub-G1 phase was accompanied by a decrease of cells in the G1, and S phases (Table 3). It has been reported that cells with these features are those dying of apoptosis. The number of cells in the G2-M phase remained constant.

Morphological observation of nuclear change: There are several morphological characteristics for apoptotic cells, such as cell shrinkage, nuclear fragmentation and chromatin condensation. To examine cell death due to exposure to 12, we investigated the nuclear morphological changes in KB-3-1 cells treated with 1 μM 12 for 6, 18 and 24 h (Figure 4). Nuclear staining with Hoechst 33258 demonstrated that control KB-3-1 cells had regular and round-shaped nuclei (Figure 4A). In contrast, condensation and fragmentation of nuclei, characteristic of apoptotic cells, were observed in cells treated with 12. After 6 h exposure, patches of localized partially condensed chromatin were found on the inner face of the nuclear membrane (Figure 4B), while the nuclei appeared slightly deformed. The number of such nuclei dramatically increased up to 18 h (Figure 4C and D).

| Table 3. Effect of 12 on the cell cycle of KB-3-1 cells. |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| Conc. 12 [μM]  | Sub-G1 [%]      | G1 [%]          | S [%]           | G2/M [%]        |
| 0              | 1.7             | 50.6            | 22.9            | 24.8            |
| 0.3            | 19.2            | 36.5            | 14.7            | 29.6            |
| 1.0            | 51.8            | 11.8            | 8.3             | 28.1            |

[a] KB-3-1 human epidermoid cells were seeded into six-well plates to ensure that they had not reached confluency. After 24 h they were incubated either in the absence (control) or presence of 0.3 or 1 μM 12. After 18 h the percentage of cells in each phase of the cell cycle was determined by flow cytometry as described in the Experimental Section. Data were obtained from at least three separate experiments in duplicate.
Figure 4C). Incubation of cells for 24 h led to severe damage to the vast majority of nuclei, with the formation of apoptotic bodies (Figure 4D).

Quantification of apoptosis by annexin V binding and flow cytometry: Increases in morphologically changed cells, and in the number of cells in the sub-G0/G1 phase, are usually associated with apoptosis. We examined whether cell death was apoptotic when induced by the glycyrrhetinic acid derivatives by using annexin V and propidium iodide analysis (Figure 5). KB-3-1 cells were exposed to 12, then subjected to flow cytometric analysis. Annexin V binds phosphatidylserine residues, which are asymmetrically distributed toward the inner plasma membrane, and migrate to the outer plasma membrane during apoptosis.[24] The data show that 12 induced apoptotic cell death in 50% of KB-3-1 cells at concentrations equal to the IC50 values. The number of apoptotic cells increased with the time of incubation, and with increasing compound concentration. 89.2% of KB-3-1 cells were detected as apoptotic following 24 h of incubation in the presence of 1 μM 12, so 12 induces dose- and time-dependent apoptotic cell death. Taken together, these data indicate that the decrease in viability of cancer cells exposed to the novel glycyrrhetinic acid derivatives occurred by apoptosis, and that 12 had the greatest potency.

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Dissipation of the mitochondrial transmembrane potential: We investigated whether 12 utilizes the mitochondrial "intrinsic" pathway in the apoptotic death of KB-3-1 cells, as the pivotal role of mitochondria in the triggering of apoptosis is well established. We evaluated the mitochondrial transmembrane potential (ΔΨm) in KB-3-1 cells exposed to 12, and compared this to that for 1, whose ability to decrease ΔΨm has been documented.[10, 19, 25–29] Changes in ΔΨm were evaluated by cytofluorometric analysis. Cells were stained with the mitochondria-specific cationic dye JC-1 (5,5′,6,6′-tetrachloro-1,1′,3,3′-tetraethyl benzimidazole carbocyanine iodide), which accumulates in the transmembrane region of polarized mitochondria where it forms "J-aggregates". These emit orange fluorescence that can be recorded on channel 2 of a cytofluorometer, or visualized via a red filter on a fluorescence microscope. A decrease in ΔΨm results in a decrease in J-aggregates and increase in JC-1 monomers, which emit a greenish-yellow fluorescence. The cytometric analysis of KB-3-1 cells stained with JC-1 is shown in Figure 6B. In the control cells (incubated in the presence of 0.1% DMSO) the majority of cells showed a high emission of fluorescence in both channels, because of the equilibrium between J-aggregates and monomers. The exposure of KB-3-1 cells to 12 leads to a significant decrease in fluorescence compared to the control (0.1% DMSO). In fluorescent

Figure 5. Quantification of apoptosis by annexin V binding to KB-3-1 cells. Cells were incubated in the presence of 0.3 or 1 μM 12, or in the presence of 0.1% (v/v) DMSO for the indicated times. Annexin V binding was carried out with the Annexin V–FITC detection kit as described in the Experimental Section. Annexin V/PI staining was analyzed by flow cytometry. The lower-right quadrant of each cytometry scattergram shows the annexinV+PI– cells; the upper-right quadrant shows the annexinV–PI+ cells. The results are representative of one of three independent experiments.

