Abstract. Although progress has been made in chemotherapeutic strategies against pancreatic cancer, overall survival has not significantly improved over the past decade. Thus, the development of better therapeutic regimens remains a high priority. Pancreatic cancer cell lines were treated with tamoxifen, a novel antitumor fusicoccin derivative (ISIR-042), and anticancer drugs, and their effects on cell growth, signaling and gene expression were determined. Xenografts of Panc-1 cells were treated with tamoxifen, ISIR-042 and 5-fluorouracil (5FU) to determine the effects on tumor growth. The inhibition of the growth of pancreatic cancer cells induced by tamoxifen was effectively reduced by \( \alpha \)-tocopherol, a membrane stabilizer. ISIR-042 produced synergistic effects with tamoxifen in inhibiting cell growth. Tamoxifen elevated lipid peroxidation and the release of cytochrome c, and these effects of tamoxifen were reduced by \( \alpha \)-tocopherol. ISIR-042 significantly inhibited colony formation and the expression of stemness-related genes of pancreatic cancer cells. The triple combination of tamoxifen, ISIR-042, and 5FU or gemcitabine was effective at inhibiting cell growth and the appearance of drug-resistant cells. This combined treatment significantly inhibited the growth of Panc-1 cells as xenografts without apparent adverse effects. The triple combination of tamoxifen and ISIR-042 with 5FU or gemcitabine may be highly effective against pancreatic cancer by overcoming resistance to therapy.

Introduction

Pancreatic cancer remains a highly lethal neoplasm, and even with multimodality therapy for localized disease, patient survival is measured in months. 5-Fluorouracil (5FU) and gemcitabine are the mainstays of therapy. However, the response rates to either are <30%, and their toxic effects are significant (1,2). Although the FOLFIRINOX regimen has produced substantial benefits in the treatment of metastatic pancreatic cancer, it is associated with more adverse effects (3). Chemoresistance usually develops in patients who respond initially, and an effective salvage therapy is currently unavailable. Thus, the development of better therapeutic regimens for pancreatic cancer remains a high priority.

At the start of therapy, large pancreatic cancers always contain mixed populations of malignant cells that are resistant to anticancer agents, and these cells expand clonally once therapy is initiated. Tumor recurrence is thus a fait accompli when single agents are used. Combination therapy eliminates cells that are resistant to either drug alone, if there are no single mutations that cause cross-resistance to both drugs. Therefore, combination therapy with two drugs greatly increases the chance of success, since the likelihood of a doubly mutated cell emerging in such a population is low. However, mathematical analyses have revealed that triple therapy may be needed in patients with a large tumor burden (4).

Cotylenins, fusicocci and some diterpene glucosides are modulators of 14-3-3 proteins, and have been shown to exhibit antitumor effects in vitro and in vivo. We previously synthesized many fusicoccin derivatives and found that ISIR-042 was the most potent at inhibiting the proliferation of tumor cells (5). ISIR-042 in combination with 5FU or gemcitabine was observed to have synergistic effects at inhibiting the growth of pancreatic cancer cells. In pancreatic cancers, late diagnosis and the early development of metastasis to the regional lymph nodes and liver limit curative resection to <10% of patients. Since most of the patients have a large tumor burden, combination therapy with two drugs may be insufficient in these patients (1,2). While the FOLFILINOX regimen has been introduced, this regimen is more toxic and few patients can receive it. Triple combination therapy with less toxicity may be required for an effective therapy against pancreatic cancers. Therefore, we examined the synergistic effects of various compounds and ISIR-042 on the growth of pancreatic cancer cells to identify the most potent and clinically applicable drugs. The

 Combined treatment with tamoxifen and a fusicoccin derivative (ISIR-042) to overcome resistance to therapy and to enhance the antitumor activity of 5-fluorouracil and gemcitabine in pancreatic cancer cells

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most effective agent was tamoxifen. Tamoxifen is a selective estrogen receptor modulator that is used as the first-line treatment for estrogen receptor-positive breast cancer. However, multiple non-estrogen receptor-mediated mechanisms have been implicated in antitumor effects induced by tamoxifen in estrogen receptor-negative tumors (6). There seems to be a consistent relationship between higher doses of tamoxifen and longer survival in patients with largely inoperable and recurrent malignant glioma (7). Therefore, in the present study, we sought to clarify the synergistic effects of ISIR-042 and tamoxifen on human pancreatic cancer cells and to examine the therapeutic effects on xenografts of human pancreatic carcinoma cells in the presence or absence of 5FU treatment.

