Cotylenin A, a Differentiation-inducing Agent, and IFN-α Cooperatively Induce Apoptosis and Have an Antitumor Effect on Human Non-Small Cell Lung Carcinoma Cells in Nude Mice

Yoshio Honma, Yuki Ishii, Yuri Yamamoto-Yamaguchi, Takeshi Sassa, and Ken-ichi Asahi

Saitama Cancer Center Research Institute, Saitama 362-0806 [Y. H., Y. I., Y. Y.-Y., K. A.], and Department of Bioresource Engineering, Yamagata University, Tsuruoka [T. S.], Japan

ABSTRACT

Cotylenin A, a novel inducer of the differentiation of leukemia cells, and IFN-α synergistically inhibited the growth of and induced apoptosis in several human non-small cell lung carcinoma cell lines. Tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) and its receptor DR5 were the early genes induced by the combination of cotylenin A and IFNα in lung carcinoma cells. Neutralizing antibody to TRAIL inhibited apoptosis, suggesting that cotylenin A and IFNα cooperatively induced apoptosis through the TRAIL signaling system. This combined treatment preferentially induced apoptosis in human lung cancer cells while sparing normal lung epithelial cells and significantly inhibited the growth of human lung cancer cells as xenografts without apparent adverse effects, suggesting that this combination may have therapeutic value in treating lung cancer.

INTRODUCTION

NSCLC is one of the most common malignant diseases in the world. Surgical resection is the only treatment modality with a reasonable chance of offering cure when applied to appropriately selected patients. Only ~40% of patients have resectable disease, and only one-quarter of these (10–12%) are still alive at 5 years and apparently cured of their disease. Chemotherapy is reserved solely for patients with advanced stages of NSCLC and until now has only brought marginal benefits (1). Efforts are underway to optimize chemotherapeutic strategies and discover new agents. Attempts to translate recent findings regarding the biology of lung cancer into therapeutic strategies, such as the use of biological response modifiers, monoclonal antibodies, and inhibitors of signal transduction of oncogenes, may some day lead to significant progress.

IFNs are pleiotropic cytokines that block viral infection, inhibit cell proliferation, induce apoptosis, and modulate cell differentiation (2). IFNα has therapeutic activity as a single agent in some types of hematological malignancies but is less effective in the therapy of solid tumors, including NSCLC (2–4). To overcome this resistance, various therapeutic strategies, such as the use of biological response modifiers, monoclonal antibodies, and inhibitors of signal transduction of oncoproteins, may some day lead to significant progress.

Chemicals. Cotylenin A was purified from the culture filtrate of Cladosporium fungus sp. 501–7W by flash chromatography on silica gel with 18 U.S.C. Section 1734 solely to indicate this fact. The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.
Cells and Cell Culture. Human lung carcinoma cell lines were maintained in RPMI 1640 supplemented with 10% FBS at 37°C in a humidified atmosphere containing 5% CO2 in air (14). A549 was established from a human lung carcinoma with properties of type II alveolar epithelial cells. PC9 and PC14 were well- and poorly differentiated adenocarcinoma cell lines, respectively. Human normal bronchial epithelial and SAE cells, which were derived from human lung bronchi and the small airway of healthy donors, respectively, were purchased from Clonetics (San Diego, CA) and grown in serum-free Bronchial Epithelial Cell and Small Airway Epithelial Cell Growth Medium Bullet kits (Clonetics), respectively.

Assay of Cell Growth and Apoptosis. The cells were seeded at 1 x 10^4/ml in a 24-well multidish. After culture with or without the test compounds for the indicated times, viable cells were examined by the modified MTT assay. Briefly, 100 µl of MTT solution (1 mg/ml in PBS) were added to each well. After incubation with MTT for 4 h, the cells were centrifuged at 1000 x g for 10 min. The precipitates were dissolved in 1 ml of DMSO, and their absorption at 560 nm was determined. Assay of the cumulative cell number was determined as described elsewhere (21). The cellular DNA content was analyzed using propidium iodide-stained nuclei (21). Caspase activity in intact cells was measured using PhiPhLux by flow cytometric analysis according to the manufacturer’s instructions (Oncolimmunin, Inc., Gaithersburg, MD).