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microscopy (Figure 6A), one can see that most of the cells turn green. After incubation in the presence of 1, cells can be seen to be somewhere between the control and 12-treated cells, both in the fluorescent micrograph and in the flow-cytometry histogram. One can conclude that 12 causes a dramatic dissipation of mitochondrial potential, and that this effect, consistent with the results of the MTT assay, is more pronounced than that for 1.

**Activation of the caspase cascade in apoptosis induced by glycyrrhetinic acid derivatives:** To determine whether activation of the caspase cascade is involved in 12-induced apoptosis, we used the fluorescein isothiocyanate (FITC)-labeled pan-caspase inhibitor FITC-VAD-FMK (FITC-valyl-alanyl-aspartyl-[O-methyl]fluoromethylketone). The conjugated compound is cell-permeable and binds irreversibly to activated caspase molecules, and thus serves as an in situ marker for apoptosis. We compared the abilities of 12 and 1 to activate caspase (Figure 7). In control cells (18 h incubation in the presence of 0.1 % DMSO) only a faint green signal was seen: this equates to 9 % of cells with activated caspase (Figure 7 A and C). With the addition of 12 (0.3 and 1 μM), the number of cells with activated caspase increased (51 and 85 %, respectively; Figure 7 C), and green fluorescence was observed in the flow cytogram. Similar assays with 1 yielded data that lay between those for the control and 12, as had been the case for mitochondrial transmembrane potential dissipation.

The results provide evidence that the most-active glycyrrhetinic-acid derivative 12 induces caspase-dependent apoptosis in cancer cells. Caspase involvement in cell death is suggested also by the higher IC50 for MCF-7 cells (Table 2)—cells that are known to be caspase-3-deficient.

**Conclusions**

In this report we describe the synthesis of the new glycyrrhetinic acid derivative methyl 2-cyano-3,12-dioxo-18β-H-olean-
9(11),1(2)-dien-30-oate (12), obtained by the direct modification of the A and C rings of glycyrrhetinic acid. We provide a detailed description of synthesis and physicochemical characteristics of the end product, 12, and of the intermediate compounds. The modifications converted the well-known triterpenoid (exhibiting weak antitumor activity) to derivative 12, which displays high antiproliferative activity toward cancer cells. The intermediate products 7–10 did not display this activity.

We have shown that human epidermoid cancer cells are sensitive to 12, as are other tumor cell types, including cells exhibiting the multidrug-resistant phenotype. Compound 12 displays potent single-agent activity, at micromolar concentrations, against different human cancer cells in culture. The mechanism of action (MOA) of triterpenoids on cancer cells is not fully understood. Different mechanisms have been proposed for the cytotoxic activity of synthetic triterpenoids in various types of cancer and leukemia cells; this suggests that cellular context is important. Several studies point to an MOA dependent on the extrinsic apoptotic pathway (DR4/DR5/caspase-8 activation), whereas other studies point to involvement of the intrinsic apoptotic pathway. Our studies imply that the apoptotic MOA of 12 includes components of intrinsic pathways in epidermoid cancer cells.

We have compared the ability of 12 to cause cancer-cell death with that of CDDO-Me, a well known compound that is currently in late-stage clinical trials for the treatment of chronic kidney disease in type 2 diabetes mellitus patients. The antiproliferative activity of 12 exceeds that of CDDO-Me: the IC₅₀ value was lower for all tested cell lines, and the IC₅₀ value was lower for all tested cell lines, and from 0.8 to 4.3% in characteristic of the end product, detailed description of synthesis and physicochemical characteristics of the compounds was determined by NMR from proton spin–spin coupling constants in ¹H, ¹³C, ¹H double-resonance spectra, and by analyzing ¹³C NMR proton-selective and off-resonance saturation spectra. 2D ¹³C, ¹H correlated spectroscopy on CH constants (COSY, JCH = 135 Hz; and COLOC, JCH = 10 Hz, correspondingly), and 1D ¹³C, ¹H long-range J modulation difference (LURMD, JCH = 10 Hz). Flash column chromatography was performed with silica gel (Merck, 60–200 mesh) and neutral alumina (Chemapol, 40–250 mesh).