Materials and methods

Materials. ISIR-042 was prepared as previously described (5). Gemcitabine, tamoxifen, 5FU, α-tocopherol and 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyltetrazolium bromide (MTT) were purchased from Sigma-Aldrich Japan (Tokyo, Japan). Taurine, L-nitroarginine methyl ester (L-NAME), N-acetyl cysteine (NAC), and protease inhibitor cocktail were obtained from Wako Pure Chemical (Tokyo, Japan).

Cells and culture. Human pancreatic cancer cell lines CFPAC-1, BxPC-3 and Capan-2 were purchased from American Tissue Culture Collection (Manassas, VA, USA). Cells were cultured in RPMI-1640 medium (Sigma-Aldrich Japan) supplemented with 10% fetal bovine serum and 80 µg/ml gentamycin at 37˚C in a humidified atmosphere of 5% CO2 in air.

Colony-forming assay. MIAPaCa-2 cells (1x104 per dish) were plated into 1.1 ml of a semisolid methylcellulose medium with 0.8% methylcellulose and 20% fetal bovine serum in triplicate for 14 days. A solution of 0.2 ml of PBS containing various concentrations of drugs was added to the semisolid medium. Colonies were photographed under an inverted microscope. Colonies in enlarged photographs were measured and counted. Colonies >0.4 mm in diameter were counted. Bone marrow plugs from a single femur of three BALB/c mice were pooled in RPMI-1640 medium. A single cell suspension was prepared by vigorous pipetting. Nucleated cells (1,000 cells/ml/dish) were placed into a semi-solid medium containing hematopoietic growth factors (MethoCult® GF M3434, Stem Cell Technology Inc., Vancouver, BC, Canada) and incubated for 10 days.

Assay of cell growth. Human cancer cells were seeded into 24-well multidishes at 1.0x104 cells/ml/well. The cells were cultured with various concentrations of drugs for 4-6 days. After culture with or without drugs, viable cells were examined by the MTT assay (5).

Analysis of the effects of combinations of drugs. An isobologram analysis was used to determine the effects of combinations of drugs on Panc-1 cells. Dose-dependent effects were determined for each compound and for one compound with fixed concentrations of another. The interaction of two compounds was quantified by determining the combination index (CI), in accordance with the following classic isobologram (8).

Assay of the cumulative cell number. The cell density of the drug-treated cells was kept at 2.8x104/ml to maintain the growth phase in a 24-well multidish (Falcon). The medium of treated cultures was replaced by fresh medium with drugs at least every 4 or 5 days to remove cell debris of dead cells. The viable cell number was measured by MTT assay. The cumulative cell number was calculated from the MTT values and the dilution used when feeding the culture.

Western blotting. Cells were packed after being washed with cold PBS and then lysed at 1.5x106 cells/ml in sample buffer. The resultant lysate was resolved on SDS-polyacrylamide gel, as described elsewhere (5). Primary antibodies were obtained from the following sources: K-ras (Abnova, Taiwan); acid ceramidase (BD Biosciences Japan, Tokyo, Japan); and cytochrome c and actin (Cell Signaling Technology Japan, Tokyo, Japan).

Mitochondria preparation. After cells were washed twice with PBS, protease inhibitor cocktail was added and cells were homogenized by 30 strokes with a glass homogenizer. The homogenate was centrifuged at 1,000 x g and the pellet was discarded. The supernatant was centrifuged at 10,000 x g for 30 min and the cytoplasmic (supernatant) and mitochondrial (pellet) fractions were separated.