Analysis of TRAIL-, DR4-, and DR5-positive Cells. We detected the expression of TRAIL and its receptors by flow cytometry. Total RNA was extracted using Isogen (Nippon Gene, Toyama, Japan) according to the manufacturer’s instructions. Total RNA (1 µg) from lung cancer cells was converted to first-strand cDNA primed with random hexamer in a 20-µl reaction using an RNA PCR kit (Takara Shuzo Co., Ltd., Tokyo, Japan), and 4 µl of this reaction were used as a template in the PCR. The oligonucleotide oligomers used in PCR amplification were as described elsewhere (22, 23), and a quantitative RT-PCR reaction was performed as described in the literature (24).

Transplantation of Lung Cancer Cells into Nude Mice and Treatment. Female athymic nude mice with a BALB/c genetic background were supplied by CLEA Japan (Tokyo, Japan). They were housed under specific pathogen-free conditions. The in vivo experiments were performed in accordance with the guidelines of our institute (Guide for Animal Experimentation, Saitama Cancer Center). Mice were inoculated s.c. with 6 x 10^5 PC14 cells. A stock solution of cotylenin A for administration was prepared in DMSO at 100 mg/ml. Mice were given a daily s.c. injection of 0.1 ml of PBS, including 3 x 10^4 IU of IFNα, and/or s.c. injections every other day of 0.2 ml of PBS, including 100 µg of cotylenin A (6.7 mg/kg body weight) at a site distant to the tumors, with the first injection given 7 days after the inoculation of tumor cells. Tumor size was measured with vernier calipers every other day. Statistical analysis was performed using Student’s t test.

RESULTS

Combined Effects of IFNα and Various Drugs on the Growth of NSCLC Cells. To measure the effects of various drugs on the growth of lung carcinoma A549 cells, the number of viable cells was determined by the MTT assay after 6 days of exposure to various concentrations of drugs with or without 300 IU/ml IFNα. The growth-inhibiting effects of the drugs were examined by determining the concentrations of drugs required to reduce the cell number to one-half of that in untreated cells (IC50). The sensitivity to anticancer agents, such as 5-fluorouracil, cis-platin, or doxorubicin, was not affected by IFNα, whereas the sensitivity to hydroxyurea was significantly enhanced (Table 1). Most of the differentiation-inducing agents for myeloid leukemia cells were not toxic toward the lung carcinoma cell line when used within a range of concentrations that were effective at inducing the differentiation of human myeloid leukemia cells, suggesting that the lung cancer cells were less sensitive to agents that inhibited cell growth. Retinoids and vitamin D3 did not affect the growth of A549 cells even at high concentrations (Table 1). The growth-inhibitory effect of DMSO was significantly enhanced by IFNα, although IFNα alone hardly inhibited cell growth (Table 1). We next examined the effects of >50 compounds that induce the differentiation of myeloid leukemia cells (25) and found that although the effects of some agents were strongly affected by IFNα, the effects of other agents were not. Among the differentiation-inducing agents tested, the sensitivity of lung cells to cotylenin A was most strongly affected by IFNα (Table 1). The synergistic effects of cotylenin A and IFNα were also observed in other lung carcinoma cell lines, although the sensitivity of lung carcinoma cell lines to IFNα varied among the cell lines (Fig. 1 and Table 2).

Next, we examined the combined effects of cotylenin A and various cytokines on the growth of A549 cells. Unlike IFNα, IFNγ did not affect cotylenin A-induced growth inhibition (Fig. 2b). TNFα and TGFβ also showed cooperative effects with cotylenin A in the inhibition of cell proliferation (Fig. 2, c and d), whereas other cytokines, such as interleukin 1, interleukin 4, leukemia-inhibitory factor, granulocyte macrophage colony-stimulating factor, granulocyte colony-stimulating factor, granulocyte macrophage colony-stimulating factor, and keratinocyte growth factor, did not show such cooperation with cotylenin A (data not shown). IFNα was the most potent cytokine at inhibiting cell growth in the presence of cotylenin A. These results suggest that the combination of cotylenin A and IFNα is the most potent for inhibiting the growth of human lung cancer cells.