Methyl 18H-Glycyrrhetinate acetate (3) [42] A solution of diazomethane in ether was added dropwise to 0°C to a stirred suspension of 2 (10 g, 19.0 mmol) in methanol (200 mL) until the originally colorless mixture turned yellow. The resulting mixture was allowed to stand at room temperature overnight. The solvent was removed and the product was purified by crystallization (chloroform/methanol; yield = 91.1 g, 89%). M.p. 303–304 C; ¹H NMR (CDCl₃); δ = 0.76 (dd, JH₂₃H₂₄ = 12.5, JH₂₃H₂₄ = 1.5 Hz; H₂₃), 0.76 (s, 3 H; C₂₈-H₃), 0.84 (s, 3 H; C₂₂-H₂₃, C₂₄-H₂₃), 0.97 (dd, JH₂₃H₂₄ = 13.8 Hz; H₂₄), 1.01 (dd, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 3.7 Hz; H₂₄), 1.08 (s, 3 H; C₂₆-H₂₃), 1.10 (s, 3 H; C₂₉-H₂₃), 1.12 (s, 3 H; C₂₅-H₂₃), 1.14 (ddm, 2 H, 2 H; H3₁, H3₂), 1.41 (dddd, 2 H, 2 H; H3₃, H3₄), 1.42 (dd, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 12.0, JH₂₃H₂₄ = 3.7 Hz; H₂₅), 2.15–1.66 (m, 3 H; H₂₃, H₂₄, H₂₅), 1.75 (dd, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 11.7, JH₂₃H₂₄ = 3.7 Hz; H₂₆), 1.78 (dd, JH₂₃H₂₄ = 13.8, JH₂₃H₂₄ = 13.8, JH₂₃H₂₄ = 4.5 Hz; H₂₇), 1.88 (dd, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 4.2, JH₂₃H₂₄ = 2.7 Hz; H₂₈), 1.95 (dm JH₂₃H₂₄ = 10 Hz; H₂₉), 1.98 (dd, JH₂₃H₂₄ = 13.8, JH₂₃H₂₄ = 13.8, JH₂₃H₂₄ = 4.8 Hz; H₂₉), 2.00 (s, 3 H; C₃₃-H), 2.04 (dd, JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 4.2, JH₂₃H₂₄ = 4.2 Hz; H₂₉), 2.32 (s, 1 H; H₂₉), 2.76 (dd JH₂₃H₂₄ = 13.5, JH₂₃H₂₄ = 3.7, JH₂₃H₂₄ = 3.0 Hz; H₂₉), 3.64 (s, 3 H; OC₃₁-H₃), 4.47 (dd, JH₂₃H₂₄ = 11.7, JH₂₃H₂₄ = 4.7 Hz; H₂₉), 5.62 (s, 1 H; H₂₉), ¹³C NMR (CDCl₃); δ = 38.63 (t, C₁), 23.41 (t, C₂), 80.45 (d, C₃), 37.88 (t, C₄), 54.88 (d, C₅), 17.22 (t, C₆), 32.55 (t, C₇), 43.03 (s, C₈), 61.56 (d, C₉), 36.78 (s, C₁₀), 199.85 (s, C₁₁), 128.34 (d, C₁₂), 169.01 (s, C₁₃), 45.23 (s, C₁₄), 26.31 (t, C₁₅), 26.26 (t, C₁₆), 31.67 (s, C₁₇), 48.25 (d, C₁₈), 40.93 (t, C₁₉), 43.87 (s, C₂₀), 30.08 (t, C₂₁), 37.59 (t, C₂₂), 27.89 (q, C₂₃), 16.52 (q, C₂₄), 16.24 (q, C₂₅), 18.52 (q, C₂₆), 23.17 (q, C₂₇), 28.36 (q, C₂₈), 28.15 (q, C₂₉), 176.73 (s, C₃₀), 51.38 (q, C₃₁), 170.77 (s, C₃₂), 21.13 (q, C₃₃); HRMS: m/z calcd for C₉₉H₇₆O₂₇: 526.7471; found: 526.3658.