Cytochrome c release and lipid peroxidation. Cytochrome c was determined by western blot analysis using monoclonal anti-cytochrome c antibody. Lipid peroxidation (LPO) was determined by measuring thiobarbituric acid-reactive substances (TBARS) at 535 nm, as described elsewhere (9).

Reverse transcription-polymerase chain reaction (RT-PCR). Total RNA was extracted from cells using TRI reagent (Sigma). Total RNA was converted to first-strand cDNA primed with random hexamer in a reaction volume of 20 µl using an RNA PCR kit (qPCR RT Master Mix, Toyobo Co. Ltd., Osaka, Japan), and 2 µl of this reaction was used as a template in real-time PCR. The primers used were previously described (10).

Transplantation of human cancer cells into nude mice. Four-week-old female athymic nude mice with a BALB/c genetic background were obtained from CLEA Japan (Tokyo). They were housed under specific pathogen-free conditions. Mice were subcutaneously inoculated with 2x106 Panc-1 cells. The adjusted 2x104 cells/ml were mixed with an equal volume of Matrigel (BD Sciences) and 0.2 ml of the cell solution was injected subcutaneously (11). Mice were given daily intraperitoneal injections of 3 mg/kg ISIR-042 in 0.2 ml PBS and/or 2 mg/kg tamoxifen in 0.1 ml corn oil, with the first treatment given 2 days after the inoculation of tumor cells (11). Treatment with 5FU was initiated when tumors in each group achieved an average volume of 80 mm3. Treatment with 5 mg/kg 5FU was performed three times per week. Tumor volume was measured with vernier calipers. Our protocol was approved by the animal ethics committee at Shimane University.

Statistical analysis. The results are expressed as means ± standard deviation (SD). Pairs of data were compared using Student’s t-test. Significant differences were considered to exist for probabilities <5% (P<0.05). For the in vivo experiment, an
F-test was performed to demonstrate statistical significance. Again, significant differences were considered to exist for probabilities <5%.

**Results**

**Combined effects of tamoxifen and ISIR-042 on the growth of pancreatic cancer cells.** Tamoxifen inhibited the growth of Panc-1 cells in a concentration-dependent manner, and the combined effects of tamoxifen and ISIR-042 are shown in Fig. 1. ISIR-042 produced synergistic effects with tamoxifen (Fig. 1A) and the results were confirmed by isobologram analysis. Fig. 1B shows isoboles for the combination of tamoxifen with ISIR-042 that were isoeffective (IC₅₀) for inhibiting the proliferation of Panc-1 cells. The dashed line indicates zero interaction of the isobole. (C) Combined effects on other human pancreatic cancer cells. Cells were treated with various concentrations of tamoxifen in the presence of 0 (◆), 2 (◼), 4 (▲), or 6 (●) μg/ml of ISIR-042 for 5 days.

Tamoxifen enhances LPO and α-tocopherol prevents the action of tamoxifen. Multiple non-estrogen receptor-mediated mechanisms have been implicated in cell death induced by tamoxifen (6). These mechanisms include changes in intracellular calcium, modulation of protein kinase C, changes in calmodulin activity, signaling though mitogen-activated protein kinases including ERK, K-ras degradation (12) and inhibition of acid ceramidase activity (13). To understand the growth-inhibitory effect of tamoxifen on pancreatic cancer cells, we examined the effects of various compounds including estrogen and inhibitors of signaling on the growth of MIAPaCa-2 cells in the presence of tamoxifen and/or ISIR-042. Estrogen did not affect the growth-inhibitory effect.
Several inhibitors of protein kinases also had no effect, whereas α-tocopherol effectively reduced tamoxifen-induced growth inhibition but not ISIR-042-induced growth inhibition (Fig. 2A). Taurine, another membrane stabilizer and antioxidant, also reduced the growth inhibition induced by tamoxifen, although this effect was less than that of α-tocopherol (Fig. 2B). When MIAPaCa-2 cells were incubated with tamoxifen in the presence of the antioxidant NAC, tamoxifen-induced growth inhibition was not affected by NAC (Fig. 2C). However, treatment with NAC effectively inhibited the growth inhibition induced by the Japanese apricot extract MK615 (11) (Fig. 2C). Although the inhibitor of nitric oxide synthase L-NAME alone did not significantly affect tamoxifen-induced growth inhibition (Fig. 2D), L-NAME effectively reduced tamoxifen-induced growth inhibition in the presence of a suboptimal concentration of α-tocopherol (Fig. 2E). These results suggest that tamoxifen-induced changes in membrane dynamics are associated with growth inhibition in pancreatic cancer cells.