Induction of Apoptosis in A549 Cells Treated with IFNα Plus Cotylenin A. When exposed to cotylenin A in the presence of 600 IU/ml IFNα for 4 days, the number of viable A549 cells decreased in a dose-dependent manner. After exposure for 4 days, a morphological analysis showed shrunken cells, chromatin condensation, nuclear fragmentation, and cytoplasmic blebbing (data not shown). Induction of apoptosis (cells in sub-G1 phase) in treated A549 cells was confirmed by an analysis of DNA histograms (Fig. 3A) and caspase-3 activation (Fig. 3B). Treatment with cotylenin A alone also inhibited

Table 1 Potentiation of the growth-inhibitory activities of various agents in human lung carcinoma A549 cells by IFNα

<table>
<thead>
<tr>
<th>Compound</th>
<th>IFNα</th>
<th>IFNα + Cotylenin A</th>
<th>Ratio (−/+)</th>
</tr>
</thead>
<tbody>
<tr>
<td>5-Fluorouracil (µg/ml)</td>
<td>0.81 ± 0.06</td>
<td>0.59 ± 0.04</td>
<td>1.4</td>
</tr>
<tr>
<td>Hydroxyurea (µM)</td>
<td>407 ± 38</td>
<td>149 ± 11</td>
<td>2.7</td>
</tr>
<tr>
<td>Doxorubicin (ng/ml)</td>
<td>31.4 ± 2.8</td>
<td>22.4 ± 1.9</td>
<td>1.4</td>
</tr>
<tr>
<td>Cis-platin (ng/ml)</td>
<td>312 ± 30</td>
<td>259 ± 22</td>
<td>1.2</td>
</tr>
<tr>
<td>Etoposide (µg/ml)</td>
<td>5.13 ± 0.3</td>
<td>3.03 ± 0.2</td>
<td>1.7</td>
</tr>
<tr>
<td>Camptotecin (µg/ml)</td>
<td>124 ± 8.1</td>
<td>72.2 ± 4.8</td>
<td>1.7</td>
</tr>
<tr>
<td>Actinomycin D (ng/ml)</td>
<td>0.31 ± 0.02</td>
<td>0.24 ± 0.02</td>
<td>1.3</td>
</tr>
<tr>
<td>All-trans retinoic acid (µM)</td>
<td>&gt;36</td>
<td>&gt;36</td>
<td></td>
</tr>
<tr>
<td>13-Cis retinoic acid (µM)</td>
<td>&gt;36</td>
<td>&gt;36</td>
<td></td>
</tr>
<tr>
<td>1a,25-Dihydroxyvitamin D3 (ng/ml)</td>
<td>&gt;90</td>
<td>&gt;90</td>
<td></td>
</tr>
<tr>
<td>Cotylenin A (µg/ml)</td>
<td>15.4 ± 9.1</td>
<td>11.6 ± 1.8</td>
<td>0.86</td>
</tr>
<tr>
<td>Sodium butyrate (µM)</td>
<td>521 ± 43</td>
<td>164 ± 11</td>
<td>3.2</td>
</tr>
<tr>
<td>Dimethyl sulfoxide (µM)</td>
<td>257 ± 21</td>
<td>78.2 ± 7.3</td>
<td>3.3</td>
</tr>
<tr>
<td>Hexamethylene bisacetamide (µM)</td>
<td>12.7 ± 1.1</td>
<td>4.4 ± 0.4</td>
<td>2.9</td>
</tr>
<tr>
<td>Isopenylenaldehyde (µg/ml)</td>
<td>86.1 ± 8.2</td>
<td>69.3 ± 5.9</td>
<td>1.3</td>
</tr>
<tr>
<td>8-C-AMP (µM)</td>
<td>24.7 ± 1.9</td>
<td>18.8 ± 1.6</td>
<td>1.3</td>
</tr>
<tr>
<td>Butafil (µM)</td>
<td>1.6 ± 0.1</td>
<td>0.8 ± 0.1</td>
<td>0.9</td>
</tr>
<tr>
<td>Trichostatin A (ng/ml)</td>
<td>71.5 ± 7.6</td>
<td>48.2 ± 4.1</td>
<td>1.5</td>
</tr>
<tr>
<td>Genistein (µg/ml)</td>
<td>8.2 ± 0.7</td>
<td>6.3 ± 0.6</td>
<td>1.3</td>
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<tr>
<td>Wortmannin (µM)</td>
<td>4.9 ± 0.5</td>
<td>3.2 ± 0.3</td>
<td>1.5</td>
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</table>