Methyl 3β-Acetoxy-18H-olean-12-en-30-oate (4) [43] A solution of conc. hydrochloric acid (50 mL) was added dropwise to 10°C to a stirred suspension of 3 (9.1 g, 17.3 mmol) and zinc powder (18.2 g, 280 mmol) in dioxane (300 mL) over 2 h. The reaction mixture was stirred for a further 3 h at 5–10°C, concentrated in a vacuum, diluted with water (1 L), and filtered. The solid was dried and subjected to flash column chromatography (silica gel; benzene followed by chloroform) to give crude 4 (yield = 66.8 g, 77%). This material was used for the next reaction without further purification. An analytically pure sample was obtained by recrystallization from a mixture chloroform/methanol. M.p. 265–267°C; ¹H NMR (CDCl₃); δ = 0.74 (s, 3 H; C₂₈-H₃), 0.81 (dd, JH₂₃H₂₄ = 12.0, JH₂₃H₂₄ = 1.6 Hz; H₂₃), 0.82–0.86 (m, 6 H, H₂₄), 0.83 (s, 3 H; C₂₄-H₂₅), 0.84 (s, 3 H; C₂₅-H₂₆), 0.93 (s, 6 H; C₂₆-H₂₇, C₂₇-H₂₈), 0.94 (ddm JH₂₃H₂₄ = 13.5 Hz; H₂₄), 1.02 (m, H₂₅), 1.09 (s, 3 H; C₂₉-H₃), 1.10 (s, 3 H; C₃₀-H₃), 1.17–1.17 (m, 4 H; H₂₃, H₂₄, H₂₅, H₂₆), 1.39 (m, 3 H), 1.44–1.64 (m, 7 H; H₂₃, H₂₄, H₂₅, H₂₆, H₂₇, H₂₈), 1.79–1.93 (m, 5 H; H₂₃, H₂₄, H₂₅, H₂₆, H₂₇).
Methyl 3β-acetoxy-12-oxo-18-holean-30-oate (5): A mixture of hydrogen peroxide (30–30%, 25 mL) and acetic acid (25 mL) was added dropwise at 80°C to a stirred suspension of 4 (3.0 g, 5.7 mmol) in acetic acid (100 mL) over 1 h. The reaction mixture was stirred for a further 1 h at 80°C, cooled to room temperature, and diluted with water (500 mL). The solid was filtered, washed with water, and dried to give crude 5 (yield = 6.8 g, 96%). This material was used for the next reaction without further purification. A analytically pure sample was obtained by recrystallization from a mixture chloroform/methanol. M. P. 296–299°C; 1H NMR (CDCl3): δ = 0.79–0.88 (m, 2 H; H6), 0.80 (s, 3 H; C28-H3), 0.83 (s, 3 H; C24-H8), 0.84 (s, 3 H; C23-H3), 0.86 (s, 3 H; C25-H5), 0.90 (s, 3 H; C27-H2), 0.92–0.101 (m, 2 H; H7, 9 η), 1.09 (s, 3 H; C29-H11), 1.10 (s, 3 H; C26-H16), 1.19 (dd, J=13.0, 3.4 Hz; H3), 1.74 (m, 1 H; H4), 1.76 (m, 1 H), 1.85 (dd, J=13.2, 3.4 Hz; H3), 1.91 (dd, J=4.7, 3.4 Hz; H2). 2.00 (s, 3 H; C33-H3), 2.12 (dd, J=17.0, 13.0 Hz; H5), 2.23 (dd, J=17.0, 13.0 Hz; H5), 2.54 (dd, J=13.4, 4.7 Hz; H2), 2.87 (d, J=4.7 Hz; H3), 2.94 (s, 3 H; OC31-H3), 3.05 (m, 3 H; C26-H16), 3.08 (m, 2 H; H21e, H18), 3.18 (s, 3 H; OC31-H3), 3.69 (s, 3 H; OC31-H3), 4.43 (dd, J=13.4, 3.4 Hz; H3), 4.56 (s, 1 H; H11), 4.90 (s, 1 H; H11), 4.92 (s, 1 H; H11), 4.94 (s, 1 H; H11), 5.76 (s, 1 H; H11); 13C NMR (CDCl3): δ = 35.86 (t, C1), 23.66 (t, C2), 79.49 (d, C3), 37.95 (s, C4), 50.08 (d, C5), 17.67 (t, C6), 32.66 (t, C7), 45.29 (s, C8), 177.61 (s, C9), 39.59 (s, C10), 122.89 (d, C11), 201.14 (s, C12), 47.70 (d, C13), 41.61 (s, C14), 26.03 (t, C15), 26.03 (t, C16), 31.87 (t, C17), 37.81 (t, C18), 33.69 (t, C19), 43.88 (s, C20), 31.10 (t, C21), 38.13 (t, C22), 27.77 (t, C23), 16.46 (q, C24), 23.79 (q, C25), 23.85 (q, C26), 21.84 (q, C27), 26.90 (q, C28), 28.45 (q, C29), 177.22 (s, C30), 51.32 (s, C31), 170.59 (s, C32), 21.01 (q, C33); HRMS: m/z calcd for C39H42O5: 572.3673; found: 572.3668.