A recent study indicated that tamoxifen induced autophagic K-Ras degradation in many cancer cells including pancreatic cancer cells (12). Since pancreatic cancers exhibit an extremely high mutation rate of K-Ras (>90%) (14), we examined the effect of α-tocopherol on the degradation of K-Ras protein induced by tamoxifen in the presence or absence of ISIR-042. Tamoxifen concentration-dependently induced K-Ras degradation, and α-tocopherol effectively counteracted this tamoxifen-induced degradation (Fig. 3A). ISIR-042 did not essentially affect the effects of α-tocopherol and tamoxifen. These results suggest...
that K-ras degradation is closely associated with growth inhibition in tamoxifen-treated cells. However, no significant K-Ras degradation was observed until 8 h after treatment with tamoxifen, suggesting that K-Ras degradation is not the cause of growth inhibition, but rather the result of growth inhibition, in tamoxifen-treated cells (Fig. 3b).

Acid ceramidase (AC) is a central player in ceramide metabolism. Since AC regulates the levels of pro-apoptotic ceramide and mitogenic sphingosine-1-phosphate, it is considered to be an apt target in cancer therapy. A recent study indicated that tamoxifen induces the downregulation of AC protein in cancer cells including pancreatic cancer cells (13). Therefore, we examined the effect of tamoxifen on AC protein in MIAPaCa-2 cells (Fig. 3B).

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α-tocopherol is a membrane stabilizer and a lipophilic antioxidant (17). Therefore, we examined the effects of tamoxifen on lipid peroxidation and cytochrome c release in the presence or absence of α-tocopherol. Stimulation of mitochondrial NOS increases mitochondrial peroxynitrite, which releases cytochrome c from the mitochondria (18). The stimulation of mitochondrial NOS also increases lipid peroxidation (LPO), which is a widely used biomarker of peroxynitrite (18,19). Fig. 4A shows that tamoxifen increased LPO within 4 h of treatment in MIAPaCa-2 cells and that the effects of tamoxifen were prevented when samples were co-treated with α-tocopherol. The increase in LPO in mitochondria induced by tamoxifen was more evident than that in...
whole cells (Fig. 4B and C). Furthermore, tamoxifen released cytochrome c from the particulate fraction which included mitochondria in MIAPaCa-2 cells (Fig. 4D).

**Effect of ISIR-042 on colony formation by MIAPaCa-2 cells and the expression of stemness-related genes.** Pancreatic cancer stem cells express the cell surface markers CD24 and CD44 (20,21). Gemcitabine showed an increased percentage of CD24⁺CD44⁺ cells in MIAPaCa-2 cells, whereas ISIR-042 significantly reduced the percentage of CD24⁺CD44⁺ cells, suggesting that ISIR-042 preferentially inhibits stem/progenitor cells in pancreatic cancer cells (5). Therefore, we examined the effects of ISIR-042 on colony formation by MIAPaCa-2 cells in semi-solid culture and the expression of stemness-related genes. ISIR-042 effectively inhibited colony formation by MIAPaCa-2 cells in semi-solid culture and the expression of stemness-related genes. ISIR-042 effectively inhibited colony formation by MIAPaCa-2 cells in semi-solid culture and the formation of large colonies was inhibited more evidently than that of small colonies (Fig. 5A). Colony formation in semi-solid culture was more efficiently inhibited by ISIR-042 than cell proliferation in liquid culture, whereas the inhibition of colony formation by tamoxifen or 5FU was similar to the inhibition of cell proliferation (Fig. 5B).