*Cells were incubated with various concentrations of the test compounds in the presence or absence of 600 IU/ml IFNα for 6 days. Ratio (−/+), IC50 without IFNα/IC50 with IFNα. The values are the mean ± SD of four determinations.*
were obtained when the cells were treated with anti-Fas monoclonal antibody CH-11, which was highly effective at inducing apoptosis in lymphoid cells. Although TRAIL significantly enhanced the growth inhibition induced by DMSO, the enhancing effects were less than those of cotylenin A (Fig. 5c). Similar results were observed when cell growth in a concentration-dependent manner. Cotylenin A alone induced G₁ arrest, but not apoptosis, even at a high concentration (Fig. 3A). Combined treatment with 300 IU/ml IFNα plus 4 μg/ml cotylenin A caused significant growth inhibition, and this effect was similar to that with 24 μg/ml cotylenin A alone (Fig. 4). When the cells were treated with cotylenin A and IFNα for 7 days, and then washed and cultured without the drugs, cell growth was greatly inhibited at day 16, indicating that the growth-inhibitory effect of cotylenin A plus IFNα was irreversible in long-term culture. On the other hand, cotylenin A-treated cells began to grow within a few days after removal of the drug (Fig. 4). These results were consistent with the finding that the combination of cotylenin A and IFNα induced apoptosis, whereas a high concentration of cotylenin A induced G₁ arrest in A549 cells (Fig. 3), and suggest that this combined treatment may have therapeutic value in the chemotherapy of some lung cancers.

Fas and/or TRAIL signaling pathways are involved in IFN-induced apoptosis in some malignant cells (26–29). Therefore, we examined the effects of Fas ligand and TRAIL on the growth of A549 cells in the presence of IFNα or cotylenin A. The combination of IFNα and Fas ligand or TRAIL scarcely inhibited the proliferation of A549 cells (data not shown), whereas cotylenin A cooperated with TRAIL at inhibiting cell growth (Fig. 5a). However, Fas ligand did not affect the growth inhibition induced by cotylenin A (Fig. 5b). Similar results

Table 2: Synergistic effects of IFNα and cotylenin A on the growth of human lung carcinoma cell lines

<table>
<thead>
<tr>
<th>Cell line</th>
<th>IC₅₀ for cotylenin A (μg/ml)</th>
<th>− IFNα</th>
<th>+ IFNα</th>
<th>Ratio (+/−)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adenocarcinoma</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A549</td>
<td>15.4 ± 1.1</td>
<td>1.8 ± 0.1</td>
<td>8.6</td>
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<tr>
<td>PC7</td>
<td>37.6 ± 3.1</td>
<td>3.6 ± 0.3</td>
<td>10.4</td>
<td></td>
</tr>
<tr>
<td>PC9</td>
<td>34.8 ± 2.9</td>
<td>7.9 ± 0.5</td>
<td>4.4</td>
<td></td>
</tr>
<tr>
<td>PC14</td>
<td>28.6 ± 2.2</td>
<td>2.1 ± 3.2</td>
<td>13.6</td>
<td></td>
</tr>
<tr>
<td>ABC-1</td>
<td>16.5 ± 1.8</td>
<td>2.2 ± 0.2</td>
<td>7.5</td>
<td></td>
</tr>
<tr>
<td>Squamous cell carcinoma</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>EBC-1</td>
<td>6.3 ± 4.2</td>
<td>0.8 ± 0.1</td>
<td>7.9</td>
<td></td>
</tr>
<tr>
<td>LK2</td>
<td>26.8 ± 2.4</td>
<td>6.1 ± 0.8</td>
<td>4.4</td>
<td></td>
</tr>
<tr>
<td>Large cell carcinoma</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lu65</td>
<td>16.3 ± 1.4</td>
<td>3.1 ± 0.3</td>
<td>5.3</td>
<td></td>
</tr>
<tr>
<td>Lu99</td>
<td>7.5 ± 0.7</td>
<td>0.8 ± 0.1</td>
<td>9.4</td>
<td></td>
</tr>
<tr>
<td>Small cell carcinoma</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lu135</td>
<td>4.8 ± 0.4</td>
<td>1.5 ± 0.1</td>
<td>3.2</td>
<td></td>
</tr>
<tr>
<td>H69</td>
<td>14.8 ± 1.2</td>
<td>3.4 ± 0.2</td>
<td>4.4</td>
<td></td>
</tr>
</tbody>
</table>