Methyl 3β-hydroxy-12-oxo-9(11)-en-30-oate (7): A mixture of 6 (6.2 g, 11.8 mmol) and KOH (41 g, 732 mmol) in methanol (400 mL) was heated under reflux for 1.5 h. The resulting solution was cooled to room temperature, concentrated in vacuo, and 10% aqueous hydrochloric acid solution was added. The mixture was extracted with ether. An analytically pure sample was obtained by recrystallization from a mixture chloroform/methanol. M. P. 202–203°C; 1H NMR: δ = 0.80 (s, 3 H; C24-H8), 0.89 (s, 3 H; C28-H2), 0.94 (s, 3 H; C27-H2), 1.01 (s, 3 H; C23-H3), 1.08 (s, 3 H; C29-H11), 1.16 (s, 3 H; C25-H5), 1.35 (s, 3 H; C26-H16), 0.85–0.91 (m, 2 H; H7, H9), 1.04 (dm, J=12.8 Hz; H8), 1.23 (dd, J=13.2, 3.2 Hz; H8), 1.18–1.25 (m, 1 H; H7), 1.27 (dm, J=14.0 Hz; H8), 1.31 (m, 1 H; H8), 1.44 (dm, J=9.8 Hz; H8), 1.49 (dd, J=14.0, 14.0 Hz; H8), 4.25 (H2), 1.55–1.76 (m, 5 H; 2 H 6, H7), 2.18 (m, 1 H; H8), 1.85 (m, 1 H; H8), 1.89–2.00 (m, 3 H; H7, H8, H9), 2.19 (dd, J=19.9, 13.2, J=19.9 Hz; H9), 3.29 (J=13.2, J=3.2 Hz; H10), 2.73 (H2), 2.95 (J=13.2, J=3.2 Hz; H10); 13C NMR (CDCl3): δ = 35.86 (t, C1), 23.66 (t, C2), 79.49 (d, C3), 37.95 (s, C4), 50.08 (d, C5), 17.67 (t, C6), 32.66 (t, C7), 45.29 (s, C8), 177.61 (s, C9), 39.59 (s, C10), 122.89 (d, C11), 201.14 (s, C12), 47.70 (d, C13), 41.61 (s, C14), 26.03 (t, C15), 26.03 (t, C16), 31.87 (t, C17), 37.81 (t, C18), 33.69 (t, C19), 43.88 (s, C20), 31.10 (t, C21), 38.13 (t, C22), 27.77 (t, C23), 16.46 (q, C24), 23.79 (q, C25), 23.85 (q, C26), 21.84 (q, C27), 26.90 (q, C28), 28.45 (q, C29), 177.22 (s, C30), 51.32 (s, C31), 170.59 (s, C32), 21.01 (q, C33); HRMS: m/z calcd for C39H42O5: 572.3674; found: 572.3668.
Methyl 2-hydroxyethylene-3,12-dioxo-18-h-olean-9(11)-en-30-olate (9): Ethyl formate (3.75 mL, 39.5 mmol) and sodium methanolate (2.1 g, 38.9 mmol) were added to a solution of ketone (2.17 g, 38.9 mmol) in dry benzene (70 mL). The mixture was stirred at room temperature for 2 h. The reaction mixture was diluted with a mixture of chloroform/diethyl ether (1:3, 100 mL), and 5% HCl was added to achieve pH < 7. The organic layer was separated, and the aqueous layer was extracted with chloroform/diethyl ether (1:3; 3 × 50 mL). The combined organic layers were washed with saturated sodium hydrogen carbonate solution (3 × 50 mL), and brine (3 × 50 mL), and dried over magnesium sulfate. The solvent was removed to give an amorphous solid 9 (yield = 4.5 g, 95%). This material was used for the next reaction without further purification.