The growth of pancreatic cancer cell lines was effectively inhibited by ISIR-042, whereas only weak growth-inhibition was seen in normal endothelial HUVEC cells (5). Cotylenin A, the parent compound of ISIR-042, preferentially induced apoptosis in human cancer cells, while sparing normal cells (22). We examined the effects of ISIR-042 on colony formation by normal mouse bone marrow cells in the presence or absence of tamoxifen. Although ISIR-042 alone hardly affected colony formation by bone marrow cells, the combination of ISIR-042 and 4 µM tamoxifen only weakly inhibited colony formation by normal bone marrow cells (Fig. 5C). These results suggest that pancreatic cancer cells were more sensitive to combined treatment with ISIR-042 and tamoxifen than normal bone marrow cells.

Since the expression of pluripotency-associated transcription factors has been implicated in the maintenance of cancer stem cells, we evaluated mRNA expression of Sox-2, Oct3/4 and Nanog (Fig. 5D). The expression of Sox-2 and Nanog mRNAs was significantly suppressed by treatment with ISIR-042. This reduction was observed within 4 days. ISIR-042 did not significantly affect the expression of Oct3/4 mRNA under these conditions. Tamoxifen also reduced the expression level
of Nanog mRNA, although the effects were less than those of ISIR-042 (Fig. 5D).

**Combined effects of tamoxifen, ISIR-042 and anticancer drugs on growth inhibition and the appearance of drug-resistant cells.** The combined effects of ISIR-042, tamoxifen and anticancer drugs were examined. A triple combination (ISIR-042, tamoxifen and 5FU) cooperatively inhibited the growth of MIAPaCa-2 cells (Fig. 6A). 5FU at 1,000 ng/ml did not completely inhibit cell viability, and the combinations of 5FU and 6 µg/ml of ISIR-042 and the combination of 5FU and 4 µM tamoxifen did not produce 100% killing either. However, treatment with the triple combination led to no viable cells in the culture (as measured by MTT assay). Similar results were obtained when cells were treated with gemcitabine. Next, we examined the growth-inhibitory effects of triple combinations in long-term culture. The growth of Panc-1 cells was markedly reduced for 12 days by treatment with 500 ng/ml of 5FU alone. However, after ~16 days, growing cells reappeared and the proliferation rate of the treated cells became similar to that of untreated cells (Fig. 6B). Drug-resistant cells were not observed when cells were cultured for 28 days with 500 ng/ml 5FU, 3 µM tamoxifen and 3 µg/ml ISIR-042, and dual combinations (5FU+tamoxifen or 5FU+ISIR042) were less effective.
Similar results were obtained in the long-term culture of cells treated with gemcitabine (Fig. 6C). These results suggest that the triple combination therapy is effective for overcoming the emergence of drug-resistant cells in pancreatic cancers.
with 5FU alone. Treatments were started at day 2 after the inoculation of Panc-1 cells. We administered both tamoxifen and ISIR-042 daily by intraperitoneal injection. The combined treatment had no appreciable adverse effects in mice, including weight loss. Although neither ISIR-042 nor tamoxifen alone retarded tumor growth, combined treatment was more effective (Fig. 6D). Next, we examined the antitumor effects of the combination of 5FU with tamoxifen and ISIR-042. Treatment was initiated when tumors in each group achieved an average volume of 80 mm³. Mice were treated with 5 mg/kg 5FU three times per week.

While treatment with 5FU alone induced a slight loss of body weight (6-8% loss), treatment with the triple combination did not induce a further reduction of body weight. Although 5FU alone retarded tumor growth, combined treatment with 5FU, tamoxifen and ISIR-042 was more effective (Fig. 6E). Statistical analysis (F-test) revealed that the difference was significant (P<0.05 at 6 weeks). There were no significant differences between treatment with 5FU and combination treatment with 5FU+tamoxifen or 5FU+ISIR042. These results indicate that the combination of tamoxifen and ISIR-042 is effective therapeutically, consistent with the in vitro findings.