*IC₅₀ for cotylenin A was determined by the drug concentration required for 50% growth inhibition. The values are mean ± SD of four determinations.

1. Cells were cultured with various concentrations of cotylenin A in the presence or absence of 600 IU/ml IFNα for 6 days. IC₅₀ of Lu65 cells for IFNα alone was 3402 ± 194 IU/ml, and the IC₅₀ of the other cells were >6000 IU/ml. The values are the mean ± SD of four determinations.

2. p53 status is wild type in A549 and Lu99 cell lines and mutated in the other cell lines.

Fig. 1. Synergistic effects of IFNα and cotylenin A on the growth of A549 and PC9 cells. Cells were cultured with various concentrations of cotylenin A in the presence of 0–300 IU/ml IFNα for 6 days. The values are mean ± SD of four determinations.

Fig. 2. Effect of cytokines on growth of A549 cells in the presence of cotylenin A. Cells were cultured with various concentrations of IFNα (a), IFNγ (b), TGFβ (c), or TNFα (d) in the presence of cotylenin A for 6 days. The values are mean of four determinations.

Fig. 3. A, induction of G₁ arrest and apoptosis in A549 cells treated with a high concentration of cotylenin A and a combination of cotylenin A and IFNα, respectively. Cells were cultured without (1) or with 24 μg/ml cotylenin A (2) or a combination of 300 IU/ml IFNα and 4 μg/ml cotylenin A (3) for 4 days, and DNA histograms were then analyzed. The apoptotic cell population is shown according to the sub-G₁ fraction. B, increase in cells with caspase-3 activity by cotylenin A plus IFNα but not by cotylenin A. Cells were treated with 24 μg/ml cotylenin A (1) or 300 IU/ml IFNα plus 4 μg/ml cotylenin A (2) for 4 days. Cells were then incubated with a fluorogenic caspase substrate PhiPhiLux-G1D2 for 60 min and then analyzed by flow cytometry. Faint line, untreated controls.

Fig. 4. A, cell growth in the presence of IFNα and cotylenin A. Cells were treated with 24 μg/ml cotylenin A (1), 600 IU/ml IFNα (2), or cotylenin A plus IFNα (3) for 4 days, and DNA histograms were then analyzed. The apoptotic cell population is shown according to the sub-G₁ fraction. B, increase in cells with caspase-3 activity by cotylenin A plus IFNα but not by cotylenin A. Cells were treated with 24 μg/ml cotylenin A (1) or 300 IU/ml IFNα plus 4 μg/ml cotylenin A (2) for 4 days. Cells were then incubated with a fluorogenic caspase substrate PhiPhiLux-G1D2 for 60 min and then analyzed by flow cytometry. Faint line, untreated controls.

Fig. 5. A, induction of G₁ arrest and apoptosis in A549 cells treated with a high concentration ofcotylenin A and a combination of cotylenin A and IFNα, respectively. Cells were cultured without (1) or with 24 μg/ml cotylenin A (2) or a combination of 300 IU/ml IFNα and 4 μg/ml cotylenin A (3) for 4 days, and DNA histograms were then analyzed. The apoptotic cell population is shown according to the sub-G₁ fraction. B, increase in cells with caspase-3 activity by cotylenin A plus IFNα but not by cotylenin A. Cells were treated with 24 μg/ml cotylenin A (1) or 300 IU/ml IFNα plus 4 μg/ml cotylenin A (2) for 4 days. Cells were then incubated with a fluorogenic caspase substrate PhiPhiLux-G1D2 for 60 min and then analyzed by flow cytometry. Faint line, untreated controls.