Analitically pure sample was obtained by flash column chromatography (silica gel; hexane/ethyl acetate (9:1) followed by hexane/ethyl acetate (3:1)). 1H NMR (CDCl3, δ): 0.90 (3 H, 7H; C28-H 3), 0.96 (3 H, C27-H 2), 1.09 (3 H, C29-H 1), 1.13 (3 H, C24-H 4), 1.15 (3 H, C25-H 3), 1.20 (3 H, C23-H 1), 1.38 (3 H, C26-H 2), 0.91 (m, H 15), 0.97 (m, H 16), 1.23 (dd, 3 J(H2e,H1a)=3.3, 3 J(H19e,H19a)=4.6, 3 J(H21a,H1)=3.3 Hz; H 21), 2.21 (dd, 3 J(H21a,H1)=3.3, 3 J(H19e,H21e)=4.7 Hz; H 21), 2.26 (d, 3 J(H1,H1)=14.5 Hz; H 1 and 2.58 (d, 3 J(H19e,H18)=14.5 Hz; H 18) - Ab- system, 3.02 (d, 3 J(H21a,H1)=4.6 Hz; H 21), 3.71 (3 H, C31-OH), 5.90 (m, H 17), 8.70 (m, 3 J OH=2.4 Hz; H 17), 14.81 (d, 3 J OH=2.4 Hz; H 17); 13C NMR (CDCl3) δ: 36.85 (t, C1), 104.80 (s, C2), 188.08 (s, C3), 40.32 (s, C4), 48.07 (d, C5), 18.80 (t, C6), 31.25 (t, C7), 45.55 (s, C8), 175.36 (s, C9), 38.89 (s, C10), 124.37 (d, C11), 200.73 (s, C12), 47.86 (d, C13), 41.80 (s, C14), 26.22 (d, C15), 31.93 (s, C17), 37.82 (d, C18), 33.76 (t, C19), 43.92 (s, C20), 31.19 (t, C21), 35.81 (s, C22), 28.14 (q, C23), 20.65 (s, C24), 23.36 (q, C25), 23.23 (q, C26), 21.82 (q, C27), 26.96 (q, C28), 28.48 (q, C29), 177.22 (s, C30), 51.42 (q, C31), 189.65 (d, C32); HRMS: m/z calcd for C39H60O7: 548.46954; found: 548.43396.

The mixture of tautomer (11): Sodium methanolate (11 g, 204 mmol) was added to 0 °C to a solution of isoxazole (10) (3.0 g, 5.9 mmol) in methanol (85 mL) and diethyl ether (170 mL). The mixture was stirred at room temperature for 1 h. The resulting mixture was diluted with a mixture of chloroform/diethyl ether (1:3; 100 mL), and 5% HCl was added to achieve pH < 7. The organic layer was separated, and the aqueous layer was extracted with chloroform/diethyl ether (1:3; 3 × 50 mL). The combined organic layers were washed with saturated sodium hydrogen carbonate solution (3 × 50 mL) and brine (3 × 50 mL), and dried over magnesium sulfate. The solvent was removed to give a mixture of tautomers 11 (yield = 3.0 g, 100%). This material was used for the next reaction without further purification.

HRMS: m/z calcd for C39H59NO7: 517.7040; found: 507.3349.