Discussion

Tamoxifen induces apoptosis in both ER-positive and ER-negative cells (6). Treatment of human pancreatic cancer cells with 2-8 μM tamoxifen, concentrations which are 6-25 times greater than the therapeutic concentration (the average therapeutic blood concentration in patients with breast carcinoma was 0.3 μM) inhibited cell growth. Treatment with α-tocopherol prevented the growth-inhibition induced by tamoxifen (Fig. 3A). Submicromolar concentrations of tamoxifen released cytochrome c from isolated mitochondria and increased LPO in isolated mitochondria (16). The release of cytochrome c and increased LPO in tamoxifen-treated cells was inhibited by α-tocopherol (Fig. 5). Since the action of tamoxifen on cancer cells is counteracted by the potent membrane stabilizer α-tocopherol, the main mechanism of action of tamoxifen is considered to be membrane damage via LPO. These results suggest that tamoxifen is an anticancer drug that induces oxidative stress and apoptosis via a mitochondria-dependent pathway.

The novel fusicoccin derivative ISIR-042 was synthesized from fusicoccin A as an antitumor drug (5). A receptor of fusicoccin has been reported to be a member of a family of 14-3-3 proteins that are found commonly in a huge array of signaling and regulatory pathways (23). The 14-3-3 proteins bind to discrete phosphoserin-containing motifs that are present in many signaling molecules. The 14-3-3 proteins are associated with dynamic nucleocytoplasmic shuttling. Phosphorylation-dependent binding of 14-3-3 proteins to telomerase promotes its nuclear localization (24). However, the reverse has more often been the case, in that many nuclear proteins can become phosphorylated, bind to 14-3-3 proteins and accumulate in the cytoplasm. The 14-3-3 proteins negatively regulate histone deacetylase 4 (25) or the cyclin-dependent kinase inhibitor p27 (26) by preventing its nuclear localization. Fusicoccins bind to inhibitory 14-3-3 interaction sites of c-RAF, pSer259, but not to the activating interaction site, pSer621 (27). The modification by fusicoccins may lead to the upregulation of apoptosis-regulating genes and the downregulation of cell cycle- and stemness-regulating genes. Our previous findings indicate that the pancreatic cancer stem cell marker CD24+CD44+ is significantly reduced by ISIR-042 in a MiaPaCa-2 cell population (5).

In the present study, ISIR-042 preferentially inhibited the population that formed large colonies, which may be cells with high self-renewal activity. Treatment with ISIR-042 effectively downregulated the stemness-associated transcription factors Sox-2 and Nanog. Expression of these genes is important for the development of tumors, as shown in in vivo experiments. Knockdown of Nanog expression inhibits the stemness of pancreatic cancer cells (28). Sox-2 was highly expressed in the clonogenic compartment of pancreatic cancer cells and targeting of Sox-2 could be a worthy strategy for pancreatic cancer therapy (29). Pancreatic carcinoma remains incurable for the vast majority of patients, suggesting that pancreatic cancer stem cells are relatively resistant to chemotherapy. ISIR-042 effectively inhibits clonogenic growth and the expression of stemness-related genes in pancreatic cancer cells. ISIR-042 effectively sensitizes pancreatic cancer cells to 5FU or gemcitabine (5). These results suggest that ISIR-042 is a useful drug for cancer stem cell-targeted therapy against pancreatic carcinoma.

After estrogen receptors were detected in human pancreatic carcinomas (30-32), tamoxifen was evaluated in several trials. These studies reported an increased survival time in patients with resected and non-resected pancreatic cancer after treatment with tamoxifen (33-35). Furthermore, phase II trials of tamoxifen with 5FU or gemcitabine have been performed in pancreatic cancer (36,37). These data suggest that tamoxifen has some positive effects in the treatment of pancreatic cancer, although the effect is modest. The present results suggest that the addition of ISIR-042 to tamoxifen potentiates its anti proliferative action, which provides a synergistic effect. Toxicity studies with mice and rats have suggested that ISIR-042 is well-tolerated (5). Furthermore, combined treatment with tamoxifen and ISIR-042 did not further affect the toxicity of 5FU, as judged by body weight loss. Supplementation of 5FU- or gemcitabine-based chemotherapy against pancreatic cancers with tamoxifen and ISIR-042 may provide a stronger beneficial effect.

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References