Fig. 6. A, cell growth in the presence of IFNα and cotylenin A. Cells were treated with 24 μg/ml cotylenin A (1), 600 IU/ml IFNα (2), or cotylenin A plus IFNα (3) for 4 days, and DNA histograms were then analyzed. The apoptotic cell population is shown according to the sub-G₁ fraction. B, increase in cells with caspase-3 activity by cotylenin A plus IFNα but not by cotylenin A. Cells were treated with 24 μg/ml cotylenin A (1) or 300 IU/ml IFNα plus 4 μg/ml cotylenin A (2) for 4 days. Cells were then incubated with a fluorogenic caspase substrate PhiPhiLux-G1D2 for 60 min and then analyzed by flow cytometry. Faint line, untreated controls.
and cotylenin A in two types of normal human lung epithelial cells, normal human bronchial epithelial and SAE cells, which were derived from human lung bronchus and the small airway of healthy donors, respectively, with those in lung cancer cell lines (Fig. 7). The optimal culture conditions for the normal cells were serum-free medium, whereas lung cancer cells grow in the presence of 10% FBS. In serum-free culture conditions, cells were more sensitive to the combined treatment with IFNα and cotylenin A. Therefore, we examined the effects on the growth of normal and cancer cells in both the presence and absence of FBS. Most of the cancer cells also grew in the serum-free medium for normal epithelial cells. In this condition, cancer cells were more sensitive to treatment, whereas weak growth inhibition was seen in normal lung epithelial cells (Fig. 7). Normal lung epithelial cells were still less sensitive to this treatment in the presence of FBS (data not shown). When cultured with 4 μg/ml cotylenin A and 300 IU/ml IFNα for 6 days, A549 cells underwent morphological changes characteristic of apoptosis, such as rounding, detachment, and floating, whereas the morphology of normal lung epithelial cells did not change, although they did show a slight decrease in cell number. Basal mRNA levels of DcR1 and DcR2, two decoy receptors for the death ligand TRAIL (32–34), which antagonize its action, were higher in both normal lung epithelial cells than in cancer cells. The expression of DcR1 mRNA in A549 cells was about one-tenth that in SAE cells, whereas the DcR2 mRNA level was one-half that in SAE cells (Fig. 8). mRNA levels of death receptors DR4 and DR5 in normal lung epithelial cells were also higher than those in cancer cells (Fig. 8).

Up-Regulation of TRAIL and DR5 Receptor mRNA Expression by Cotylenin A and IFNα in Cancer Cells. To determine whether cotylenin A and IFNα affected mRNA levels, quantitative RT-PCR analysis was carried out on normal lung epithelial and lung carcinoma A549 cells. The amounts of DcR1 and DcR2 mRNA were not essentially affected by IFNα, cotylenin A, or IFNα plus cotylenin A, whereas the osteoprotegerin mRNA level was significantly reduced by these treatments in A549 cells (Fig. 8c). The mRNA expression in normal lung epithelial cells was not affected by these treatments (data not shown). The expression of DR5 mRNA was significantly induced by the combination of IFNα and cotylenin A and was comparable with that in normal lung epithelial cells (Fig. 8a), whereas the up-regulation of the expression of DR4 mRNA was moderate (Fig. 8c). IFNα alone significantly up-regulated TRAIL mRNA expression, although combined treatment with IFNα and cotylenin A was more effective (Fig. 8b). Similar results were seen in PC9 lung carcinoma cells (data not shown). These results are compatible with those from flow cytometric analysis of the cell surface expression of TRAIL and its receptors (Fig. 6).