Methyl 2-cyano-3,12-dioxo-18-h-olean-9(11),12-dien-30-olate (12): Mixture 11 (2.8 g, 5.5 mmol) and 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) (1.5 g, 6.5 mmol) in dry benzene (160 mL) were heated under reflux for 4 h. Insoluble matter was removed by filtration, and the filtrate was evaporated in a vacuum to give a solid. The solid was subjected to flash column chromatography (silica gel; benzene followed by benzene/acetone (10:1)) to give crude 12. The crude product was purified by recrystallization from methanol/chloroform to give crystals 12 (yield 1.7 g, 61%). M. p. 247–249 C; 1H NMR (CDCl3, δ): 0.90 (3 H, C28-H 3), 0.96 (3 H, C27-H 2), 1.09 (3 H, C29-H 1), 1.13 (3 H, C24-H 4), 1.15 (3 H, C25-H 1), 1.20 (3 H, C23-H 1), 1.38 (3 H, C26-H 2), 0.91 (m, H 15), 0.97 (m, H 16), 1.23 (dd, 3 J(H2e,H1a)=3.3, 3 J(H19e,H19a)=4.6, 3 J(H21a,H1)=3.3 Hz; H 21), 2.21 (dd, 3 J(H21a,H1)=3.3, 3 J(H19e,H21e)=4.7 Hz; H 21), 2.26 (d, 3 J(H1,H1)=14.5 Hz; H 1 and 2.58 (d, 3 J(H19e,H18)=14.5 Hz; H 18) - Ab- system, 3.02 (d, 3 J(H21a,H1)=4.6 Hz; H 21), 3.71 (3 H, C31-OH), 5.90 (m, H 17), 8.70 (m, 3 J OH=2.4 Hz; H 17), 14.81 (d, 3 J OH=2.4 Hz; H 17); 13C NMR (CDCl3) δ: 36.85 (t, C1), 104.80 (s, C2), 188.08 (s, C3), 40.32 (s, C4), 48.07 (d, C5), 18.80 (t, C6), 31.25 (t, C7), 45.55 (s, C8), 175.36 (s, C9), 38.89 (s, C10), 124.37 (d, C11), 200.73 (s, C12), 47.86 (d, C13), 41.80 (s, C14), 26.22 (d, C15), 31.93 (s, C17), 37.82 (d, C18), 33.76 (t, C19), 43.92 (s, C20), 31.19 (t, C21), 35.81 (s, C22), 28.14 (q, C23), 20.65 (s, C24), 23.36 (q, C25), 23.23 (q, C26), 21.82 (q, C27), 26.96 (q, C28), 28.48 (q, C29), 177.22 (s, C30), 51.42 (q, C31), 189.65 (d, C32); HRMS: m/z calcd for C39H60O7: 548.46954; found: 548.43396.
Cell culture and glycyrrhetinic acids derivatives: Human KB-3-1 epidermoid carcinoma cell line, HeLa cervical epithelial carcinoma cell line, MCF-7 breast adenocarcinoma cell line, SKNMC neuroblastoma cell line (Russian Cell Culture Collection, St. Petersburg), KB-8-5 multidrug resistant cancer cell line (kindly provided by Professor M. Gottesman (NIH, USA)), were cultured in DMEM supplemented with 10% (v/v) heat-inactivated fetal bovine serum, penicillin (100 U mL\(^{-1}\)), streptomycin (100 \mu g mL\(^{-1}\)) and amphotericin (250 \mu g mL\(^{-1}\)). Cells were maintained in a humidified atmosphere (5% CO\(_2\), 37 °C). The KB-8-5 cell line was incubated in the additional presence of vinblastine (300 nmol L\(^{-1}\)).

Glycyrrhetinic acids derivatives were dissolved in DMSO (10 mmol L\(^{-1}\)), and stock solution were stored at −20 °C.

After treatments, both floating and adherent scraped cells were collected by centrifugation, and used for further analysis.

Cell viability analysis by MTT assay: Cancer cells, growing in log phase, were seeded in triplicate 96-well plates at a density of 5 \times 10\(^4\) cells per well for HeLa cells, 7 \times 10\(^4\) for KB-3-1, KB-8-5 and MCF-7 cells, and 30 \times 10\(^4\) for SKNMC cells. The plates were incubated at 37 °C in humidified 5% CO\(_2\) atmosphere. Cells were allowed to adhere to the surface for 24 h, then treated with varying doses of the compounds for 24 h. Aliquots of [3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide] (MTT) solution (10 \mu L, 5 mg mL\(^{-1}\)) were added to each well, and the incubation was continued for an additional 3 h. The dark blue formazan crystals (formed within healthy cells) were solubilized with DMSO, and the absorbance was measured at 570 nm in a Multiscan RC plate reader (Thermo LabSystems, Finland). The IC\(_{50}\) was determined as the compound concentration required to decrease the A\(_{492}\) to 50% of the control (no compound, DMSO), and was determined by interpolation from dose-response curves.

Analysis of antioxidant effect on the cytotoxicity of compound 12: KB-3-1 cells growing in the log phase were seeded in triplicate in 96-well plates (7 \times 10\(^4\) cells per well). The plates were incubated at 37 °C in a humidified 5% CO\(_2\) atmosphere. Cells were allowed to adhere to the surface for 24 h, then treated with GSH (1, 5, 15 or 45 mm) or with ascorbic acid (1, 3 or 5 mm), both alone and in combination with 12 (1 \mu M). Cells were incubated with the compounds for 24 h and cell viability was analyzed by the MTT assay as described above.

Morphological observation of nuclear change: KB-3-1 cells were seeded into 24-well plates (10\(^4\) cells per well) containing glass cover slips. Cells were allowed to adhere to the surface for 24 h. Cells were treated with 12 (1 \mu M) or with DMSO (0.1% (v/v)) for 6, 18 or 24 h at 37 °C in a humidified 5% CO\(_2\) atmosphere. After incubation, cells were fixed with 4% formaldehyde for 15 min, and then stained for 30 min with Hoechst 33258 (200 nmol L\(^{-1}\)). Cells were analyzed for the presence of fragmented nuclei and condensed chromatin by fluorescent microscopy.