Effects of Cotylenin A and IFNα on the in Vivo Growth of PC14 Cells as Xenografts. The in vitro studies described above suggested that combined treatment with cotylenin A and IFNα should be more effective therapeutically than treatment with cotylenin A or IFNα alone. At day 7 after the inoculation of human lung adeno carcinoma PC14 cells, the mean tumor volume was 33.4 ± 12.8 mm³ (±SD), and treatments were then started. Because of the low solubility of cotylenin A, a dose-escalating effect was not observed in the therapeutic experiments (20). Therefore, we administered 100 μg/mouse cotylenin A, which had no appreciable adverse effects on mice, even in the presence of IFNα. The mice were injected with 3 × 10⁶ IU of IFNα every day. This dose is equivalent to ∼3 × 10⁶ IU/m² daily in humans using the calculations of Freireich et al. (35). The combined treatment significantly inhibited the growth of PC14 cells as xenografts (Fig. 9). At day 12 after treatment, the mean tumor volumes of untreated, cotylenin A-, IFNα-, and cotylenin A plus IFNα-treated nude mice were 520.4 ± 82.7, 318.5 ± 65.4, 391.6 ± 61.5, and

![Graph](image_url)

Fig. 4. Proliferation of A549 cells in long-term culture with IFNα and cotylenin A. Cells were cultured without (●) or with 300 IU/ml IFNα (○), 4 μg/ml cotylenin A (△), 24 μg/ml cotylenin A (■), or 300 IU/ml IFNα plus 4 μg/ml cotylenin A (●), 8 and ▼ represent cells that were cultured with 300 IU/ml IFNα plus 4 μg/ml cotylenin A and 24 μg/ml cotylenin A for 7 days and then washed and cultured further without drugs, respectively. The culture medium was replaced by fresh medium at least once every 3 days. The cell density was kept at 1–8 × 10⁴/ml. The values are means of four separate experiments.
Although cotylenin A and IFNα each significantly retarded tumor growth (P < 0.05), combined treatment induced tumor regression. The treatment was continued for 12 days and then stopped, with a follow-up on day 26. All of the untreated mice had a large tumor burden at day 26. On the other hand, >50% of the treated mice escaped from the disease (13 of 20 mice), and the rest had only a small tumor burden, suggesting that the therapeutic effects were still maintained after treatment was terminated. These results indicate that the combination of cotylenin A and IFNα is more effective therapeutically than treatment with cotylenin A or IFNα alone, and the combined treatment had a significant antitumor effect (P < 0.001).

Fig. 6. Combined treatment with cotylenin A and IFNα up-regulates TRAIL and DR5 expression in A549 cells. Cells were treated with various concentrations of cotylenin A in the presence of 0 (□), 150 (●), 300 (▲), or 450 (▲) IU/ml IFNα for 4 days. Expression was assayed by flow cytometry. The values are mean ± SD of four determinations.
DISCUSSION

IFNα alone had no significant effects on the growth or viability of human lung cancer cell lines, even at 2,000 IU/ml. However, we found previously that several human lung cancer cell lines similarly expressed IFNα receptors (3,000–10,000 receptors/cell) with high affinity, and activation of the transcription factor ISGF-3, which has been shown to be required for the transcriptional activation of IFN-induced genes, was induced by IFNα alone in lung carcinoma PC9 cells (14). Moreover, IFNα rapidly induced the expression of some IFNα-inducible genes, such as IRF-1 and PML, in PC9 cells (data not shown). The present study also showed that the expression of TRAIL mRNA was induced by IFNα alone (Fig. 8), suggesting that resistance and its restoration by differentiation-inducing agents, such as cotylenin A and DMSO, in some lung carcinoma cells act downstream of the activation of ISGF-3. Although IFNα alone up-regulated the expression of some genes associated with apoptosis and growth inhibition, it might be inadequate for activating some important genes to exert significant effects on apoptosis. Expression of the death receptor DR5 was not essentially induced by IFNα or cotylenin A alone, but combined treatment greatly up-regulated its mRNA and cell surface expression in lung cancer cells. Wild-type p53 has been shown to up-regulate DR5 gene expression (36). However, expression of DR5 protein expression in A549 cells was not up-regulated by some anticancer drugs that up-regulate p53 protein expression, and the combined effects of cotylenin A and IFNα were observed in both cancer cells with wild-type and mutant/deleted p53. A receptor of fusicoccin, closely related to cotylenin A, has been reported to be a member of the family of 14-3-3 proteins that are commonly found in a huge array of signaling and regulatory pathways (37). A special subfamily of 14-3-3 proteins may bind cotylenin A and affect the interaction with some signaling molecules, including ISGF-3. This modification may lead to up-regulation of DR5 gene expression. However, further investigation is required to explain how up-regulation of DR5 is related to its effects on the 14-3-3 signaling pathway.

To investigate the role of IFNα in the induction of apoptosis in human lung carcinoma cells, we examined the effects of IFNα and/or cotylenin A...
and IFN\(^\text{c}\) the expression of TRAIL mRNA, and the combination of cotylenin A in cells, even in the presence of IFN\(^\text{c}\)/H9251 resembling in structure to some extent, the antitumor triterpenoids including NSCLC (39, 40). Because cotylenin A is a diterpenoid and effective in synergism with IFN\(^\text{c}\)/H9251 by treatment with IFN\(^\text{c}\) and cotylenin A (data not shown). The expression of caspase-4 mRNA was induced by treatment with IFN\(^\text{c}\) alone but not by treatment with cotylenin A. The combination of IFN\(^\text{c}\) and cotylenin A did not cause an additional increase in the expression of caspase-4 mRNA. However, treatment with an inhibitor of caspase-4 did not affect the apoptosis induced by IFN\(^\text{c}\) and cotylenin A (data not shown). The expression of other caspase, Apaf-1, and bcl-2 family genes was unchanged by IFN\(^\text{c}\) and/or cotylenin A (data not shown). IFN\(^\text{c}\) alone significantly induced the expression of TRAIL mRNA, and the combination of cotylenin A and IFN\(^\text{c}\) caused an additional increase in expression. The gene expression of the TRAIL receptor DR5 was greatly induced by the combination of cotylenin A and IFN\(^\text{c}\) (Fig. 8). TRAIL is one of the early genes induced by IFN in apoptosis-sensitive melanoma and lymphoma cells, and apoptosis is mediated by the autocrine and/or paracrine loop involving TRAIL and its receptors (26, 27). These results suggest that the activation of TRAIL and DR5 genes is an important process in the induction of apoptosis in NSCLC cells by cotylenin A plus IFN\(^\text{c}\).

Although natural retinoids did not affect the growth of NSCLC cells, even in the presence of IFN\(^\text{c}\), some synthetic retinoids might be effective in synergism with IFN\(^\text{c}\). CD347 (6-[3-(1-adamantyl)-4-hydroxyphenyl]-2-naphthalene carboxylic acid) induces gene expression of DR4 and DR5 and then the apoptosis of NSCLC cells (38). However, normal lung epithelial cells are less sensitive to the retinoid. CD347 and IFN\(^\text{c}\) might cooperatively induce apoptosis in lung cancer cells. Some triterpenoids induce apoptosis of several tumor cell lines, including NSCLC (39, 40). Because cotylenin A is a diterpenoid and resembling in structure to some extent, the antitumor triterpenoids may also cooperate with IFN\(^\text{c}\) in inducing apoptosis of cancer cells. Recent published results indicate that TRAIL and chemotherapeutic drugs act synergistically to kill cancer cells (22, 41). In the present study, the cells were more sensitive to the combination of cotylenin A and IFN\(^\text{c}\) than to chemotherapeutic drugs and TRAIL. We did not investigate the maximal tolerable dose or dose-limiting toxicity of cotylenin A in the absence or presence of IFN\(^\text{c}\), because the maximal concentration of cotylenin A was 100 μg/0.2 ml saline. This treatment has no apparent effects on mice (body weight and behavior). Potent derivatives of cotylenin A that are readily soluble in saline will be required to further develop this therapeutic strategy.

REFERENCES

Cotylenin A, a Differentiation-inducing Agent, and IFN-α Cooperatively Induce Apoptosis and Have an Antitumor Effect on Human Non-Small Cell Lung Carcinoma Cells in Nude Mice

Yoshio Honma, Yuki Ishii, Yuri Yamamoto-Yamaguchi, et al.


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