Apoptosis detection by Annexin V staining: Log-phase KB-3-1 cells in six-well plates (5 \times 10\(^4\) cells per well) were treated with 12 (0.3 \mu M or 1 \mu M) or with DMSO (0.1% (v/v)) for 4, 18 or 24 h. The cells were stained with Annexin V-FITC and propidium iodide by using the ApopNexin-FITC apoptosis detection kit (Chemicon Millipore) according to the manufacturer's instructions. Briefly, cells were collected by scraping, washed twice with cold PBS, and centrifuged (400 g, 5 min). Cells were resuspended in binding buffer (1 mL with a concentration of 1 \times 10\(^5\) cells per mL, then a sample (200 \mu L) was transferred to a 5 mL culture tube, and Annexin V-FITC (3 \mu L) and 100 \times PI (2 \mu L) were added. Cells were incubated for 15 min at room temperature in the dark. Finally, binding buffer (300 \mu L) was added to each tube, and the quantity of apoptotic cells in samples was analyzed by flow cytometry (FCS500, Beckman Coulter, USA). For each sample, 10,000 ungated events were acquired. Annexin V+/PI− cells represented early apoptotic populations. Annexin V−/PI+ cells represented either late apoptotic or secondary necrotic populations.

Mitochondria depolarization analysis: Mitochondria involvement in apoptosis was measured by the mitochondrial depolarization that occurs early during the onset of apoptosis. KB-3-1 cells were treated with 1 (1 \mu M), 12 (1 \mu M) or DMSO (0.1% (v/v)) for 6 h, and loss of mitochondrial potential was determined by using the mitochondrial potential sensor JC-1 (Molecular Probes, Invitrogen).

Flow cytometry assay: Cells were incubated for the appropriate time with the compounds, then collected, incubated in complete media in the dark with JC-1 (5 \mu M) at 37 °C for 15 min, and washed with PBS. At the end of the incubation period the cells were washed twice with cold PBS, and resuspended in PBS (400 \mu L). J-aggregate and J-monomer fluorescence were recorded in the channels 2 (FL2) and 1 (FL1), respectively, of an FC500 flow cytometer. Necrotic fragments were electronically gated out, on the basis of morphological characteristics on the forward light scatter versus side light scatter dot plot.

Fluorescent microscopy assay: Cells were plated into 24-well plates (10\(^3\) cells per well) containing glass cover slips, and allowed to adhere to the surface for 24 h. Cells were incubated for the appropriate time with the compounds. After incubation the cell culture media was removed and replaced with JC-1 reagent (5 \mu M) diluted in PBS. Cells were incubated at 37 °C in a 5% CO\(_2\) incubator for 15 min, and analyzed by fluorescence microscopy.

Cytosfluorimetric analysis of DNA content: Exponentially growing KB-3-1 cells in 6-well plates (5 \times 10\(^4\) cells per well) were treated with 12 (0.3 \mu M or 1 \mu M) or DMSO (0.1% (v/v)) for 18 h. After incubation, the cells were collected by centrifugation (400 g, 10 min), fixed with ice-cold 70% ethanol for at least 1 h at 4 °C and treated with RNase A from bovine pancreas (1 mg mL\(^{-1}\)) for 30 min at 37 °C. PI (50 \mu g mL\(^{-1}\)) was then added to the solution and the DNA content was quantitated by a flow cytometry. Cells in sub-G1 phase were considered apoptotic.

Analysis of caspase activation: After treatment of KB-3-1 cells with 1 (0.3 \mu M, 1 \mu M), or 12 (0.3 \mu M, 1 \mu M), or DMSO (0.1% (v/v)) for 18 h, caspase activation was assayed by using the CaspACE FITC-VAD-FMK in situ marker (Promega).

Flow cytometry assay: Cells were incubated for the appropriate time in the presence of the compounds, collected, suspended in PBS (0.5 mL), and FITC-VAD-FMK (1 \mu L, 5 mM) was added. The cells were gently mixed and incubated for 20 min at RT in the dark. Cells were washed twice with PBS, and the pellets resuspended in PBS (0.5 mL). Flow cytometry was conducted within 10 min.

Fluorescent microscopy assay: Cells were seeded (10\(^4\) cells per well) into 24-well plates containing glass cover slips, and allowed to adhere to the surface for 24 h. After incubation for the appropriate time with the compounds, the cell culture medium was removed and replaced with JC-1 reagent (5 \mu M) diluted in PBS. Cells were incubated at 37 °C in a 5% CO\(_2\) incubator for 15 min. The cells were
washed twice with PBS, and caspase activation was analyzed by fluorescence microscopy within 10 min.

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